

Showcasing research from Dr Thomas Sheppard's laboratory, Institute for Chemical Technology and Polymer Chemistry (ITCP), Karlsruhe Institute of Technology (KIT), Germany.

Synchrotron PXRD deconvolutes nickel particle and support changes in Ni/ZrO₂ methanation catalysts





Synchrotron radiation is applied to understand multiple modes of catalyst deactivation in a single method approach. *Operando* synchrotron powder X-ray diffraction allows to follow phase changes during activation, reaction conditions of CO₂ methanation and artificial thermal aging steps of a designed Ni/ZrO₂ catalyst series. Furthermore, the evolution of Ni and ZrO₂ crystallite sizes can be followed independently by PXRD. By correlating to activity data, the catalyst series showed deactivation dominated either by changes in Ni active metal or ZrO₂ support, depending on the thermal stability of the initial support structure.

As featured in:



See Thomas L. Sheppard *et al.*,
Catal. Sci. Technol., 2022, **12**, 6069.

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *Catal. Sci. Technol.*, 2022,
12, 6069Synchrotron PXRD deconvolutes nickel particle
and support changes in Ni/ZrO₂ methanation
catalysts†Mariam L. Schulte, ^{†ab} Sebastian Weber, ^{†ab} Linda Klag,^a
Jan-Dierk Grunwaldt ^{ab} and Thomas L. Sheppard ^{*ab}

Understanding catalyst deactivation is important for future knowledge-based design of catalysts with improved stability. Deactivation by thermal aging is particularly relevant for exothermic reactions, here demonstrated with CO₂ methanation using nickel-based catalysts. A series of five Ni/ZrO₂ catalysts is studied which differ by calcination temperature of the ZrO₂ support prior to Ni deposition, leading to different textural properties of the support. Artificial thermal aging of the Ni/ZrO₂ series is then performed and monitored by *operando* synchrotron-based powder X-ray diffraction (SPXRD). This reveals the dependence of catalyst stability on the support properties and shows that different deactivation mechanisms take place. Catalyst deactivation is mainly correlated either to changes of the support or to changes in nickel crystallite size, depending on the calcination temperature of the support before nickel deposition. By preparing a targeted series of supports with different textural properties, it is possible to deconvolute these effects. *Operando* SPXRD is shown as a powerful tool to follow both textural and structural changes during thermal catalyst deactivation, which is mostly only performed by *post mortem ex situ* analysis.

Received 29th May 2022,
Accepted 6th July 2022

DOI: 10.1039/d2cy00972b

rsc.li/catalysis

1 Introduction

Exothermic heterogeneous catalytic processes are challenging in regard to safety aspects and catalyst stability. Reliable heat management is necessary in order to maintain safe operation under process conditions. Thermal catalyst deactivation can be classified in three effects: (i) loss of active sites due to metal particle agglomeration; (ii) collapse of support structure leading to a decrease in surface area or porosity changes; (iii) formation of undesired alloy or mixed metal oxide phases, by reaction of the active phase and catalyst support.^{1,2} Distinguishing between these three effects for a given chemical process is difficult, since all effects can occur simultaneously and in addition to other types of deactivation, e.g., poisoning or fouling. Detailed investigation of temperature-induced catalyst deactivation is needed in order to unravel the impact of each degradation process.

A suitable exothermic model reaction for studying catalyst deactivation is CO₂ hydrogenation to CH₄, also known as CO₂ methanation. This process involves significant heat generation and therefore forms distinct temperature gradients (hot-spots),³ which have been simulated in fixed-bed reactors⁴ and observed experimentally using spatially-resolved measurements.⁵ A wide range of catalysts have been investigated for CO₂ methanation and the reviews of Aziz *et al.*⁶ and Frontera *et al.*⁷ provide a comprehensive overview on catalyst development. Ni-based catalysts are most frequently studied, due to relatively low cost of the metal and high activity and selectivity in CO and CO₂ methanation.⁷

Possible deactivation pathways for CO₂ methanation catalysts have also been reviewed.^{8–10} These mainly include fouling/coking, poisoning, (redox) reaction of the active metal sites, and thermal degradation. Although coking is discussed as a deactivation phenomena for CO₂ methanation, it is more relevant for CO or mixed CO/CO₂ methanation where the Boudouard reaction favors coke formation from CO-containing feed.^{8–11} Poisoning is mainly relevant for applications of CO₂ methanation associated with coal or biomass based CO₂ feeds, and can be attributed to sulfur species present in the feed.^{8,10,12} Changes in oxidation state of active metallic Ni sites are discussed in several studies especially in terms of fluctuating feeds such as H₂ dropout scenarios, which can lead to irreversible or partially

^a Institute for Chemical Technology and Polymer Chemistry, Karlsruhe Institute of Technology (KIT), Engesserstr. 20, 76131 Karlsruhe, Germany.E-mail: thomas.sheppard@kit.edu; Tel: +49 72160847989^b Institute of Catalysis Research and Technology, Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Germany† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2cy00972b>

‡ Both authors contributed equally.

irreversible Ni oxidation to NiO.^{13–17} However, the above deactivation pathways can generally be reversed by reactivation, compared to thermal degradation,¹⁸ which is a permanent feature due to the exothermic nature of the reaction.

As an example, long-term tests were reported for co-precipitated Ni/Al₂O₃ catalysts at reaction temperatures of around 380–400 °C by Abelló *et al.*¹⁹ and Koschany *et al.*,²⁰ where a slow decline in activity was observed over several hundreds of hours of operation, eventually leading to constant conversion after 320 h. This was traced back to gradual sintering of Ni particles, which Beierlein *et al.*¹⁸ studied in more detail during rapid ageing tests by exposing the catalysts to hydrothermal conditions at 600 to 700 °C. They observed increasing Ni particle sizes and additional changes in the support structure on thermal aging.¹⁸ In addition, potential deactivation by an oxidized or restructured Ni surface was discussed.¹⁸ Similar findings were reported by Mebrahtu *et al.*²¹ showing the influence of nickel hydroxide-like species formed under hydrothermal conditions, resulting in enhanced formation of mixed oxides or sintering. Ewald *et al.*²² attributed the long term deactivation of co-precipitated Ni/Al₂O₃ catalysts to Ni particle sintering, textural changes, *i.e.*, loss of surface area and reduction of medium basic sites, which is associated with changes of the CO₂ adsorption capacity. In addition, they report the potential incorporation of Ni into γ -Al₂O₃ or the beginning of mixed oxide phase formation such as NiAl₂O₄.²²

Both the support and the metal nanoparticles can change upon (hydro)thermal ageing at the same time. Hence, the challenge is to deconvolute the many different deactivation pathways which may occur, *e.g.*, sintering, oxidation, composite formation. *Post mortem* or *ex situ* analysis can hardly reveal the stage at which deactivation happens. This requires *in situ* or *operando* measurements sensitive to the aforementioned sample changes.^{23–25} It should be noted that there are mainly two different mechanisms proposed for CO₂ methanation,^{7,8,10} where aside from Ni metal as the active species, the support might also be involved in the reaction by adsorption of CO₂ as part of an associative mechanism. The behavior of both the active metal and the support may therefore

be important. In the current study, we present a systematic examination of deactivation effects of Ni/ZrO₂ methanation catalysts by *operando* synchrotron-based powder X-ray diffraction (SPXRD) during artificial thermal aging. The aim is to deconvolute contributions of support and Ni nanoparticle changes to catalyst deactivation. A designed model series of different Ni/ZrO₂ catalysts was prepared using the same ZrO₂ support, which was calcined at various temperatures prior to Ni deposition. Thus, distinct thermal stability and textural properties of the supports can be achieved, while similar Ni particles are initially present on each support, as illustrated in Fig. 1. A Ni/ZrO₂ system with the monoclinic ZrO₂ polymorph was chosen as a typically employed combination for methanation catalysts^{6,7} and because ZrO₂ exhibits a more defined crystalline structure compared to more commonly applied γ -Al₂O₃ supports. Furthermore, monoclinic ZrO₂ is typically a non-reducible support and mixed oxide formation with Ni is less likely compared to γ -Al₂O₃. ZrO₂ is therefore an ideal model support system to study thermally induced catalyst changes, as the expected changes are mainly related to textural properties. *Operando* SPXRD measurements were used to reveal structural and textural changes of the support and Ni nanoparticles based on crystallite size analysis at different deactivation temperatures during CO₂ methanation. In this way, the distinct contributions of active metal or support changes on catalyst deactivation can be deconvoluted. This allows to derive structure–deactivation relationships for the Ni/ZrO₂ catalyst series. Therefore, this study presents an approach to distinguish changes occurring in different components of the solid catalyst during artificial thermal deactivation cycles, using a single method, *i.e.*, *in situ/operando* SPXRD. The catalysts used in this study may be considered as model systems for the purpose of monitoring deactivation, without optimization regarding their catalytic performance.

2 Experimental

2.1 Catalyst synthesis

Ni/ZrO₂ catalysts were prepared using the homogeneous deposition–precipitation method adapted from Mutz *et al.*²⁶ As support material commercial ZrO₂ pellets (Saint-Gobain) were crushed and thermally aged at different temperatures (indicated in abbreviation) for 12 h (10 °C min^{−1}) resulting in samples Z350, Z500, Z650, Z800 and Z1000. The calcined powders were finely granulated to a mesh size ≤100 µm and 1.6 g of each powder was suspended in an aqueous solution (230 mL) of 2.002 g Ni(NO₃)₂·6H₂O (Merck 99%) with 2.906 g of urea (7 eq., Carl Roth, crystalline, 99.6%), which corresponds to an target Ni loading of 20 wt%. The suspension was stirred for 1 h at room temperature (pH = 4) and afterwards heated to 90 °C. The mixture was heated for 18 h under reflux, cooled to room temperature, then stirred for another 1 h (pH = 8). The solid was filtered off, washed three times with 100 mL of deionized water and dried overnight at 90 °C. Subsequently, the catalysts were calcined at 500 °C for 4 h (10 °C min^{−1}).

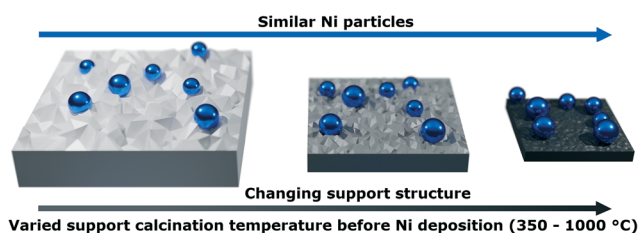


Fig. 1 Illustration of the initially prepared Ni/ZrO₂ catalysts applied in artificial thermal aging experiments. Before Ni deposition the ZrO₂ support was calcined at different temperatures to alter the support textural properties (surface area, pore volume) and to vary the thermal stability of the support. Ni blue, ZrO₂ grey, the grey color gradient illustrates changing textural properties of the support.



2.2 Laboratory catalyst characterization

Catalyst composition was determined by inductively coupled plasma optical emission spectroscopy (ICP-OES) with a Perkin Elmer Optima 8000 instrument equipped a Scott/crossflow sample injection system.

Specific surface area (A_{BET}) and pore volume (V_{p}) were obtained from N_2 physisorption using a BELSORP-mini II system (Microtrac) with the method of Brunauer, Emmett and Teller (BET).²⁷ Prior to the measurement the samples were degassed at 300 °C for 2 h under vacuum.

Mercury intrusion porosimetry (MIP) analysis was conducted using a ThermoScientific PASCAL 140 (lower pressures) and a ThermoScientific PASCAL 440 (higher pressures). The pore width (w_{p}) distribution was calculated using the Washburn equation (eqn (1)).²⁸ The used contact angle (θ) was 140° with a surface tension (σ) of 0.48 N m⁻¹. Here p_{L} and p_{G} are the pressure of liquid and gas, respectively, and d_{p} is the pore diameter.

$$p_{\text{L}} - p_{\text{G}} = -\frac{4\sigma \cos \theta}{d_{\text{p}}} \quad (1)$$

Crystalline phases and crystallite sizes (L_{vol}) of the catalysts were studied with laboratory powder X-ray diffraction (PXRD) measurements using a PANalytical X'Pert PRO diffractometer with Ni-filtered Cu K α radiation ($\lambda = 1.54060$ Å) for $2\theta = 5$ –80° (0.625° steps with 1.6 s per step). Analysis of the laboratory PXRD was performed similarly to SPXRD experiments (see section 2.5 and ESI†, chapter 4).

Transmission electron microscopy (TEM) and scanning transmission electron microscopy (STEM) images were obtained for three spent catalysts after catalytic testing on a Tecnai F20 (Philips) electron microscope with 200 kV acceleration. A copper grid was used as specimen holder and element distribution was studied energy dispersive X-ray analysis (EDX) using an EDAX s-UTW detector.

2.3 Catalyst activity testing

The catalytic performance of the catalysts was studied in a continuous flow setup with a stainless steel tubular fixed-bed reactor (length = 410 mm, inner diameter $d_{\text{i}} = 7$ mm, outer diameter $d_{\text{o}} = 9.5$ mm) (Fig. S14, ESI†), described in previous work.²⁶ The catalyst bed (length = 30 to 35 mm) contained 300 mg of catalyst (sieve fraction 300–450 μm) diluted in 1.15 g of SiC (210 μm , Carborundum, VWR Chemicals). The spare void of the reactor was densely packed with SiC (500 μm) and the different layers were separated with quartz glass wool to ensure a defined gas flow. Thermocouples (K-type) were placed in front and behind the catalyst bed for automated temperature control, using the average temperature of the front and end positions. Gases were dosed using mass flow controllers (Bronkhorst) calibrated using a gas flow meter (Bios DryCal Definer 220, MesaLabs). Gaseous product analysis was performed by gas chromatography (GC) using an Agilent 490 micro gas chromatograph (μ -GC) (channel 1: 10 m PoraPLOT Q, 0.25 mm diameter, carrier gas He; channel 2: 10

m mole sieve column with 5 Å, 0.25 mm diameter, carrier gas Ar). Prior to each reaction, the catalysts were activated in the reactor in a 50% H_2/N_2 mixture (both purity 99.9999% from Air Liquide) (600 mL min⁻¹) at 500 °C for 2 h. After cooling to reaction temperature, the gas feed was changed to 10% $\text{CO}_2/40\%$ H_2/N_2 (CO_2 purity 99.995% from Air Liquide) resulting in a weight hourly space velocity (WHSV) of 60 L g_{cat}⁻¹ h⁻¹. The catalytic performance was tested at three different temperatures (250 °C, 350 °C and 400 °C) and a hold time of 1 h each to obtain stable catalytic activity. Conversion of CO_2 (X_{CO_2}) was calculated similarly to earlier measurements by Mutz *et al.* using N_2 as internal standard (see ESI†).²⁶

2.4 Operando synchrotron powder X-ray diffraction (SPXRD) experiments

SPXRD measurements were performed at the X04SA-MS beamline of the Swiss Light Source (Paul Scherrer Institute, Villigen, Switzerland).²⁹ An X-ray energy of 30.5 keV ($\lambda = 0.406739$ Å) was used to ensure high transmission of the samples, which was required due to the strongly absorbing ZrO_2 support. 2D diffraction patterns were acquired with a Pilatus 6M detector, which were azimuthally integrated using the Dioptas software.³⁰

Operando SPXRD studies were carried out using a quartz capillary (WJM-Glas Müller GmbH) microreactor setup of 1.0 mm diameter.³¹ About 4 mg of the undiluted calcined catalyst (sieve fraction 100–200 μm) resulting in a bed length of ≈ 4 mm was used for each experiment. The reactor was heated using a hot-air blower (LE MINI SENSOR KIT, Leister Technologies), gases were dosed using mass flow controllers (Bronkhorst) and reactor outlet gases were analyzed by on-line mass spectrometry (MS) with an OMNI-Star GSD 320 (Pfeiffer Vacuum). Before the experiments, the temperature inside an empty capillary was calibrated using a type-K-thermocouple (for calibration data see Fig. S16, ESI†). It should be noted that an online temperature control was used during the whole experiment, but the recording of the temperature values failed due to an error in the control software. A LaB₆ NIST 640b standard was measured inside of a 1.0 mm quartz capillary as a calibrant for 2θ scale and to retrieve an instrumental profile function.

The initial calcined catalyst was first activated in a total flow of 10 mL min⁻¹ of 25% H_2/He at 500 °C for 2 h with a heating rate of 10 °C min⁻¹. Afterwards, it was cooled to 400 °C and switched to reaction conditions of 5% $\text{CO}_2/20\%$ H_2/He to determine the initial activity of the catalyst sample at reaction step R1 for 30 min. Subsequently, the catalysts underwent three artificial thermal deactivation cycles with 5 °C min⁻¹ ramps and a holding time of 15 min at temperatures of 500 °C, 575 °C and 625 °C, still under reaction conditions. In between each cycle, changes in catalyst activity were studied at 400 °C labelled as reaction steps R2, R3 and R4. Afterwards, the sample was cooled to room temperature with 10 °C min⁻¹. Diffraction patterns were measured with 9 s acquisition time during activation and



final cool-down as well as 3 s during reaction conditions and artificial thermal deactivation cycles.

The on-line MS data was used to estimate the conversion of CO₂ (X_{CO_2}) and H₂ (X_{H_2}) during *operando* SPXRD. X_i for the respective m/z ratio and thus component i is calculated according to eqn (2):

$$X_i = \left(1 - \frac{I_{i,\text{norm}}}{I_{i,\text{norm,baseline}}}\right) \times 100\% \quad (2)$$

with the normalized ion current ($I_{i,\text{norm}}$) as the ion current of trace i normalized to the He trace. The $I_{i,\text{norm,baseline}}$ is retrieved after cooling down to room temperature at the end of each experiment under reaction conditions, see ESI† for details. To compare relative changes of the conversion during the cycles, X_i is normalized by the initial $X_{i,\text{R1}}$ at R1:

$$X_{i,\text{rel}} = \frac{X_i}{X_{i,\text{R1}}} \quad (3)$$

2.5 Analysis of SPXRD data

Rietveld refinements^{32,33} of the integrated *operando* SPXRD data were performed using Topas (v.6, Bruker AXS).³⁴ The measured LaB₆ NIST 640b standard was used to derive an instrumental profile function, described by a pseudo-Voigt Thompson–Cox Hastings peak shape. A micro-structure analysis was performed using the double-Voigt approach by Balzar as implemented in Topas to describe crystallite size (L_{vol}) and strain (η).³⁵ The monoclinic ZrO₂ phase was described by the structure reported by Smith *et al.*,³⁶ NiO phase by a structure from Rodic *et al.*,³⁷ and the Ni phase with a structure by Rouquette *et al.*³⁸ Sequential Rietveld refinements were performed by adapting templates available from the Topas wiki website.³⁹ More details on the Rietveld refinements are provided in the ESI.†

An average $L_{\text{vol,avg}}$ of the catalyst is defined as following:

$$L_{\text{vol,avg}} = \sum L_{\text{vol},i} \omega_i \quad (4)$$

with $L_{\text{vol},i}$ as the crystallite size of the respective phase i and ω_i as the mass fraction of this phase. Additionally, the relative $L_{\text{vol,rel},i}$ of component i in reference to R1 ($L_{\text{vol},i,\text{R1}}$) and its difference $\Delta L_{\text{vol,rel},i}$ are calculated as following:

$$L_{\text{vol,rel},i} = \frac{L_{\text{vol},i}}{L_{\text{vol},i,\text{R1}}} \quad (5)$$

$$\Delta L_{\text{vol,rel},i} = \left(\frac{L_{\text{vol},i}}{L_{\text{vol},i,\text{R1}}} - 1\right) \times 100\% \quad (6)$$

For spherical particles, L_{vol} can be used with the density (ρ) to calculate the specific surface area (SSA):

$$\text{SSA} = \frac{6000}{L_{\text{vol}} \cdot \rho} \quad (7)$$

However, in practical cases and upon phase changes ρ also changes, which cannot be determined during *in situ* or *operando* SPXRD experiments. Therefore, SSA is approximated by fitting eqn (8) with the fitting parameters a and b to the measured A_{BET} and respective $L_{\text{vol,avg}}$ in the initial state of the catalysts, *i.e.*, taking into account the presence of ZrO₂ and NiO phases.

$$\text{SSA} = \frac{a}{L_{\text{vol,avg}}} + b \quad (8)$$

3 Results and discussion

3.1 Synthesis of catalysts with varying support texture

A series of five ZrO₂ supports was synthesized with different calcination temperatures ranging from 350 °C to 1000 °C (supports Z350 to Z1000, Table S1, ESI†). Following Ni deposition, the resulting Ni/ZrO₂ catalysts were named accordingly as NZ350 to NZ1000 (Table 1). The aim was to prepare a catalyst series with distinct textural properties of the support (*e.g.*, A_{BET} , V_p) dependent only on one parameter – variation in calcination temperature (Fig. 1). In addition, the supported Ni particles were intended to exhibit a similar initial size distribution across all catalysts.

The synthesized catalysts were first characterized by various methods to study the structural and textural properties as summarized in Fig. 2 and Table 1. The support textural properties were studied by N₂ physisorption to gain information about the surface area (A_{BET}) and the pore volume (V_p) (Table S1, ESI†). A steady decrease in A_{BET} from 93 m² g^{−1} for the support calcined at 350 °C to 5 m² g^{−1} for the support calcined at 1000 °C was observed (Fig. 2a, light grey). The V_p changed analogously from 0.23 cm³ g^{−1} to 0.009 cm³ g^{−1} with increasing calcination temperature (Fig. 2d, light grey). The as-prepared Ni/ZrO₂ catalysts exhibited similar tendencies as the pure supports. A decrease in A_{BET} from 66 m² g^{−1} to 18 m² g^{−1} (Fig. 2a, dark grey) and in V_p from 0.23 cm³ g^{−1} to 0.056 cm³ g^{−1} (Fig. 2d, dark grey) were observed with increasing calcination temperature of the initial support (Table 1). An increased density and therefore

Table 1 Overview of the characterization results for the different as-prepared calcined Ni/ZrO₂ catalysts. Additionally, $L_{\text{vol,Ni}}$, D and TOF for the spent catalysts after testing in CO₂ methanation at 400 °C is reported

Catalysts	$\omega_{\text{Ni}}/\text{wt}\%$	$A_{\text{BET}}/\text{m}^2 \text{ g}^{-1}$	$V_p/\text{cm}^3 \text{ g}^{-1}$	$L_{\text{vol,ZrO}_2}/\text{nm}$	$L_{\text{vol,NiO}}/\text{nm}$	$L_{\text{vol,Ni}}/\text{nm}$	$D/\%$	TOF/s ^{−1}
NZ350	14.1	66	0.23	10.4	10.7	34	11.0	2.3×10^{-2}
NZ500	13.2	53	0.14	13.6	9.6	32	11.2	1.8×10^{-2}
NZ650	14.7	33	0.070	23	10.3	26	12.9	1.2×10^{-2}
NZ800	15.2	26	0.055	45	8.1	31	11.5	7.7×10^{-3}
NZ1000	14.7	18	0.056	95	7.9	29	11.8	2.6×10^{-3}



decrease in volume of the ZrO_2 powders with higher support calcination temperatures was visually apparent following synthesis (Fig. 2i), which correlates with the N_2 physisorption results. In general, the A_{BET} values increased following Ni deposition, excluding for the NZ350 catalyst. The relatively large decrease in A_{BET} between Z350 ($93 \text{ m}^2 \text{ g}^{-1}$) and NZ350 ($66 \text{ m}^2 \text{ g}^{-1}$) can be explained by the subsequent calcination of all catalysts at 500°C after Ni deposition. For all catalysts except NZ350, 500°C was an equal or lower temperature treatment than during the initial support calcination step prior to Ni deposition, and would therefore not be expected to induce significant changes in the support. Notably for the NZ350 catalyst A_{BET} ($66 \text{ m}^2 \text{ g}^{-1}$) and V_p ($0.23 \text{ cm}^3 \text{ g}^{-1}$) remained higher than for NZ500 ($52 \text{ m}^2 \text{ g}^{-1}$, $0.14 \text{ cm}^3 \text{ g}^{-1}$). At higher support calcination temperatures (650 – 1000°C) the catalysts exhibited

larger A_{BET} and V_p than the pure support, which can be explained by additional contribution of the nickel to the surface area and pore volume. The A_{BET} of the catalysts is comparable to other Ni/ ZrO_2 catalysts in literature. Everett *et al.* reported $66 \text{ m}^2 \text{ g}^{-1}$ (calcined at 500°C) for a 5 wt% Ni/ ZrO_2 catalyst with commercial m- ZrO_2 from Saint-Gobain.⁴⁰ Martinez *et al.* presented surface areas of 53 and $19 \text{ m}^2 \text{ g}^{-1}$ for 20 wt% Ni/ ZrO_2 catalysts (calcined at 500°C) with mixed ZrO_2 phases (m- ZrO_2 and t- ZrO_2).⁴¹ Pandey *et al.* reported Ni and Ni-Fe supported on m- ZrO_2 (calcined at 500°C) with surface areas between 34 and $23 \text{ m}^2 \text{ g}^{-1}$.⁴²

The pore structure of the as-prepared catalysts was further studied by mercury intrusion porosimetry (MIP) (Fig. S1–S5†). The w_p distribution for NZ500 and NZ1000 are depicted in Fig. 2b and c. A shift to larger mean w_p values (from 17 nm

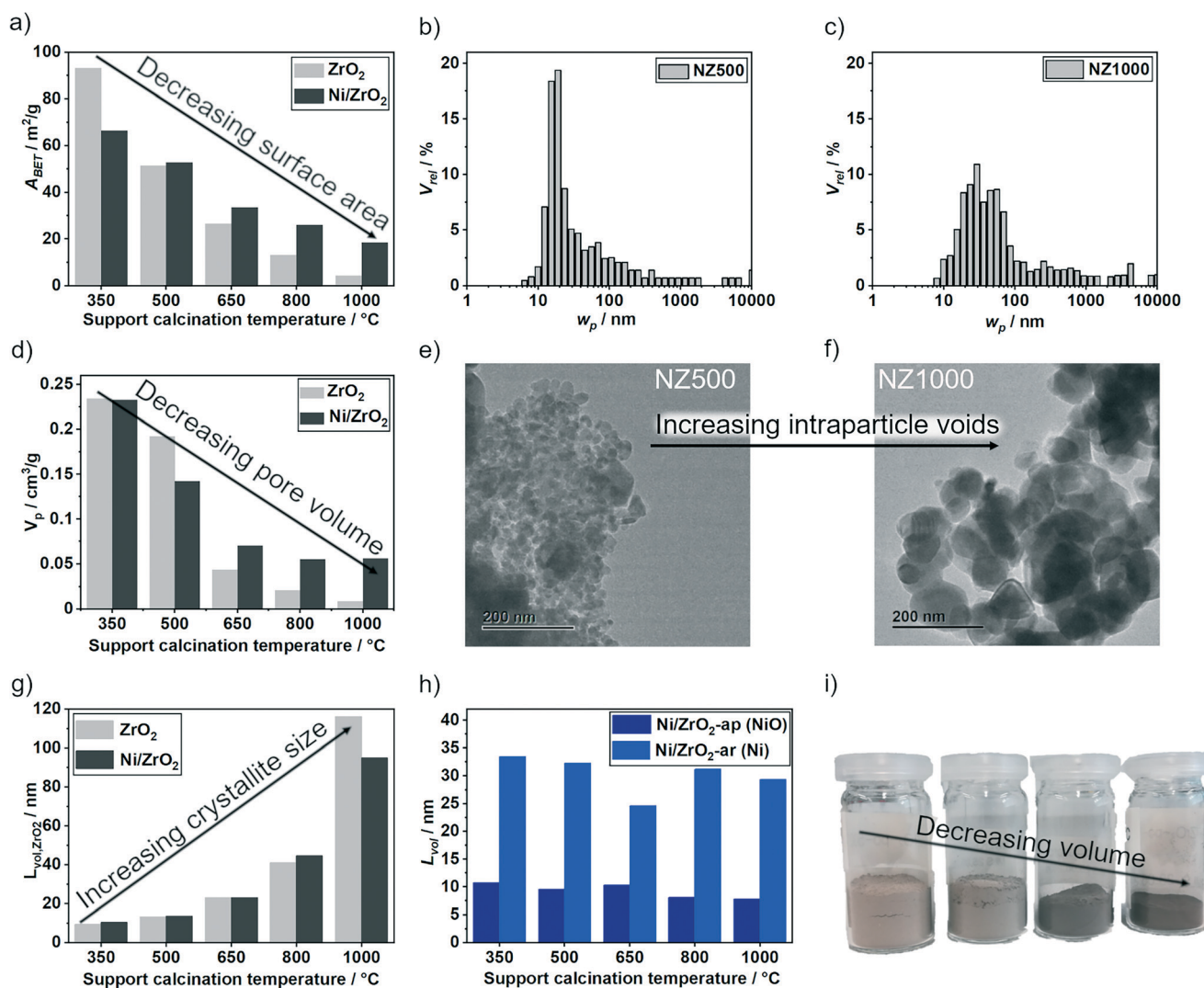


Fig. 2 (a) A_{BET} obtained by N_2 physisorption of the pure calcined supports (light grey) and as-prepared calcined catalysts (dark grey). (b and c) Pore width (w_p) distribution obtained by MIP for as-prepared calcined NZ500 and NZ1000. (d) V_p obtained by N_2 physisorption of the pure calcined supports (light grey) and as-prepared calcined catalysts (dark grey). (e and f) TEM images of the catalysts NZ500 and NZ1000 after CO_2 methanation. (g) ZrO_2 crystallite sizes ($L_{\text{vol,ZrO}_2}$) of the pure calcined supports (light grey) and the as-prepared calcined catalysts (dark grey). (h) Crystallite size L_{vol} for NiO of the as-prepared catalysts (Ni/ ZrO_2 -ap) and Ni of the spent catalysts after CO_2 methanation (Ni/ ZrO_2 -ar). (i) An equal mass of the as-prepared catalysts NZ500, NZ650, NZ800 and NZ1000 showing volume loss and colour change.



to 43 nm) was observed with rising calcination temperature. In addition, TEM analysis showed growth and agglomeration of the support particles during the aging process, leading to the formation of intraparticle voids (Fig. 2e and f). For the NZ1000 catalyst, the support structure mainly consists of large agglomerated particles with no visible internal porosity. Compared to NZ1000, the NZ500 catalyst has smaller particles, which seem to be packed more densely. Hence, an increase of intraparticle voids with rising support calcination temperature can be observed combined with particle growth, explaining the w_p changes from MIP. This further substantiates the N_2 physisorption results which showed a strong decrease of surface area, and the visually apparent increase in density of supports calcined at higher temperature.

The trend of increasing support particle sizes can further be confirmed with laboratory powder X-ray diffraction (PXRD). The PXRD for the calcined supports, as-prepared and spent catalysts are shown in Fig. S6–S9, ESI†. For the support almost all reflections can be assigned to m-ZrO₂ with small contributions of t-ZrO₂ for the supports Z25 and Z350 (Fig. S6 and S7, ESI†). The t-ZrO₂ phase is not detected in NZ350, which might be related to the calcination after Ni deposition. By using Rietveld refinement, the crystallite sizes (L_{vol}) of ZrO₂ for the pure support and the catalysts were determined (Tables S2–S7, ESI†). The zirconia crystallite size (L_{vol,ZrO_2}) increased for the pure support together with temperature from ≈ 10 nm (Z350) to ≈ 116 nm (Z1000) (Fig. 2g, light grey, Table S2, ESI†). For the catalysts (dark grey) a similar trend with increasing L_{vol,ZrO_2} from ≈ 10 nm (NZ350) to ≈ 95 nm (NZ1000) was observed (Fig. 2g, Table 1). The difference in L_{vol,ZrO_2} for Z1000 (116 nm) and NZ1000 (95 nm) could be due to support changes during the synthesis and/or calcination after Ni deposition. In summary, the textural properties A_{BET} and V_p decreased, whereas L_{vol,ZrO_2} increased with rising support calcination temperature. Therefore, the temperature can be used as a single parameter to adjust the textural properties of the initial supports prior to Ni deposition.

Despite the textural differences of the supports, it was aimed to have similar Ni loading and particle size across the catalyst series. Elemental analysis showed that all catalysts have a comparable Ni loading of ≈ 13 –15 wt% (Table 1). The crystallite sizes of NiO (as-prepared catalysts) and Ni (spent catalysts) were investigated using PXRD. A comparable $L_{vol,NiO}$ for all as-prepared catalysts of ≈ 10 nm was determined (Fig. 2h, light blue). After the reaction, $L_{vol,Ni}$ was ≈ 32 nm for all catalysts except NZ650 (26 nm) (Fig. 2h, dark blue). The crystallite growth from 10 to 32 nm could be due to agglomeration during reduction and/or reaction. The size of Ni crystallites is comparable to other Ni/ZrO₂ catalysts reported in literature. Pandey *et al.*⁴² reported $L_{vol,Ni}$ of 58 nm for a 10 wt% Ni/ZrO₂ catalyst and Ren *et al.*⁴³ 34 nm for a 30 wt% Ni/ZrO₂ catalyst.

For a 17 wt% Ni/Al₂O₃ catalyst, which was synthesized with the same method used in this study, $L_{vol,Ni}$ of 5.7 nm was reported.^{17,26} However the different support material (Al₂O₃) limits comparability to this study, while tuning the Ni

particle size towards lower values was not the focus of the present work. A common technique to obtain particle size distributions is TEM/STEM. However, the obtained contrast between ZrO₂ and Ni was too low to distinguish these from each other, which did not allow for reasonable particle size analysis using TEM/STEM. Therefore, EDX mapping was applied for specific areas of selected catalysts (Fig. S10–S13, ESI†). Both large (≥ 100 nm) and small Ni particles (10–30 nm) were found for all catalysts by EDX. With the large Ni particles, the number of obtained images was not sufficient to retrieve statistically meaningful particle size distributions. Therefore, solely PXRD is used to quantitatively estimate the particle/crystallite size based on structural refinement. In conclusion, a catalyst series (NZ350–1000) with similar Ni centers was obtained based on ZrO₂ supports with varying textural properties. This series allows to further investigate the activity and stability of the catalysts depending on the support properties.

3.2 Catalyst testing results

Catalytic activity for CO₂ methanation was monitored at three reaction temperatures (T_r : 250 °C, 350 °C, 400 °C) (Fig. 3a). At 250 °C between 1 to 8% X_{CO_2} was reached, which is far

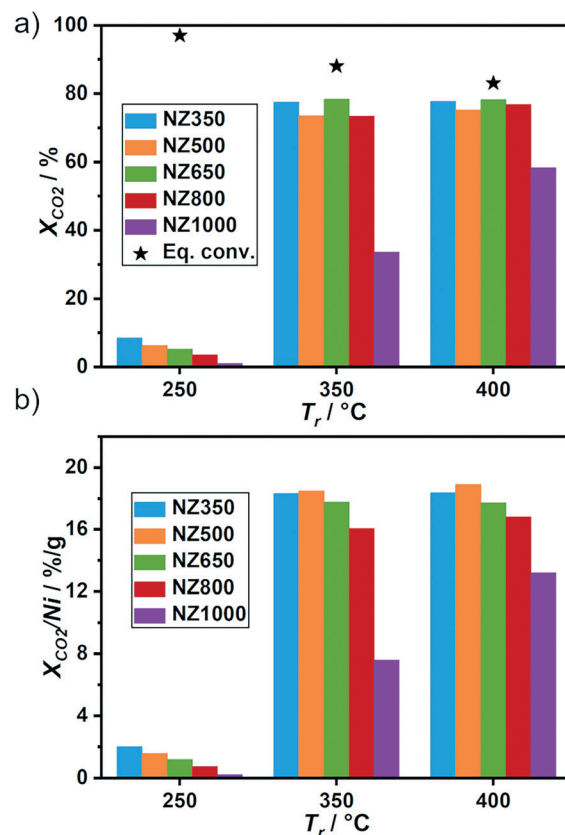


Fig. 3 Catalytic activity of the Ni/ZrO₂ series in CO₂ methanation at 250 °C, 350 °C and 400 °C. (a) CO₂ conversion (X_{CO_2}) with equilibrium conversion (black stars). (b) X_{CO_2} normalized to the Ni amount in the reactor.



below the equilibrium conversion of 97% and therefore suitable for probing catalyst stability.⁴⁴ A trend of decreasing X_{CO_2} with increasing calcination temperature of the support could be observed during reaction at 250 °C. At higher temperatures (350 °C, 400 °C) no specific trend was observed, except for the strong decrease in activity for NZ1000. At 350 °C conversions between 73% and 78% were obtained for NZ350 to NZ800 and 33% for NZ1000. At 400 °C the NZ1000 catalyst showed an increase in activity with 58% X_{CO_2} , whereas the rest of the series achieved similar X_{CO_2} as for 350 °C ($\approx 78\%$), close to but significantly below equilibrium conversion (Fig. 3a, black stars). For better comparison of the activity data, the conversion was normalized to the Ni amount of the tested catalyst (Fig. 3b). With the normalization a similar trend to the one observed at 250 °C can be found also at higher temperatures for the catalysts NZ800 and NZ1000, which exhibited decreased activity compared to NZ350, NZ500 and NZ650.

All catalysts exhibited 100% selectivity towards CH_4 at 250 °C (Fig. S15, ESI†). With increase of T_r , the formation of CO slightly increases. CO formation is probably due to the reverse water gas shift (RWGS) reaction, which is endothermic and therefore more favored at higher temperatures.⁴⁴ The catalysts showed almost no differences in CH_4 selectivity depending on the support calcination temperature except for NZ1000. At 350 °C CH_4 selectivity decreased from 99% (NZ350) to 86% (NZ1000), while at 400 °C a decrease from 97% (NZ350) to 84% (NZ1000) was found. The decline in S_{CH_4} is accompanied with a rise in S_{CO} from 1% (NZ350) to 14% (NZ1000) at 350 °C and from 3% (NZ350) to 16% (NZ1000) at 400 °C. A similar phenomena was observed by Beierlein *et al.* by applying rapid aging tests on Ni/ Al_2O_3 catalysts.¹⁸ They could observe an increase in S_{CO} for samples aged at higher temperatures. Zhao *et al.*⁴⁵ used different combustion methods to synthesize 15 wt% Ni/ ZrO_2 catalysts and tested them in CO_2 methanation. They obtained 1–15% X_{CO_2} at 250 °C, 40–80% at 350 °C and 60–82% at 400 °C. The selectivity towards CH_4 was 100% at 250 °C and at higher temperatures a slight decrease (93%) for some catalysts was observed.

For better comparison with literature the turnover frequency (TOF) was calculated (see details in ESI†) for T_r of 250 °C with X_{CO_2} far below the thermodynamic equilibrium (Table 1). The dispersion (D) was calculated using the $L_{\text{vol,Ni}}$ obtained from Rietveld refinements of the spent samples. For all catalysts a similar D ranging from 11 to 12% was determined (Table 1). The calculated TOF values ranged between $2.3 \times 10^{-2} \text{ s}^{-1}$ for NZ350 and $2.6 \times 10^{-3} \text{ s}^{-1}$ for NZ1000 (Table 1). Tan *et al.*⁴⁶ reported a comparable TOF of $3.8 \times 10^{-3} \text{ s}^{-1}$ at 200 °C for a 12 wt% Ni/ ZrO_2 catalyst with similar X_{CO_2} as at 250 °C reported here. The value is similar to NZ1000 at 250 °C with $2.6 \times 10^{-3} \text{ s}^{-1}$, which showed the lowest activity in this series compared to NZ350 with a nearly one order of magnitude higher TOF of $2.3 \times 10^{-2} \text{ s}^{-1}$. Jia *et al.*⁴⁷ obtained a TOF value of $4.0 \times 10^{-2} \text{ s}^{-1}$ for a 10 wt% Ni/ ZrO_2 catalyst at 235 °C which is similar to NZ350. At 200

°C the group of Liu *et al.*⁴⁸ found a TOF of $4.9 \times 10^{-3} \text{ s}^{-1}$ with a similar X_{CO_2} as for 250 °C in this study. In conclusion, the activity of the catalysts can be influenced by the support calcination temperature, while showing comparable performance to literature values. Notably, the loading, dispersion, and crystallite size of nickel was similar for all catalysts. A decrease in X_{CO_2} and TOF with increased aging temperature can be observed, especially at T_r of 250 °C. Following activity tests, the stability of the catalyst during thermal aging experiments was further investigated to distinguish between deactivation caused by changes of the support or Ni particles.

3.3 Operando SPXRD experiments

Operando SPXRD studies were carried out for the Ni/ ZrO_2 catalyst series to investigate various potential deactivation mechanisms. Distinctions in catalyst thermal stability were expected due to the different initial calcination temperatures of the ZrO_2 supports prior to Ni deposition, and the observed differences in catalytic performance depending on the support properties. The experimental setup and applied temperature profile are shown in Fig. 4a and b, respectively. The acquired MS data for the samples are shown in Fig. S17, S19, S21, S23 and S25, ESI†. All catalysts were first activated in 25% H_2/He at 500 °C for 2 h with the SPXRD results shown in Fig. 4d. During activation the NiO reflexes completely vanish and Ni reflexes appear in the same temperature region (≈ 200 to 250 °C) for all five samples, indicating complete reduction to Ni.

The activation was followed by switching to reaction conditions 5% $\text{CO}_2/20\% \text{ H}_2/\text{He}$ at 400 °C for 30 min, which is labelled as the first reaction step R1. R1 is used to determine the initial activity of the catalysts based on the determined X_{CO_2} and X_{H_2} (Fig. S18, S20, S22, S24 and S26, ESI†). The $X_{\text{CO}_2}/\text{Ni}$ is shown in Fig. 4c. The activity differences between the samples show comparable trends to those found during laboratory testing (Fig. 3). R1 was followed by artificial thermal deactivation cycles (Fig. 4b), with temperatures of 500 °C, 575 °C and 625 °C. After each cycle the activity of the catalysts was measured again at 400 °C for 15 min, labelled as reaction steps R2, R3 and R4, respectively. This allows to determine catalyst deactivation depending on three different applied temperatures. During thermal deactivation cycles no changes of the crystalline phase composition were found as shown by the SPXRD in Fig. 4e. Minor changes of the reflex positions (Fig. 4e), can be attributed to temperature induced lattice expansion or contraction, which matches the temperature program applied to the capillary.

3.4 Crystallite size changes during operando SPXRD

In general, textural properties such as surface area are strongly connected to catalyst performance. In particular adsorption of reactants or products takes place on the surface, but the available surface area is also linked to stabilization of metal nanoparticles. The specific surface area



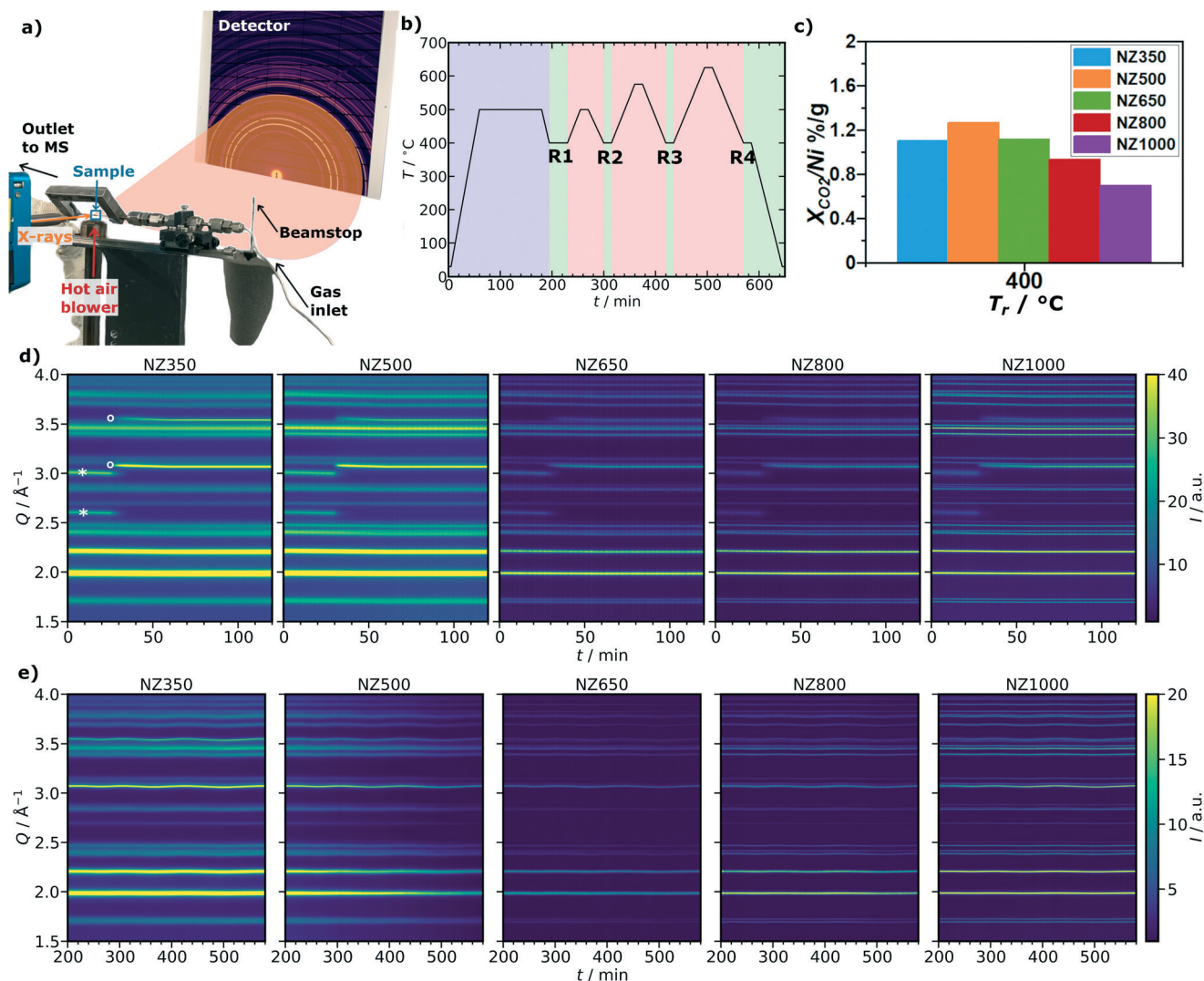


Fig. 4 (a) Scheme of the experimental setup used at the beamline. (b) Overview of the experimental conditions during *operando* SPXRD experiments with activation in 25% H_2/He (blue), reaction conditions 5% $\text{CO}_2/20\% \text{H}_2/\text{He}$ (green) and thermal deactivation cycles under reaction conditions (red) with each reaction condition step labelled as R1–4. (c) CO_2 conversion normalized by the Ni mass inside the reactor for *operando* SPXRD at reaction step R1. (d) 2D overview plots of the diffraction patterns showing the Ni (o) formation from NiO (*) during activation of the five samples as well as (e) under reaction conditions and thermal deactivation cycles, showing no crystalline phase changes.

(SSA) can be calculated from particle sizes as shown in eqn (7), where for crystalline compounds the particle size can be approximated by the crystallite size. However, this relation is only valid for spherical particles. In the present case, two distinct crystalline phases are present, which limits the definition of an overall particle size based on the crystallite sizes as required for eqn (7). Additionally, exact knowledge about ρ is hardly possible during *operando* experiments due to textural changes of the sample, which might lead to density changes. Therefore, in a first step an average crystallite size ($L_{\text{vol,avg}}$) is calculated for the catalysts in the initial calcined state based on eqn (4) taking into account $L_{\text{vol,ZrO}_2}$ and $L_{\text{vol,NiO}}$. Details on the respective refinements are provided in the ESI† This $L_{\text{vol,avg}}$ is then correlated to A_{BET} (Fig. 5a), which is available for the catalysts in the initial state. This correlation is fitted by eqn (8), which is similar to

eqn (7) without the requirement of knowing the exact value of ρ and approximating it *via* a fitting parameter. The fitting results are shown in Fig. 5a with the theoretical curve for pure monoclinic ZrO_2 (eqn (7), $\rho = 5.86 \text{ g cm}^{-3}$ (ref. 36)) showing a similar trend.

Based on the fitted values it is possible to calculate the SSA of the catalysts during the *operando* SPXRD experiments using $L_{\text{vol,avg}}$ of the ZrO_2 and Ni phases present after activation. However, it should be noted that the values are an approximation as the correlation was obtained for the calcined state and not the activated state. Nevertheless, as all catalysts show a very similar phase composition and no crystalline phase changes occur during reaction conditions and thermal deactivation cycles, SSA can be approximated in this way to unravel textural changes of the catalysts. The same correlation was also studied for the pure supports as



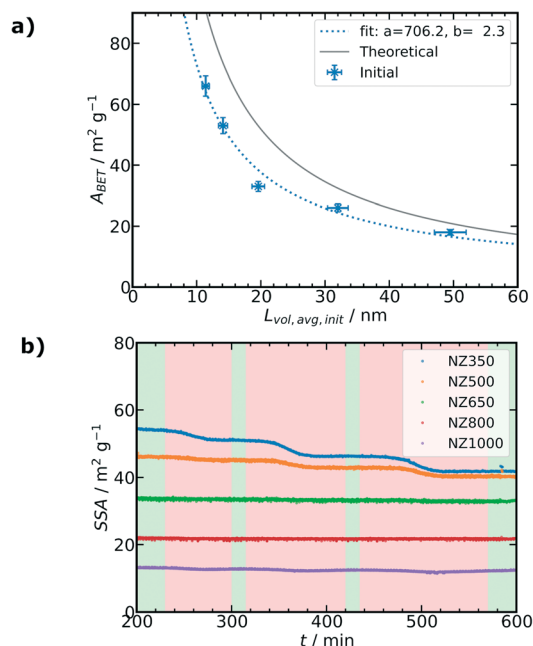


Fig. 5 (a) Fitted correlation (blue dotted line) of the measured A_{BET} to $L_{\text{vol,avg}}$ (blue crosses, mass fraction weighted average of $L_{\text{vol,ZrO}_2}$ and $L_{\text{vol,NiO}}$) initially at the start of the *operando* SPXRD experiments based on eqn (8) with fitting variables a and b , error bars estimated as 5% standard deviation. Theoretical relation of A_{BET} and L_{vol} for spherical particles based on eqn (7) (grey line). (b) Changes of SSA during reaction conditions and thermal deactivation cycles of the *operando* SPXRD experiments as calculated from $L_{\text{vol,avg}}$ (mass fraction weighted average of $L_{\text{vol,ZrO}_2}$ and $L_{\text{vol,Ni}}$) and the fit values from (a) according eqn (8) (shaded areas as Fig. 4b).

shown in Fig. S39, ESI†, resulting in similar fit values supporting the correlation approach. The SSA starting from R1 evolves differently for the catalysts as shown in Fig. 5b. Similarly as for A_{BET} analysis (Fig. 2a), it can be observed that the initial SSA of the catalysts (at $t = 200$ min) decreased with increasing calcination temperature of the ZrO_2 support before Ni deposition. The calculated initial SSA for all catalysts is lower compared to the measured A_{BET} values of the initial calcined samples (Table 1), which can be attributed to the increased crystallite sizes of Ni compared to NiO by a factor of ≈ 3 (for details see Tables S8–S13 ESI†). For NZ350 a pronounced step-wise decrease of the SSA can be observed, with each step related to one of the thermal deactivation cycles. NZ500 exhibits a decrease only significantly with the second and third thermal deactivation step, while NZ650, NZ800 and NZ1000 show no significant reduction of SSA following the thermal deactivation cycles. Besides the study of the overall textural changes based on the calculated SSA, Rietveld refinement together with microstructural analysis allows to investigate crystallite size changes of each crystalline phase individually. Fig. 6a and b show the evolution of the selected (111) reflex for ZrO_2 and Ni phases during *operando* SPXRD, respectively. The different peak shapes of the ZrO_2 (111) reflection (Fig. 6a) illustrate the different

ZrO_2 crystallite sizes depending on the calcination temperature of the support prior to Ni deposition. A representative pattern during each of the reaction steps R1 to R4 is highlighted. For NZ350 and NZ500 significant changes of the reflex depending on the reaction step can be observed, which are more pronounced for NZ350. This indicates crystallite growth of the ZrO_2 phase during thermal deactivation cycles. For NZ650, NZ800 and NZ1000 no significant variation of the reflex can be directly observed. In the case of the Ni (111) reflex (Fig. 6b), all samples show significant changes depending on the reaction step. The variations become more pronounced with higher calcination temperature of the ZrO_2 support. Additionally to the qualitative description of the peak shape, a microstructural refinement as part of Rietveld refinement of the SPXRD provides quantitative information about crystallite size changes during the thermal deactivation cycles. Fig. 6c–e show the evolution of $L_{\text{vol,ZrO}_2}$, $L_{\text{vol,Ni}}$ and the mass fraction weighted average $L_{\text{vol,avg}}$ of both, respectively. Complementary to the absolute values, Fig. 6f–h show the relative changes compared to the initial reaction step R1, respectively, indicating sequential changes during the subsequent thermal aging cycles.

$L_{\text{vol,ZrO}_2}$ depends on the calcination temperature of the support prior to Ni deposition and ranges from about 9 nm to 70 nm in the initial reaction step R1. During thermal deactivation cycles a step-wise increase of $L_{\text{vol,ZrO}_2}$ can be observed for each temperature step for NZ350. Although all catalysts were initially activated for 2 h at 500 °C, for NZ350 a crystallite growth can be observed already at the first thermal deactivation temperature of 500 °C. For NZ500 a significant increase of the crystallite size is only detected with the second and third thermal deactivation steps. The catalysts NZ650, NZ800 and NZ1000 show no significant increase of $L_{\text{vol,ZrO}_2}$ during those cycles. The reversible increase of $L_{\text{vol,ZrO}_2}$ of NZ1000 during the thermal deactivation cycles (Fig. 6c and f) might be explained by larger crystalline domains due to lattice expansion or reversible ordering/disordering, but the differences are small compared to the refinement error and could also be caused by refinement artifacts. The evolution of $L_{\text{vol,Ni}}$ is contrary to the ZrO_2 phase. The initial $L_{\text{vol,Ni}}$ at reaction step R1 range from about 25 to 35 nm for the different catalysts (Fig. 6d). For NZ350, NZ500 and NZ650, the $L_{\text{vol,Ni}}$ only slightly increased with thermal deactivation after the second reaction step R2 (Fig. 6d and g). This is different for the NZ800 and NZ1000 samples where $L_{\text{vol,Ni}}$ already increased with the first thermal deactivation step, while the behavior is most pronounced for NZ1000. The changes can be also observed for the $L_{\text{vol,avg}}$ in Fig. 6e and h. As $L_{\text{vol,avg}}$ represents the mass fraction weighted average for ZrO_2 and Ni it mainly reflects changes of the ZrO_2 phase. Therefore, here the NZ350 and NZ500 samples show the most pronounced changes. On the other hand, it can be observed that the strong changes of $L_{\text{vol,Ni}}$ result in a significant increase of the $L_{\text{vol,avg}}$ for NZ1000. The strong Ni



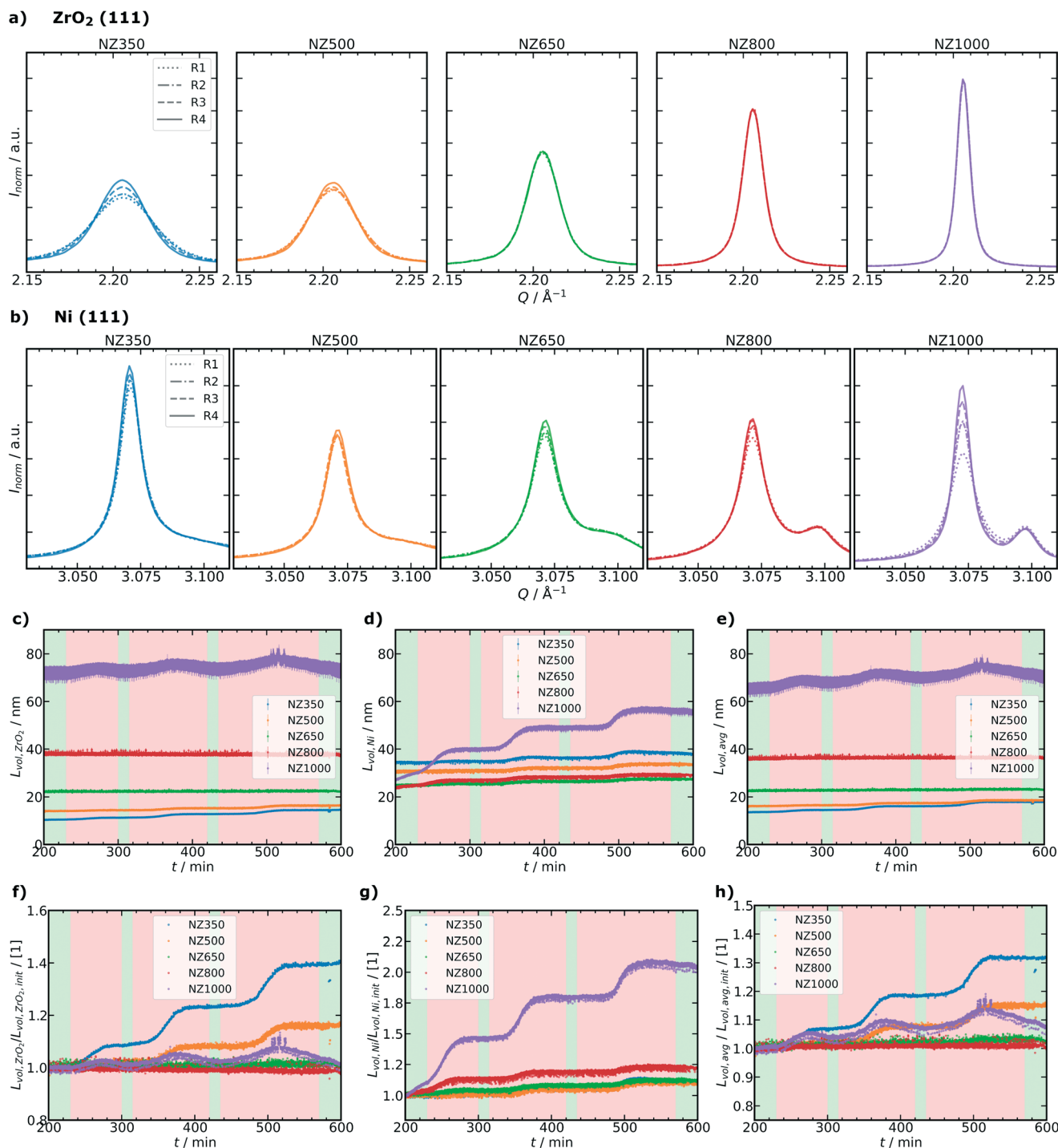


Fig. 6 (a) Changes of the ZrO_2 (111) reflex during thermal deactivation cycles highlighted by R1, R2, R3 and R4 steps, (b) analogous changes of the Ni (111) reflex. Evolution of different crystallite sizes obtained from sequential Rietveld refinements for the different catalysts during *operando* SPXRD experiments: (c) $L_{\text{vol,ZrO}_2}$, (d) $L_{\text{vol,Ni}}$ and (e) $L_{\text{vol,avg}}$ (mass fraction weighted average of $L_{\text{vol,ZrO}_2}$ and $L_{\text{vol,Ni}}$). Relative changes of L_{vol} normalized to the initial reaction step R1 according eqn (5): (f) ZrO_2 phase, (g) Ni phase and (h) average crystallite size of the catalysts. Shaded areas as Fig. 4b.

crystallite size increase in NZ1000 might be explained by the support reaching such a small SSA that Ni particle growth is strongly favored. In contrast, NZ650 and NZ800 were less impacted as a whole by the thermal deactivation procedure.

In summary, the crystallite size changes during thermal deactivation cycles could be quantified and show for NZ350 and NZ500 mainly changes of $L_{\text{vol,ZrO}_2}$, while for NZ800 and NZ1000 especially changes of $L_{\text{vol,Ni}}$ were observed.



3.5 Correlation of crystallite size and thermal deactivation

The conversion of CO₂ and H₂ obtained from MS analysis were used to determine the activity of the catalysts during *operando* SPXRD. The absolute values for X_{CO_2} and X_{H_2} at the different reaction steps R1–4 are shown in Fig. S27 and S28, ESI† respectively. The initial conversions for the catalysts at R1 are in the range of 57 to 67%. Therefore, it can be reasonably assumed that the reaction is performed far from thermodynamic equilibrium conditions or full conversion ($\approx 80\%$, Fig. 3a), which is a prerequisite to detect possible catalyst deactivation. The changes of X_{CO_2} and X_{H_2} in comparison to the initial activity at R1 (ΔX) are shown in Fig. 7a and d, respectively. All studied catalysts show a step-wise deactivation during the thermal deactivation cycles. Therefore, the deactivation can be clearly related to the temperature applied during each of the cycles. Less pronounced deactivation occurs for the NZ650 and NZ800 catalysts, which suffer a loss of $\approx 1\%$ in X_{CO_2} and X_{H_2} after thermal deactivation cycles. The other three catalysts lost more than 2% in X_{CO_2} and X_{H_2} , while there are differences in the extent of activity loss between X_{CO_2} and X_{H_2} .

Besides the activity differences, the changes of $L_{\text{vol,rel,ZrO}_2}$ and $L_{\text{vol,rel,Ni}}$ in reference to the initial reaction step R1 are shown in Fig. 7b and e, respectively. The $L_{\text{vol,rel,ZrO}_2}$ of NZ350 is increased by about 40% and for NZ500 by about 17% compared to the initial values at R1, while the other three catalysts show only minor changes of the $L_{\text{vol,rel,ZrO}_2}$ after thermal deactivation cycles of less than 3%.

for $L_{\text{vol,rel,Ni}}$, where NZ1000 shows an increase of almost 100% followed by the NZ800 sample of $\approx 20\%$. The other three samples show increases of $\approx 10\%$ or below.

To estimate the influence of the different L_{vol} changes on catalyst deactivation, the relative conversions ($X_{\text{CO}_2,\text{rel}}$, $X_{\text{H}_2,\text{rel}}$, eqn (3)) are correlated with the relative crystallite sizes ($L_{\text{vol,rel},i}$ eqn (5)) with respect to the initial reaction step R1 in Fig. 7c and f, respectively. This correlation analysis allows to retrieve information regarding which of the two phases might have a stronger influence on catalyst deactivation. For both evaluated conversions, the strongest observed correlations are for the changes of $L_{\text{vol,Ni}}$ of the NZ1000 catalyst, followed by the correlations of $L_{\text{vol,ZrO}_2}$ for NZ350. In general, it can be observed that there is a greater correlation between $L_{\text{vol,Ni}}$ and the conversions for NZ1000, NZ800 and NZ650. This is different for NZ350 and NZ500, where the correlation of $L_{\text{vol,ZrO}_2}$ with the conversions is larger than $L_{\text{vol,Ni}}$.

3.6 Overall discussion of the thermal deactivation

The five different studied catalysts essentially differ by the calcination temperature and consequently the surface area and pore volume of the ZrO₂ support prior to Ni deposition, while the following calcination and activation steps are identical for all samples. In addition, the initial properties of the active metal Ni particles in terms of loading, dispersion, and crystallite size were similar for all catalysts following synthesis and calcination. As observed in the *operando* SPXRD the activity trends of the catalysts after activation at

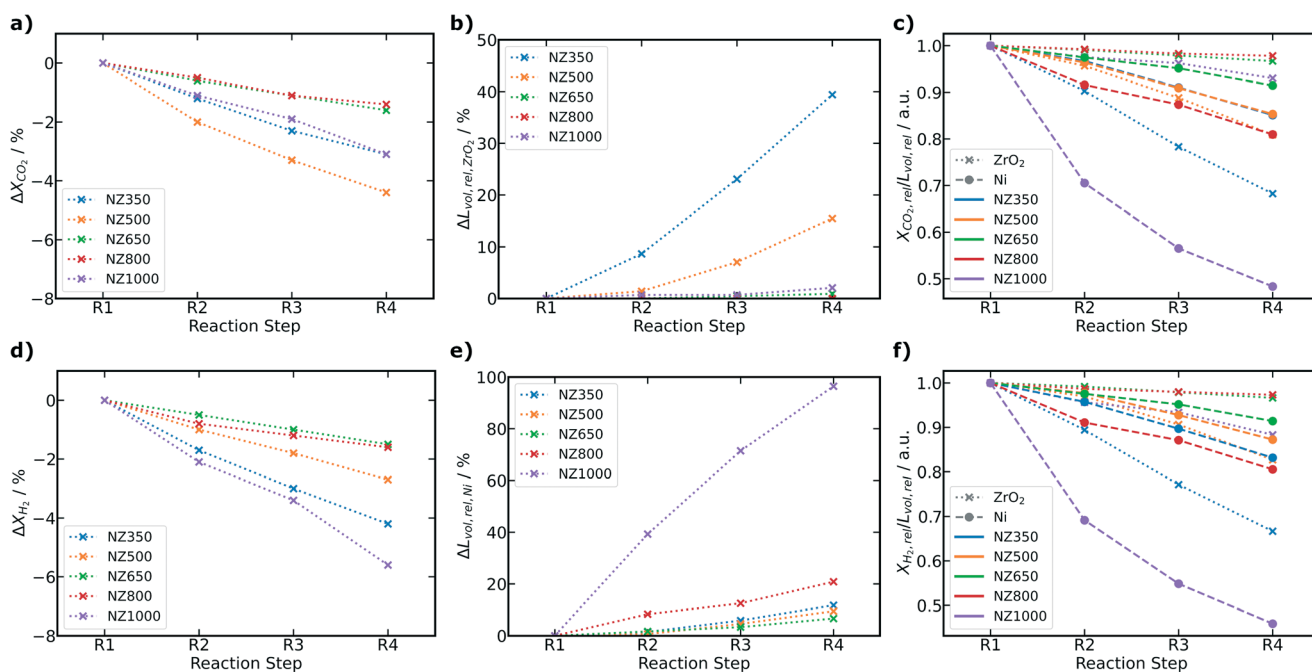


Fig. 7 (a) Changes of the X_{CO_2} for each reaction step as difference to the initial activity at R1. (b) Relative changes $\Delta L_{\text{vol,rel,ZrO}_2}$ of the ZrO₂ crystallite size according eqn (6) depending on the reaction step referenced to R1. (c) Relative CO₂ conversion $X_{\text{CO}_2,\text{rel}}$ (eqn (3)) divided by the relative crystallite size $L_{\text{vol,rel}}$ (eqn (5)) for Ni and ZrO₂ phases. (d) Changes of the X_{H_2} for each reaction step as difference to the initial activity at R1. (e) Relative changes $\Delta L_{\text{vol,rel,Ni}}$ of the Ni crystallite size according eqn (6) depending on the reaction step with R1 as reference. (f) Relative H₂ conversion $X_{\text{H}_2,\text{rel}}$ (eqn (3)) divided by the relative crystallite size $L_{\text{vol,rel}}$ (eqn (5)) for Ni and ZrO₂ phases.



R1 are similar compared to laboratory tests, showing generally decreasing activity with higher calcination temperature of the support, especially for NZ1000. The activated catalysts at R1 are mainly distinguished by their $L_{\text{vol,ZrO}_2}$, while the $L_{\text{vol,Ni}}$ is in a comparable range for all catalysts with minor differences. The $L_{\text{vol,Ni}}$ is relatively large at around 30 nm. Therefore, it can be reasonably assumed that no major differences in the reaction mechanism depending on the Ni particle size as shown by Vogt *et al.*¹⁵ are present, since the type of surface sites present are similar and deactivation as reported by Mutz *et al.*¹⁴ should be negligible. The resulting activity should mainly depend on the available active surface area of the catalysts.

Typically, the active surface area and the specific surface area of catalysts are determined *via* gas sorption methods in *post mortem* analysis.^{49,50} However, applying these methods under *operando* conditions is hardly possible. Contrary to this, PXRD with microstructural refinement as shown here allows to determine textural changes of the ZrO₂ support and Ni nanoparticles as long as there are no complex phase changes occurring during thermal deactivation. In the present case, the five catalysts show different crystallite size changes and deactivation behavior depending on the calcination temperature of the support. The most pronounced catalyst deactivation was observed for NZ350, NZ500 and NZ1000 catalysts (Fig. 7a and d), while for those three samples also the strongest crystallite size changes were found. For NZ350 and NZ500, mainly an increase of $L_{\text{vol,ZrO}_2}$ and thus sintering of the support was detected, while for NZ1000 the $L_{\text{vol,Ni}}$ after thermal aging was much larger and thus Ni particle sintering occurred. Based on the correlation analysis shown in Fig. 7c and f the major reason for catalyst deactivation might be divided into: (i) dominant changes of the ZrO₂ support, (ii) dominant changes of the Ni phase and (iii) an intermediate region with both effects as schematically illustrated in Fig. 8.

Despite that all catalysts were activated at 500 °C for 2 h, NZ350, NZ500 and NZ1000 show a slight deactivation after only 15 min at 500 °C under reaction conditions. The deactivation of NZ350 and NZ500 can be mainly related to textural changes of the ZrO₂ support. For NZ650, NZ800 and NZ1000 sintering of Ni particles appears to be the main reason for deactivation, with NZ650 and NZ800 showing highest stability among the tested samples. The stability of NZ650 and NZ800 might be related to the calcination temperature of the supports during catalyst preparation, which were higher than the subsequent deactivation temperatures. The deactivation due to Ni sintering can be explained by a decrease in the active metal surface area. Literature studies have discussed changes of Ni nanoparticles, the support material and a complex interplay of the effects as reasons for deactivation.^{18–22} However, deconvolution of these effects was not previously possible due to entirely *post mortem* analysis and thus the missing link with activity data provided by the *operando* analysis shown here. Sintering of the ZrO₂ support as shown in the present work might influence the catalyst stability in different ways. For Ni/ZrO₂ the associative mechanism for CO₂ methanation would require CO₂ adsorbates on the ZrO₂ support to take part in the reaction.^{7,8,10} Therefore the amount of CO₂ adsorbed on the support is lower with decreasing support surface area leading to lower catalytic activity. Furthermore, it is possible that the textural changes of the support can result in blocking of active Ni surface sites.

Additionally, SPXRD did not reveal formation of bulk NiO species and the quantitative phase analysis from Rietveld refinement matches the values obtained from elemental analysis. Thus, a fully reduced state of Ni throughout the experiment was assumed. The formation of surface NiO cannot be fully excluded by the applied methods, however due to the relatively large and similar Ni particle sizes in the

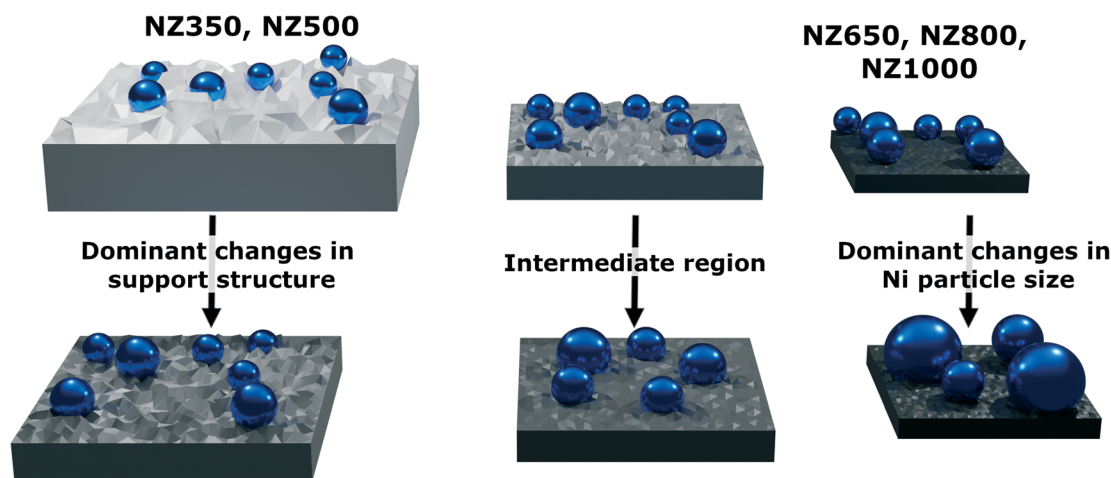


Fig. 8 Summary of the different proposed deactivation mechanism found by *operando* SPXRD. NZ350 and NZ500 show predominant ZrO₂ sintering and NZ650, NZ800 and NZ1000 dominant Ni sintering and an expected intermediate region as origin of deactivation. Ni blue, ZrO₂ grey. The grey color gradient illustrates changing textural properties of the support associated to a decrease in surface area and pore volume due to sintering.



catalyst series, similar redox properties are assumed for all catalysts. Possible deactivation due to formation of surface oxide species, if this occurs, should therefore be similar for all five catalysts due to identical reaction conditions. The significant quantity of H₂ in the feed makes formation of NiO species unlikely in any case, also according to previous studies using *operando* XAS.^{13,14,16,17} In these previous studies, NiO was only significantly present during H₂-dropout tests. However, to unravel the different potential effects in more detail, further kinetic and spectroscopic studies of the active and deactivated catalysts would be required. In the present study, the focus was on presenting SPXRD as a single powerful and relatively routinely available method to differentiate thermal deactivation effects sensitive towards textural and crystallographic changes under *operando* conditions, with the possibility to distinguish support and metal nanoparticle changes.

4 Conclusions

In the present study, a bottom-up designed series of five different Ni/ZrO₂ catalysts was prepared *via* an adapted homogeneous deposition precipitation method. The catalysts are based on a commercial ZrO₂ support material, which was calcined at different temperatures prior to Ni deposition in order to change the textural properties, *i.e.*, specific surface area and pore volume of the support systematically. The catalysts were characterized by conventional laboratory methods confirming a decrease in surface area and pore volume with increasing support calcination temperature. Laboratory catalytic tests for CO₂ methanation showed comparable activity towards literature reported samples with TOF ranging from $2.3 \times 10^{-2} \text{ s}^{-1}$ to $2.6 \times 10^{-3} \text{ s}^{-1}$ at 250 °C. The catalysts exhibit similar Ni crystallite size ($\approx 30 \text{ nm}$), dispersion and Ni loading. As the five catalysts essentially only differ by a single parameter, the support calcination temperature, the series is well suited for thermal deactivation studies.

During *operando* SPXRD studies with artificial thermal deactivation cycles, the catalysts showed different extent of deactivation with NZ350, NZ500 and NZ1000 showing the strongest activity loss and NZ650 and NZ800 the least. No crystalline phase changes were observed during thermal deactivation, but rather crystallite size changes of the ZrO₂ support and Ni nanoparticles. These changes were dependent on the ZrO₂ calcination temperature along with the related textural properties. The five catalysts can be distinguished into samples that mainly deactivate due to changes of the support (NZ350 and NZ500), or changes of the Ni crystallite size (NZ650, NZ800 and NZ1000). The activity loss associated with Ni sintering can be explained by a decrease of the specific metal surface area. For the samples with dominant support sintering, either loss in CO₂ adsorption sites due to decreased support surface area or blocking of accessible metal surface area can be an explanation for the observed deactivation.

The presented approach of using *operando* SPXRD to study catalyst deactivation due to textural changes of the active site (metal nanoparticles) or the support is a valuable tool to understand temperature induced catalyst deactivation. In general, it can be emphasized to not only focus on changes of the metal nanoparticles, but also to consider the role of the support. This approach can be used in general to differentiate influences on catalyst stability under reaction conditions, which are not directly accessible with established methods including gas sorption or electron microscopy. The possibility to follow catalysts changes while they occur is especially important for potential dynamic process operation related to fluctuating renewable energy supply.⁵¹ There, (ir)reversible changes of the catalyst structure are difficult to follow by *ex situ* characterization, but are important to reveal for a fundamental understanding of catalyst behavior.⁵¹ In future, promising developments of 3D spatially-resolved techniques, *e.g.*, *operando* XRD or PDF tomography,^{52,53} may provide additional information about the location of the textural changes with respect to the reactor position (*e.g.*, due to hot-spots) or within specific catalyst particles. Additionally, recent progress in hard X-ray nanotomography methods allows to investigate textural sample changes *in situ* approaching routinely sub 50 nm resolution, which can be sufficient to reveal the processes observed here.⁵⁴

Author contributions

Conceptualization, SW; data curation, MS, SW, TS; formal analysis, MS, SW; funding acquisition, TS; investigation, MS, SW, LK, TS; methodology, MS, SW; project administration, SW, JDG, TS; resources, JDG, TS; supervision, SW, JDG, TS; validation, MS, SW, JDG, TS; visualization, MS, SW; writing—original draft preparation, MS, SW; writing—review and editing, all authors.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Gefördert durch die Deutsche Forschungsgemeinschaft (DFG)-406914011, funded by the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation)-406914011. Dr. Marc-André Serrer is acknowledged for support of the laboratory catalyst testing experiments. We thank Dr. Eduardo Barbosa and Dr. Sumeet Kale (Karlsruhe Institute of Technology) for performing N₂ sorption experiments. Thanks to Heike Rudzik, Ken Luca Abel and Prof. Roger Gläser (Universität Leipzig) for providing ICP-OES analysis. We acknowledge Maximilian Franz and Prof. Dirk Enke (Universität Leipzig) for carrying out MIP experiments. We thank Dr. Florian Maurer for assistance in the graphical preparation of the schemes. This work was partly carried out with the support of the Karlsruhe Nano



Micro Facility (KNMF), a Helmholtz Research Infrastructure at Karlsruhe Institute of Technology (KIT), which provided access to TEM instruments *via* proposal 2019-022-026980. The authors thank Xiaohui Huang, Dr. Reihaneh Pashminehazar and Prof. Christian Kübel (Karlsruhe Institute of Technology) for support with the TEM experiments. The authors acknowledge the Paul Scherrer Institut, Villigen, Switzerland for provision of synchrotron radiation beamtime at beamline X04SA-Materials Science of the Swiss Light Source and thank Dr. Nicola Casati, Michael Lange (Paul Scherrer Institut) and Shweta Sharma (Karlsruhe Institute of Technology) for support during the beamtime. Open access funding enabled by the Read & Publish agreement between Karlsruhe Institute of Technology and the Royal Society of Chemistry.

Notes and references

- 1 C. H. Bartholomew, *Appl. Catal., A*, 2001, **212**, 17–60.
- 2 M. D. Argyle and C. H. Bartholomew, *Catalysts*, 2015, **5**, 145–269.
- 3 J. Bremer and K. Sundmacher, *React. Chem. Eng.*, 2019, **4**, 1019–1037.
- 4 J. Bremer, K. H. G. Rätze and K. Sundmacher, *AIChE J.*, 2017, **63**, 23–31.
- 5 M. Serrer, M. Stehle, M. L. Schulte, H. Besser, W. Pfleging, E. Saraci and J.-D. Grunwaldt, *ChemCatChem*, 2021, **13**, 3010–3020.
- 6 M. A. Aziz, A. A. Jalil, S. Triwahyono and A. Ahmad, *Green Chem.*, 2015, **17**, 2647–2663.
- 7 P. Frontera, A. Macario, M. Ferraro and P. Antonucci, *Catalysts*, 2017, **7**, 59.
- 8 S. Rönsch, J. Schneider, S. Matthischke, M. Schlüter, M. Götz, J. Lefebvre, P. Prabhakaran and S. Bajohr, *Fuel*, 2016, **166**, 276–296.
- 9 M. Younas, L. Loong Kong, M. J. Bashir, H. Nadeem, A. Shehzad and S. Sethupathi, *Energy Fuels*, 2016, **30**, 8815–8831.
- 10 B. Miao, S. S. K. Ma, X. Wang, H. Su and S. H. Chan, *Catal. Sci. Technol.*, 2016, **6**, 4048–4058.
- 11 B. Mutz, P. Sprenger, W. Wang, D. Wang, W. Kleist and J.-D. Grunwaldt, *Appl. Catal., A*, 2018, **556**, 160–171.
- 12 R. P. Struis, T. J. Schildhauer, I. Czekaj, M. Janousch, S. M. Biollaz and C. Ludwig, *Appl. Catal., A*, 2009, **362**, 121–128.
- 13 B. Mutz, H. W. Carvalho, S. Mangold, W. Kleist and J.-D. Grunwaldt, *J. Catal.*, 2015, **327**, 48–53.
- 14 B. Mutz, A. M. Gänzler, M. Nachtegaal, O. Müller, R. Frahm, W. Kleist and J.-D. Grunwaldt, *Catalysts*, 2017, **7**, 279.
- 15 C. Vogt, E. Groeneveld, G. Kamsma, M. Nachtegaal, L. Lu, C. J. Kiely, P. H. Berben, F. Meirer and B. M. Weckhuysen, *Nat. Catal.*, 2018, **1**, 127–134.
- 16 M. A. Serrer, K. F. Kalz, E. Saraci, H. Lichtenberg and J.-D. Grunwaldt, *ChemCatChem*, 2019, **11**, 5018–5021.
- 17 M.-A. Serrer, A. Gaur, J. Jelic, S. Weber, C. A. Fritsch, A. H. Clark, E. Saraci, F. Studt and J.-D. Grunwaldt, *Catal. Sci. Technol.*, 2020, **10**, 7542–7554.
- 18 D. Beierlein, D. Häussermann, Y. Traa and E. Klemm, *Catal. Lett.*, 2021, **1**, 3.
- 19 S. Abelló, C. Berrueto and D. Montané, *Fuel*, 2013, **113**, 598–609.
- 20 F. Koschany, D. Schlereth and O. Hinrichsen, *Appl. Catal., B*, 2016, **181**, 504–516.
- 21 C. Mebrahtu, S. Perathoner, G. Giorgianni, S. Chen, G. Centi, F. Krebs, R. Palkovits and S. Abate, *Catal. Sci. Technol.*, 2019, **9**, 4023–4035.
- 22 S. Ewald, M. Kolbeck, T. Kratky, M. Wolf and O. Hinrichsen, *Appl. Catal., A*, 2019, **570**, 376–386.
- 23 M. A. Bañares, *Catal. Today*, 2005, **100**, 71–77.
- 24 R. Portela, S. Perez-Ferreras, A. Serrano-Lotina and M. A. Bañares, *Front. Chem. Sci. Eng.*, 2018, **12**, 509–536.
- 25 I. L. Buurmans and B. M. Weckhuysen, *Nat. Chem.*, 2012, **4**, 873–886.
- 26 B. Mutz, M. Belimov, W. Wang, P. Sprenger, M. A. Serrer, D. Wang, P. Pfeifer, W. Kleist and J.-D. Grunwaldt, *ACS Catal.*, 2017, **7**, 6802–6814.
- 27 S. Brunauer, P. H. Emmett and E. Teller, *J. Am. Chem. Soc.*, 1938, **60**, 309–319.
- 28 E. W. Washburn, *Phys. Rev.*, 1921, **17**, 273–283.
- 29 P. R. Willmott, D. Meister, S. J. Leake, M. Lange, A. Bergamaschi, M. Böge, M. Calvi, C. Cancellieri, N. Casati, A. Cervellino, Q. Chen, C. David, U. Flehsig, F. Gozzo, B. Henrich, S. Jäggi-Spielmann, B. Jakob, I. Kalichava, P. Karvinen, J. Krempasky, A. Lüdeke, R. Lüscher, S. Maag, C. Quitmann, M. L. Reinle-Schmitt, T. Schmidt, B. Schmitt, A. Streun, I. Vartiainen, M. Vitins, X. Wang and R. Wulschleger, *J. Synchrotron Radiat.*, 2013, **20**, 667–682.
- 30 C. Prescher and V. B. Prakapenka, *High Pressure Res.*, 2015, **35**, 223–230.
- 31 J.-D. Grunwaldt, M. Caravati, S. Hannemann and A. Baiker, *Phys. Chem. Chem. Phys.*, 2004, **6**, 3037–3047.
- 32 H. M. Rietveld, *Acta Crystallogr.*, 1967, **22**, 151–152.
- 33 H. M. Rietveld, *J. Appl. Crystallogr.*, 1969, **2**, 65–71.
- 34 DIFFRAC.SUITE TOPAS – XRD Software Bruker, 2020, <https://www.bruker.com/de/products/x-ray-diffraction-and-elemental-analysis/x-ray-diffraction/xrd-software/topas.html>.
- 35 D. Balzar, *Int. Union Crystallogr. Monogr. Crystallogr.*, 1999, **10**, 94–126.
- 36 D. K. Smith and W. Newkirk, *Acta Crystallogr.*, 1965, **18**, 983–991.
- 37 D. Rodic, V. Spasojevic, V. Kusigerski, R. Tellgren and H. Rundlof, *Phys. Status Solidi B*, 2000, **218**, 527–536.
- 38 J. Rouquette, J. Haines, G. Frayssé, A. Al-Zein, V. Bornand, M. Pintard, P. Papet, S. Hull and F. Gorelli, *Inorg. Chem.*, 2008, **47**, 9898–9904.
- 39 <https://topas.dur.ac.uk/topaswiki/doku.php?id=topas>, accessed on 18/10/2021.
- 40 O. E. Everett, P. C. Zonetti, O. C. Alves, R. R. de Aveliz and L. G. Appel, *Int. J. Hydrogen Energy*, 2020, **45**, 6352–6359.
- 41 J. Martínez, E. Hernández, S. Alfaro, R. López Medina, G. Valverde Aguilar, E. Albiter and M. Valenzuela, *Catalysts*, 2018, **9**, 24.
- 42 D. Pandey and G. Deo, *J. Ind. Eng. Chem.*, 2016, **33**, 99–107.
- 43 J. Ren, X. Qin, J. Z. Yang, Z. F. Qin, H. L. Guo, J. Y. Lin and Z. Li, *Fuel Process. Technol.*, 2015, **137**, 204–211.



- 44 J. Gao, Y. Wang, Y. Ping, D. Hu, G. Xu, F. Gu and F. Su, *RSC Adv.*, 2012, **2**, 2358–2368.
- 45 K. Zhao, W. Wang and Z. Li, *J. CO₂ Util.*, 2016, **16**, 236–244.
- 46 J. Tan, J. Wang, Z. Zhang, Z. Ma, L. Wang and Y. Liu, *Appl. Surf. Sci.*, 2019, **481**, 1538–1548.
- 47 X. Jia, X. Zhang, N. Rui, X. Hu and C. J. Liu, *Appl. Catal., B*, 2019, **244**, 159–169.
- 48 Z. Liu, X. Ding, R. Zhu, Y. Li, Y. Wang, W. Sun, D. Wang, L. Wu and L. Zheng, *ChemistrySelect*, 2022, **7**, e202103774.
- 49 M. Králik, *Chem. Pap.*, 2014, **68**, 1625–1638.
- 50 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 51 K. F. Kalz, R. Kraehnert, M. Dvoyashkin, R. Dittmeyer, R. Gläser, U. Krewer, K. Reuter and J.-D. Grunwaldt, *ChemCatChem*, 2017, **9**, 17–29.
- 52 A. Vamvakeros, S. D. Jacques, M. Di Michiel, D. Matras, V. Middelkoop, I. Z. Ismagilov, E. V. Matus, V. V. Kuznetsov, J. Drnec, P. Senecal and A. M. Beale, *Nat. Commun.*, 2018, **9**, 1–11.
- 53 S. D. Jacques, M. Di Michiel, S. A. Kimber, X. Yang, R. J. Cernik, A. M. Beale and S. J. Billinge, *Nat. Commun.*, 2013, **4**, 1–7.
- 54 S. Weber, A. Diaz, M. Holler, A. Schropp, M. Lyubomirskiy, K. L. Abel, M. Kahnt, A. Jeromin, S. Kulkarni, T. F. Keller, R. Gläser and T. L. Sheppard, *Adv. Sci.*, 2022, **9**, 2105432.

