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Operando X-ray absorption spectroscopy (XAS), diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) and mass spectrometry (MS) provide complementary information on the catalyst structure, surface reaction mechanisms and activity relationships. The powerful combination of the techniques has been the driving force to design and engineer suitable spectroscopic *operando* reactors that can mitigate limitations inherent to conventional reaction cells and facilitate experiments under kinetic regimes. Microreactors have recently emerged as effective spectroscopic *operando* cells due to their plug-flow type operation with no dead volume and negligible mass and heat transfer resistances. Here we present a novel microfabricated reactor that can be used for both *operando* XAS and DRIFTS studies. The reactor has a glass-silicon-glass sandwich-like structure with a reaction channel (3000 $\mu\text{m} \times 600 \mu\text{m}$; width $\times$ depth) packed with a catalyst bed (ca. 25 mg) and placed sideways to the X-ray beam, while the infrared beam illuminates the catalyst bed from the top. The outlet of the reactor is connected to MS for continuous monitoring of the reactor effluent. The feasibility of the microreactor is demonstrated by conducting two reactions: i) combustion of methane over 2 wt% Pd/Al₂O₃ studied by *operando* XAS at the Pd K-edge and ii) CO oxidation over 1 wt% Pt/Al₂O₃ catalyst studied by *operando* DRIFTS. The former shows that palladium is in an oxidised state at all studied temperatures, 250, 300, 350, 400 °C and the latter shows the presence of linearly adsorbed CO on the platinum surface. Furthermore, temperature-resolved reduction of palladium catalyst with methane and CO oxidation over platinum catalyst are also studied. Based on these results, the catalyst structure and surface reaction dynamics are discussed, which demonstrate not only the applicability and versatility of the microreactor for combined *operando* XAS and DRIFTS studies, but also illustrate the unique advantages of the microreactor for high space velocity and transient response experiments.

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1. Introduction

The relationship between catalyst structure, activity and product selectivity for a given reaction has been extensively studied to design and develop catalytic processes.^{1–4} Catalyst structure and activity relationships can be explored using a variety of spectroscopic and microscopic techniques and combination of two or more spectroscopic methods in a single reactor can offer complementary information.^{2,4} X-ray absorption spectroscopy (XAS) is an element-specific technique, which provides information on the geometric, electronic and chemical state of an atom in the catalyst.^{1,5–9} One of the most commonly used techniques to study catalysts is infrared (IR) spectroscopy, which interrogates the vibrational state of the catalyst and the adsorbed molecules on the catalyst surface, providing useful information on the reaction and deactivation mechanisms.^{10,11} In particular,

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diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) is suitable for studying catalysts in the form of a powder under reaction conditions.¹² Therefore, the combination of XAS, DRIFTS and MS (mass spectrometry) is proven to offer a holistic approach to study catalyst structure and activity relationships.^{2,13–18}

In the last two decades, there has been a surge in the application of *operando* techniques.^{19–23} However, *operando* cells often suffer from a range of issues including poor sealing, contamination, large dead volume, and temperature and concentration gradients along the catalyst bed, which can lead to biased conclusions.^{19,24,25} Furthermore, in order to reduce mass transfer resistances across the catalyst bed and to derive reliable intrinsic kinetic data, the *operando* cells must be capable of handling high space velocities.^{26,27} Different configurations of *operando* and *in situ* reactor cells have been reported.^{28–34} Among these, it is common to find modified Harrick cells, which have been used for combined DRIFTS and XAS studies.^{32–34} However, these cells have a large dead volume that can lead to a broad residence time distribution, by-pass and inhomogeneous reaction conditions. Agostini *et al.* proposed a spectroscopic cell with a body-dome configuration, which was optimised for time-resolved combined XAS/DRIFTS/MS studies.³⁵ This consisted of a body containing the sample holder and the heater, and a dome including the X-ray and IR windows. The catalyst in powder form was hosted in a crucible, placed in between two carbon-glass windows transparent to X-rays and below a CaF₂ window for IR radiation. The reactor could be operated at up to 600 °C and 5 bar. The dead volume in the reaction chamber was around 0.5 cm³ which increased to 1 cm³ when a dome for higher pressure experiments had to be employed. In other studies, different configurations were reported as alternatives.^{27,36,37} Among these, Chiarello *et al.* demonstrated a spectroscopic cell for *in situ* transmission XAS with DRIFTS.³⁶ In this design, the X-ray and IR beams were focussed on the same optical window and were perpendicular to the catalyst bed. Furthermore, the gas inlet and the outlet shared the same direction and the dead volume was reduced compared to a modified Harrick³³ or SpectraTech cell.³⁸ However, the gas flow was replaced in the cell in 5 s, due to the dead volume of the inlet and outlet tubes and for combined IR-XAS studies the authors had to drill a hole in the CaF₂ window for IR and fill it with carbon based glue to allow X-ray passing through. Dann *et al.*, developed a new spectroscopic packed-bed reactor to study the kinetic oscillations of CO oxidation over a Pd/Al₂O₃ catalyst with combined energy dispersive EXAFS (EDE) and DRIFTS.³⁷ The reactor cell was made of pure aluminium with a square cross sectional channel (5 mm × 5 mm) that hosted the catalyst in the form of pellets (250–335 μm). It had thin walls on two opposite sides, to allow X-ray spectroscopy analysis to be performed in transmission mode, while on top a rectangular CaF₂ window was installed for IR transmission. The heating plate was placed underneath the reactor and the temperature was measured near the outlet region. This cell

was tested up to 140 °C and allowed to perform simultaneous *operando* EDE/DRIFTS at 8 different positions along the packed bed.

The use of microfluidic and microfabricated reactors which can offer accurate control of process conditions, homogeneous temperature distribution and a distinct flow pattern, has been reported for *operando* spectroscopic studies under realistic reaction conditions.^{39–47} Such microreactors are valuable and compact tools for spectroscopic studies^{24,39} including Raman,^{40,48,49} IR,^{50–53} X-ray diffraction (XRD)^{45,54,55} and XAS.^{42,45,55,56} Among such reactors, silicon microfabricated reactors have unique advantages due to their high thermal conductivity, chemical inertness and mechanical stability. Micromachining allows fabricating microreactors with accurate and versatile microchannel geometries. Silicon microreactors, sealed with glass through an anodic bonding process, can be operated up to 500 °C. In particular, the transparency of silicon to X-rays and IR beams (also glass for Raman) makes the glass-bonded silicon microreactor ideal for combined *operando* XAS and DRIFTS studies.⁵⁵ In our previous work, a silver sputter-coated microchannel of a silicon-glass microreactor was employed for *operando* Raman spectroscopic studies of continuous oxidative dehydrogenation of methanol to formaldehyde.⁴⁰ The silver sputter-coated microchannel of the microreactor served itself as a catalyst surface and it enabled us to evaluate the effect of temperature and reaction feed composition on the catalyst structure and activity, and to identify reaction intermediates. The same reactor was also employed for *in situ* XAS studies at the silver K-edge in fluorescence mode.⁴² This work demonstrated the applicability and versatility of the microchannel reactor for multi technique *operando* spectroscopic studies. However, the application of the sputter coated microchannel reactor is limited to unsupported metal catalysts. Therefore, here we report a novel silicon-glass microreactor for supported catalysts that can be used for both *operando* XAS and DRIFTS, addressing many of the limitations imposed by conventional *operando* cells. The applicability and versatility of the microreactor are demonstrated by investigating the Pd K-edge during methane combustion over 2 wt% Pd/Al₂O₃ and surface adsorbed species during carbon monoxide oxidation on 1 wt% Pt/Al₂O₃. The experiments provide new information on catalyst structure and dynamics for these widely studied systems.

2. Materials and methods

2.1. Microreactor cell design and fabrication

The silicon microreactor developed in this work has a glass-silicon-glass sandwich-like structure that can be employed for various spectroscopic studies including XAS, IR and Raman spectroscopy. On the two sides of the silicon layer, microchannels were designed to host a packed-bed of catalyst, inlet and outlet flow channels and slits for XAS measurements. Fig. 1 illustrates the technical drawing of the silicon layer. On the back side of the silicon layer, a

Fig. 1 Technical drawing of the silicon microreactor. Back side (left), front side (right). Dimensions are in millimetres.

serpentine-shaped gas inlet channel ($1000\ \mu\text{m} \times 600\ \mu\text{m}$; width $\times$ depth) leads to the reaction channel ($3000\ \mu\text{m} \times 600\ \mu\text{m}$; width $\times$ depth) that hosts the catalyst bed. A set of small pillars ($500\ \mu\text{m} \times 40\ \mu\text{m}$; length $\times$ width; arranged in $20\ \mu\text{m}$ intervals) is placed at the outlet of the reaction channel acting as catalyst retainer. Perpendicular to and on either side of the reaction channel, six dead-end slits ($2000\ \mu\text{m} \times 600\ \mu\text{m}$; width $\times$ depth) allow the X-ray beam to probe the catalyst bed in transmission mode at six different axial locations along the bed. The silicon wall between the six dead-end X-ray slits and the reaction channel is $250\ \mu\text{m}$ thick. Alternating with the six X-ray slits, seven thermocouple wells ($600\ \mu\text{m} \times 600\ \mu\text{m}$; width $\times$ depth) enable monitoring of the axial temperature profile of the catalyst bed using 0.25

mm-diameter thermocouples (Omega Engineering). Beyond the set of small pillars, the back side shares a common hole with the front side (see top of Fig. 1). This allows the gas coming from the catalyst bed to flow into the outlet channel in the front side of the silicon layer ($600\ \mu\text{m} \times 200\ \mu\text{m}$; width $\times$ depth). The microstructured channels on both sides (front and back) of the silicon layer are sealed with two 0.8 mm thick glass wafers (Corning 7740). The front glass cover has a hollow window ($36\ \text{mm} \times 6\ \text{mm}$; length $\times$ depth) to allow the IR beam to probe the catalyst bed through the silicon wall, and the back-side glass cover has two pre-drilled holes for the gas inlet and outlet connections. Fig. 2 shows a schematic representation of the two sides (back and front) of the microreactor with inlet and outlet flows and X-ray and IR beam paths. A representation of a 15 mm long catalyst bed, preceded by glass beads used to distribute the gas flow and pillars for retaining the catalyst particles at the outlet, is shown in the back side view. The overall thickness of the fabricated reactor is 2.6 mm, with a length and width of 72 mm and 20 mm, respectively. Pictures of the reactor are shown in Fig. S1 (see ESI†).

The microfabrication process was performed *via* double-sided photolithography (Q4000-6, Quintel) followed by deep reactive-ion etching (DRIE, STS ASE) of each side of a double-sided polished 1 mm thick and 100 mm diameter silicon wafer (undoped float zone grown, CRYSTRAN UK). For the wafer, three microfabrication steps were carried out. The front side of the reactor with its gas channel was processed first, patterned and etched to *ca.* $200\ \mu\text{m}$. Except for the inlet/outlet holes, the etched channel was then covered with a photoresist (SPR-220-7, Rohm and Haas) which was applied manually using a small pipette tip. This was followed by a soft baking at $90\ ^\circ\text{C}$ for 10 min. The front side was further etched by *ca.* $200\ \mu\text{m}$ for the holes. Afterwards, the wafer with the processed front side was cleaned thoroughly with Piranha

Back side

Front side

Fig. 2 Schematic representation of the back and front side of the microreactor for *operando* XAS/MS and DRIFTS/MS studies. A packed-bed catalyst (brown) and glass beads (light grey) upstream of the catalyst bed are depicted on the back side image. Insets show detailed view of the pillars used for retaining the catalyst particles and the thermocouple wells.

solution ($\text{H}_2\text{SO}_4:\text{H}_2\text{O}_2 = 3:1$) at 100 °C for 15 min, followed by another lithography step. The back side of the wafer was patterned with the help of infrared beam alignment. Subsequently, DRIE was used to etch for 600 μm to create the reactor channel. The holes on the silicon layer were etched through to form connections between the gas inlet channel and the reaction channel. The micro-fabricated silicon wafer was then cut into three individual reactor chips. Double-sided anodic bonding followed to seal the two sides of the silicon layer with the two glass layers by performing two steps of anodic bonding. The first step was to assemble the front glass layer to the front surface of the silicon layer on a hotplate (Stuart SD162) at a temperature of 420 °C while applying a DC voltage of 500 V across the assembly (cathode attached to the silicon and anode to the glass). After the first anodic bonding, the assembled silicon-glass chip was removed from the hot plate, cooled down and cleaned with Piranha solution. The second step was performed by placing the silicon-glass chip onto the hotplate (when the hot plate was cool enough) with the back surface of the silicon layer facing up and then putting the back glass cover on the top of the silicon chip by aligning the holes on the glass cover to the inlet and outlet holes on the silicon layer. After heating the hotplate to 420 °C, a DC voltage of 500–700 V was applied across the sandwich assembly.

The microreactor was incorporated in an in-house made heating unit (see ESI†). The heating was achieved using a ceramic heater (25 mm $\times$ 50 mm $\times$ 2.5 mm, ULTRAMIC® ceramic heaters, Watlow) which was fitted into a stainless steel holder. High temperature O-rings (Perlast G80A, O Rings Ltd) were used to seal the inlet and outlet connections to the microreactor, which was placed onto the heater holder and fixed firmly in position using the top clamp of the holder. The microreactor can be operated up to a maximum temperature of 400 °C due to the material used for insulating the electrical connections of the thermocouple embedded in the ceramic heater, which can be changed to achieve a higher reaction temperature if required.

2.2. Catalytic combustion of methane

Prior to the *operando* spectroscopic studies, the applicability of the microreactor was tested for the catalytic combustion of methane at atmospheric pressure over a catalyst with a nominal composition of 2 wt% Pd/ Al_2O_3 prepared by a modified impregnation method. The synthesis procedure for this catalyst and its characterisation have been reported elsewhere.⁵⁷ The microreactor was loaded with the catalyst (28.5 mg and 53–63 μm sieve fraction), by applying vacuum at the outlet port, which formed a catalyst bed length of *ca.* 15 mm. Glass beads (75 μm , Sigma Aldrich) were placed at the inlet of the catalyst bed in order to ensure flow equi-distribution. The catalyst was heated up at 10 °C min^{-1} under 10% O_2/He and calcined for 30 min at 400 °C. After calcination, the catalyst was cooled down to 200 °C and the gas was switched from 10% O_2/He to a mixture of 1% CH_4

and 4% of O_2 in He with a flowrate of 40 NmL min^{-1} , corresponding to a space velocity of around 84 250 $\text{cm}^3 \text{g}_{\text{cat}}^{-1} \text{h}^{-1}$. The temperature was increased from 200 °C to 400 °C (at 10 °C min^{-1}) with a steady state dwell of around 45 min every 50 °C increment. The outlet gas composition was analysed by an online gas chromatograph (GC, Agilent 7890A) equipped with a thermal conductivity detector and two columns (HP-PLOT Molesieve and GS-CarbonPLOT). The temperature profile along the catalyst bed was measured in order to assess isothermality of the bed during reaction.

2.3. Operando XAS/MS

Operando XAS/MS experiments for the catalytic combustion of methane over 2 wt% Pd/ Al_2O_3 were conducted at the B18 beamline of the Diamond Light Source (DLS), Didcot, UK. A picture of the setup is shown in Fig. 3, while the schematic of the experimental rig is presented in ESI†. XAS measurements were performed at the Pd K-edge in transmission mode using the quick EXAFS (QEXAFS) setup with a fast-scanning Si(311) double crystal monochromator.⁵⁸ The data acquisition setup consisted of three detectors measuring the incident beam intensity, I_0 , the transmitted beam through the sample, I_t , and the one through the reference, I_{ref} . A standard Pd foil reference was placed between I_t and I_{ref} and each spectrum was acquired with an exposure time of 180 s. The X-ray beam size was optimised to around 200 $\mu\text{m} \times 100 \mu\text{m}$ for the measurement through the narrow X-ray access slits of the microreactor, which was placed on an adjustable stage in order to allow the alignment of the beam inside the X-ray slits. XAS measurements were conducted at the inlet and the outlet positions of the catalyst bed (see back side in Fig. 2). The reaction mixture from the outlet of the microreactor was monitored using an online MKS mass spectrometer.

Operando XAS studies were conducted similarly to the reaction test in section 2.2. The 2 wt% Pd/ Al_2O_3 catalyst particles (27.7 mg of 53–63 μm sieve fraction) were loaded into the microreactor. The catalyst was heated up at 10 °C min^{-1} under 10% O_2/He flow and calcined at 400 °C for 30 min. The temperature was then decreased to 250 °C and the gas flow was switched from 10% O_2/He to 1% CH_4 , 4% O_2 and 5% Ar in He (total flow rate of 40 NmL min^{-1}). The combustion reaction was carried out at 250, 300, 350 and 400 °C (at 10 °C min^{-1}) with a dwell time of around 45 min at each temperature to achieve steady state conditions. Under steady state, the catalyst bed at the inlet region was probed with the X-ray beam in transmission mode.

In another experiment, a temperature-resolved reduction of 2 wt% Pd/ Al_2O_3 was performed with methane in flow. After calcination at 400 °C, the catalyst was cooled down to 225 °C under 10% O_2/He flow. Then the gas mixture was switched from 10% O_2/He to 1% CH_4/He at 40 NmL min^{-1} and the temperature of the reactor was gradually increased from 225 to 300 °C at a rate of 1 °C min^{-1} while monitoring the catalyst at the bed inlet by XAS. After the experiment, the gas mixture was switched back to 10% O_2/He , the temperature

Fig. 3 Microreactor assembly (a) at the beamline B18 at the Diamond Light Source for the *operando* XAS/MS study and (b) at the Research Complex at Harwell for the *operando* DRIFTS/MS study.

increased up to 400 °C and the catalyst was re-calcined. This time the X-ray beam was directed at the outlet bed position and a temperature-resolved reduction was performed again under 1% CH₄/He at 40 NmL min⁻¹ from 225 to 305 °C (1 °C min⁻¹). During the two temperature-resolved reductions, XAS scans were taken every 3.46 min, generating 21 and 24 spectra respectively.

The raw XAS spectra were processed using the software Athena.⁵⁹ The X-ray absorption near edge structure (XANES) spectra were pre-edge subtracted and normalised to the post-edge background. The linear combination fitting (LCF) of the normalised XANES spectra was then performed around the Pd K-edge, 24 150–24 986 eV, using the software Athena.⁵⁹ The XANES spectra of the reference Pd metal foil were used as standard for the palladium metal (Pd⁰) in the LCF, while for the PdO component the spectra at the inlet during calcination were taken as reference. Their weightings were constrained to be between 0 and 1 and their sum to be equal to 1. The fitting quality was assessed using *R*-factors, which were consistently low for all the fittings (*R*-factor = 0.00083 ± 0.00021). An example of a LCF of a XANES spectrum for the catalyst at 250 °C is presented in Fig. S4 (see ESI†). The Fourier transform (FT) of the EXAFS data was performed on the *k*³-weighted functions between 2.7 and 10 Å⁻¹.

2.4. *Operando* DRIFTS/MS

The applicability of the microreactor for *operando* DRIFTS studies was evaluated for the catalytic combustion of methane over the 2 wt% Pd/Al₂O₃ catalyst and for the oxidation of carbon monoxide over a commercial 1 wt% Pt/Al₂O₃, taking DRIFTS spectra in the middle of the catalytic bed, approximately between the inlet and the outlet region. Methane combustion was performed between 250 and 400 °C, which is directly comparable with the *operando* XAS studies. In fact, the same reaction conditions described in section 2.2 were applied in the *operando* DRIFTS study of methane combustion over 2 wt% Pd/Al₂O₃ (28 mg and 53–63

μm sieve fraction). Moreover, the same reaction was also conducted using a commercial high temperature Harrick DRIFTS cell for comparative purposes.

Oxidation of carbon monoxide was performed using a typical amount of 29.6 mg (sieve fraction of 38–53 μm) of a commercial 1 wt% Pt/Al₂O₃ catalyst (Sigma Aldrich). The catalyst was heated up at 10 °C min⁻¹ under 10% O₂/He flow and calcined at 400 °C for 60 min. Then the flow was switched from 10% O₂/He to pure He, and then to 10% H₂/He at 400 °C for 30 min in order to reduce the catalyst. Subsequently, the hydrogen flow was replaced with pure helium at the same temperature to flush the catalyst bed for 15 min and the reactor was cooled to 30 °C under the same helium flow. Temperature-resolved CO oxidation over the reduced Pt/Al₂O₃ catalyst was then conducted between 30 and 400 °C using 10% O₂/He (at 30 NmL min⁻¹) and 10% CO/He (at 10 NmL min⁻¹) reaction mixture. The reaction temperature was ramped at 5 °C/min from 30 to 400 °C and DRIFTS spectra were collected approximately every 30 s, while reaction products were simultaneously monitored by MS. At 400 °C, the reaction mixture was switched to 10% O₂/He for 30 min to calcine the catalyst. Afterwards, the catalyst was cooled to 30 °C (10 °C min⁻¹) under the same O₂/He flow. A temperature-resolved CO oxidation over the oxidised 1 wt% Pt/Al₂O₃ catalyst was carried out using the same reaction mixture and temperature ramp as described above for the CO oxidation with the pre-reduced catalyst.

After the reaction at 400 °C, the temperature of the reactor was decreased to 310 °C and the gas was switched to a stoichiometric composition of 10% O₂/He (5 NmL min⁻¹), 10% CO/He (10 NmL min⁻¹), balance He (25 NmL min⁻¹) to study possible concentration gradients of adsorbed species along the catalyst bed by DRIFTS. For this experiment, the catalyst bed was probed under steady-state conditions with DRIFTS from the inlet to the outlet of the catalyst bed, by moving (at 2 mm steps) the reactor stage.

All experiments were conducted using an Agilent Cary 680 series spectrometer equipped with a Harrick DaVinci arm

(see Fig. 3). The DaVinci arm was fitted with Praying Mantis optics, which focussed the IR beam onto the catalyst bed of the microreactor. The microreactor outlet was connected to a Hidden QGA mass spectrometer for analysis of the reactor effluent. As in the *operando* XAS experiments, the microreactor was placed onto an adjustable stage that was located under the DaVinci arm. The experimental rig schematic is shown in the ESI.† The extended IR focal length of the Praying Mantis optics allowed the microreactor to be located at about 4.7 mm below the DaVinci arm. Prior to DRIFTS experiments, alignment of the IR beam along the central line of the catalyst bed was required. DRIFTS spectra were acquired by taking 64 scans with a resolution of 4 cm⁻¹ using a liquid nitrogen cooled mercury cadmium telluride (MCT) detector.

In all the experiments, the catalyst performance was assessed using the reactant consumption rate, r , according to eqn (1), where F is the inlet molar flowrate, y_{in} and y_{out} are the gas inlet and outlet concentrations (vol%, methane or carbon monoxide) respectively and n_{cat} is the molar amount of metal catalyst.

$$r = \frac{F(y_{\text{in}} - y_{\text{out}})}{n_{\text{cat}}} \quad (1)$$

In some cases the natural logarithm of the consumption rate was plotted against the inverse of absolute temperature (Arrhenius plot). This was done in order to estimate the apparent activation energy, E_a according to the Arrhenius equation (eqn (2)), where R is the ideal gas constant, T the absolute temperature and r_0 a pre-exponential factor. The estimation was performed assuming constant concentration of the reactants on the metal catalyst.

$$\ln(r) = \ln(r_0) - \frac{E_a}{RT} \quad (2)$$

3. Results and discussion

3.1. Catalytic combustion of methane

The combustion of methane over 2 wt% Pd/Al₂O₃ was performed using the microreactor, as described in section 2.2, and the methane outlet concentration is shown in Fig. 4 at four different set temperatures: 250, 300, 350 and 400 °C. The combustion of methane starts at 250 °C with an outlet methane concentration of 0.99% which decreases to 0.48% at 400 °C, equivalent to a methane consumption rate of 107 mol_{CH₄} mol_{Pd}⁻¹ h⁻¹. Similar methane consumption rates over Pd/Al₂O₃ catalysts at 400 °C and atmospheric pressure can be found in previous studies (see Table 1).^{60–64}

Fig. 5 shows the temperature values along the reactor bed during methane combustion. It is evident that no large temperature gradient is present along the glass beads and the catalyst bed, especially below 400 °C. At 400 °C, the temperature profile along the bed shows the highest variation (±1 °C from the set temperature). Based on the conversion

Fig. 4 Methane outlet concentration, $y_{\text{out, CH}_4}$, during combustion over 2 wt% Pd/Al₂O₃ catalyst at different reactor set temperatures: 250, 300, 350 and 400 °C. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 NmL min⁻¹, atmospheric pressure, catalyst amount: 28.5 mg (53–63 μm).

and temperature profiles obtained, the microreactor design and performance was judged to be satisfactory to be employed for *operando* XAS/MS studies for the combustion of methane.

From the observed reaction rate (107 mol_{CH₄} mol_{Pd}⁻¹ h⁻¹), the catalyst particle size (53–63 μm sieve fraction) and the gas physical properties at 400 °C, the Péclet (Pe), Mears (MR) and Weisz–Prater (C_{WP}) numbers can be calculated (see Table 2). The large Péclet number suggests that axial dispersion within the packed bed can be neglected, hence the microreactor can be regarded as an ideal plug-flow reactor. The low Mears and Weisz–Prater numbers indicate that the reactor operates under no external or internal mass transfer resistances, respectively. The details on the calculation of these numbers are reported in the ESI.†

3.2. *Operando* XAS/MS

3.2.1 Combustion of methane on Pd/Al₂O₃. The *operando* XANES spectra collected during the combustion of methane at the inlet of the catalyst bed are presented along with outlet methane concentration obtained by MS in Fig. 6. To begin with, the MS data shows that methane combustion already starts at 250 °C and a methane conversion of around 50% is observed at 400 °C (Fig. 6(a)), which is in excellent agreement with laboratory tests reported in Fig. 4. From the conversion profile, the reaction rate against the inverse of temperature is presented in Fig. 6(b). The linear fitting for the four experimental points shows an apparent activation energy of methane oxidation over Pd/Al₂O₃ of around 80.7 kJ mol⁻¹, which is similar to the value reported by van Giezen *et al.* (86 kJ mol⁻¹) using a pre-calcined 7.3 wt% Pd/Al₂O₃ in a reacting mixture of 1% CH₄ and 4% O₂ in helium.⁶⁵ A slightly lower value was reported by Ribeiro *et al.* (76 kJ mol⁻¹) for a pre-calcined 8.5 wt% Pd/Al₂O₃, similarly to the 75 kJ mol⁻¹ found by Mouaddib *et al.* using a 1.93 wt% Pd/Al₂O₃ catalyst aged

Table 1 Catalytic activity of Pd/Al₂O₃ catalysts at 400 °C and atmospheric pressure found in the literature

Reference	wt% Pd	Catalyst pretreatment	$y_{\text{in,CH}_4}$, %	O ₂ /CH ₄ , –	r_{CH_4} , mol _{CH₄} mol _{Pd} ^{–1} h ^{–1}
Burch <i>et al.</i> ⁶⁰	4	Calcination	0.8	20	91
Briot <i>et al.</i> ⁶¹	1.95	Reduction	1	4	10
Briot <i>et al.</i> ⁶¹	1.95	Ageing	1	4	65
Burch <i>et al.</i> ⁶²	4	Calcination	0.3	5	35
Mouaddib <i>et al.</i> ⁶³	1.93	Reduction	1	4	5
Mouaddib <i>et al.</i> ⁶³	1.93	Ageing	1	4	70
This work	2	Calcination	1	4	107

Fig. 5 Temperature profiles along the reactor bed during methane combustion over 2 wt% Pd/Al₂O₃ catalyst at different reactor set temperatures: 250, 300, 350 and 400 °C. The thermocouple positions are at the six of the seven thermowells shown in the figure on the left.

under reaction conditions at 600 °C.^{63,66} From the overlay of XANES spectra, it appears that the white line of the Pd K-edge at 24368 eV is essentially the same at all studied reaction temperatures and the spectra are similar to that of the PdO (Fig. 6(c)). Further insights on the catalyst phase composition are derived by the LCF method (Fig. 6(d)). In general, it is evident from LCF fitting that the palladium was largely in the oxidised state in the Pd/Al₂O₃ catalyst during the reaction at all studied temperatures. However, the palladium metal component in the catalyst appeared to be higher (*ca.* 7.7%) at 250 °C than at other temperatures.

In agreement with XANES and LCF data (Fig. 6(c) and (d)), the Fourier transform (FT) of the corresponding EXAFS data shows a contribution from oxygen in the first coordination shell of the PdO like structure at *ca.* 1.5 Å (phase shift uncorrected) and another contribution from the corresponding Pd–Pd pair in the second coordination shell at around 3 Å (see Fig. 7). No other peaks attributable to metallic palladium are apparent except for a small contribution from Pd–Pd metal (*ca.* 2.4 Å) at 250 °C. These

results further indicate that the palladium was predominantly in oxidised state in the catalyst during reaction at all studied temperatures.

3.2.2 Temperature-resolved reduction of Pd/Al₂O₃ with methane. The oxidation and coordination state of palladium at the inlet of the Pd/Al₂O₃ catalyst bed was studied by XAS during the temperature-resolved reduction of the catalyst. During reduction the temperature increased by approx. 3.5 °C between two consecutive scans (3.46 min). In this dynamic situation, the microreactor, due to its excellent heat transfer and short residence time (0.04 s) of the flowing gas (40 NmL min^{–1}) in the narrow catalyst channel (0.027 mL when empty), could instantaneously adapt and provide time-dependent data.

The stacked XANES, PdO and Pd metal component extracted from LCF analysis of XANES data and the FT of the EXAFS data collected for the catalyst at the reactor inlet are plotted as function of reaction temperature in Fig. 8. It is evident from the change in the white line of the XANES spectra that the palladium reduction occurred already at around 240 °C. The palladium speciation at the catalyst bed inlet is determined in more detail from XANES spectra by application of LCF and it can be readily seen from Fig. 8(b) that the PdO and Pd⁰ components change with the increase in temperature. At 225 °C, immediately after calcination at 400 °C for 30 min and at the beginning of the reduction

Table 2 Péclet, Pe, Mears, MR, and Weisz–Prater, C_{WP} , numbers for the microreactor during the combustion of methane at 400 °C

Pe	MR	C_{WP}
61 (>6)	0.07 (<0.15)	0.003 (≪1)

Fig. 6 Methane combustion over 2 wt% Pd/Al₂O₃ catalyst: (a) methane outlet concentration, y_{out,CH_4} , (b) Arrhenius plot of the reaction rate, (c) normalised operando XANES spectra at the Pd K-edge of the inlet of the catalyst bed and (d) PdO and Pd⁰ and concentrations obtained by LCF of the spectra. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 NmL min⁻¹, atmospheric pressure, catalyst amount: 27.7 mg (53–63 μm).

process, it appears that the inlet of the catalyst bed was almost completely oxidised as evident from the low presence of a metallic Pd phase fraction (2.1%). This was determined by LCF and indicates that the calcination was effective. During the temperature-resolved reduction, the isosbestic point (*i.e.*, the point at which the phase composition of two species is equal) is observed at 246 °C. These results reveal that the reduction of palladium oxide at the inlet of the catalyst bed was gradual.

These observations are further corroborated by the FT of the EXAFS (Fig. 8(c)). At 225 °C the catalyst at the bed inlet

presents two peaks at *ca.* 1.5 Å (phase shift uncorrected) in the first and at *ca.* 3 Å in the second coordination shells, respectively, attributable to Pd–O and Pd–Pd bonds of PdO like species. The presence of these two peaks at 225 °C suggests that the catalyst was mainly in the oxidised state, consistent with XANES. During the temperature-resolved reduction of the catalyst, the characteristic peak of the Pd–Pd in the first coordination shell of metallic Pd emerges at around 2.4 Å (phase shift uncorrected), with the concomitant progressive decrease of the PdO peaks in the first coordination shell at around 1.5 Å (phase shift uncorrected), which indicates the reduction of palladium oxide to metallic Pd. The amplitude of the metallic Pd–Pd peak in the first coordination shell establishes at around 250 °C, which is in excellent agreement with XANES and LCF.

The outlet region of the catalyst bed was investigated in a similar way after re-calcination of the catalyst, and results are reported in Fig. 9. It is evident from the change in the white line of the XANES spectra (Fig. 9(a)) that the palladium reduction occurred at around 280 °C, which is at higher temperature compared to the bed inlet (Fig. 8). This could be ascribed to the re-calcination before the second temperature-resolved reduction, which might have sintered the catalyst. The metal palladium speciation at the outlet can be seen in Fig. 9(b). At the beginning of the second reduction, it appears that the catalyst bed was not completely oxidised as evident from the presence of a metallic Pd phase fraction (9.9%).

Fig. 7 Phase shift uncorrected Fourier transform magnitude of the Pd K-edge EXAFS of the 2 wt% Pd/Al₂O₃ catalyst during methane combustion. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 NmL min⁻¹, atmospheric pressure, catalyst amount: 27.7 mg (53–63 μm).

Fig. 8 XAS study at the Pd K-edge of the 2 wt% Pd/Al₂O₃ catalyst at the bed inlet position during the first temperature-resolved reduction with methane: (a) normalised XANES spectra, (b) PdO and Pd⁰ concentrations obtained by LCF of the spectra and (c) phase shift uncorrected Fourier transform of EXAFS. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 Nml min⁻¹, atmospheric pressure, catalyst amount: 27.7 mg (53–63 μm).

This suggests that the first calcination was more effective than the second one.

The isosbestic point is observed at 285 °C, unlike for the inlet bed location (246 °C). However, it is important to note that the occurrence of metallic Pd above 300 °C was the same (*ca.* 92%) at the inlet and outlet of the catalyst bed, for the first and second calcination respectively. The FT of the EXAFS (Fig. 9(c)) shows the characteristic peak of the Pd–Pd in the first coordination shell of (2.4 Å) emerging at around 290 °C and at 300 °C the Pd phase dominated at the outlet bed position. These results indicate that spatial information along the catalyst bed could be derived employing this novel microreactor, enabling the user to avoid averaging out the information along the catalyst bed, which is typically the case with conventional spectroscopic cells used for XAS measurements.

3.3. Operando DRIFTS/MS studies

3.3.1 Combustion of methane on Pd/Al₂O₃. The MS and DRIFTS data obtained during methane combustion are presented in Fig. 10. The conversion of methane starts at 250 °C and a 46% methane outlet concentration is observed at

400 °C (Fig. 10(a)), which is in line with the results from the *operando* XAS (Fig. 6(a)) and the laboratory tests (Fig. 4). The corresponding DRIFTS data collected with the microreactor present weak features at around 3000 and 2360/2300 cm⁻¹, which are attributed to gaseous methane (ν C–H), and gaseous carbon dioxide, respectively (Fig. 10(b)).^{67–70}

The gas phase features significantly dominate the spectra collected using the commercial Harrick DRIFTS cell (Fig. 10(c)) and the gas phase fingerprints of methane and carbon dioxide bands appear at 3016 cm⁻¹ and 2300–2360 cm⁻¹, respectively. As also shown in previous IR studies of methane combustion, methane and carbon dioxide are predominantly detected in their gaseous forms.^{67–69} The absence of strong gas phase bands in the infrared spectra collected with the microreactor is attributed to the lack of dead volume in the microreactor, unlike in the commercial DRIFTS cell. The lack of a gas-filled head space (*i.e.* dead volume) above the catalyst bed effectively minimises the interference of gas phase bands from the bands arising from the surface adsorbed species, making the microreactor ideal for transient response *operando* DRIFTS studies.³⁶

3.3.2 Temperature-resolved CO oxidation on Pt/Al₂O₃. The temperature-resolved CO oxidation over pre-reduced and pre-

Fig. 9 XAS study at the Pd K-edge of the 2 wt% Pd/Al₂O₃ catalyst at the bed outlet position during the second temperature-resolved reduction with methane: (a) normalised XANES spectra, (b) PdO and Pd⁰ concentrations obtained by LCF of the spectra and (c) phase shift uncorrected Fourier transform of EXAFS. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 N mL min⁻¹, atmospheric pressure, catalyst amount: 27.7 mg (53–63 μm).

oxidised Pt/Al₂O₃ was followed by DRIFTS analysis at the centre of the catalyst bed. Fig. 11 shows reactor outlet concentrations obtained by MS and DRIFTS spectra collected between 50 and 300 °C during CO oxidation over the pre-reduced catalyst. From Fig. 11(a) it is clear that the gas phase CO₂ appears at around 180 °C which coincides with a drop in the CO and O₂ concentration in the gas phase, indicating occurrence of CO oxidation. At around 240 °C, CO disappears from the gas phase, while O₂ and CO₂ concentrations plateau, suggesting complete conversion of CO. The corresponding DRIFTS data show two bands at 2060 and 2072 cm⁻¹. The bands may be ascribed to linearly adsorbed CO onto two different platinum sites.^{71,72} The band at 2060 cm⁻¹ is often attributed to CO adsorption on coordinatively unsaturated platinum sites at steps, edges and kinks of smaller particles, while the band at 2070 cm⁻¹ is assigned to CO adsorbed on terrace platinum sites of large crystals.⁷² The intensity of the bands increases with increasing temperature from 50 to 240 °C and thereafter the bands disappear abruptly, in agreement with

MS data that show complete conversion of CO at around 240 °C.

Bowker elucidated the CO “light-off”, using X-ray photoelectron spectroscopy over Pd catalyst during CO oxidation.⁷³ At low temperature the surface coverage is dominated by adsorbed CO, which poisons the catalyst and prevents oxidation. However, at a certain temperature molecular oxygen starts adsorbing and dissociating into atoms, which react with adsorbed CO to produce CO₂. This process frees four free sites for further oxygen adsorption, making the reaction self-accelerating. Therefore, the abrupt disappearance of the CO band can be ascribed to the complete conversion of adsorbed CO, which at high temperature could not be detected by DRIFTS due to its fast reaction with oxygen. Based on these results, CO adsorbed on the catalyst surface hinders the activation of oxygen below 180 °C and above this temperature oxygen activation starts to take place, with CO oxidation light-off observed in the MS results.

Fig. 12 shows reactor outlet concentrations and DRIFTS spectra collected between 50 and 300 °C during CO oxidation

Fig. 10 Methane combustion over 2 wt% Pd/Al₂O₃ catalyst: (a) methane outlet concentration, y_{out,CH_4} , (b) infrared spectra recorded using the microreactor and (c) spectra obtained using a commercial Harrick cell. Gas inlet composition: 1% CH₄ and 4% of O₂ in He, flowrate: 40 NmL min^{−1}, atmospheric pressure, catalyst amount: 28 mg (53–63 μm).

over the pre-oxidised catalyst. From Fig. 12(a) it is clear that the gas phase CO₂ appears at around 200 °C, which coincides with a drop in the CO and O₂ concentration in the gas phase, indicating the occurrence of CO oxidation. At around 250 °C,

CO disappears from the gas phase, while O₂ and CO₂ concentrations stabilise. From these results, it is evident that the gas phase conversion profiles of CO, O₂ and CO₂ shift to *ca.* 20 °C higher temperatures over the pre-oxidised catalyst as compared to the pre-reduced one reported in Fig. 11. The DRIFTS data show a band at 2086 cm^{-1} appearing between 50 and 120 °C, which is different from that on the pre-reduced platinum surface (Fig. 11). Above 120 °C, the band at 2086 cm^{-1} starts to redshift to 2078 cm^{-1} and a new band appears to emerge at 2060 cm^{-1} . These two bands at 2078 and 2060 cm^{-1} become prominent at around 135 °C and grow in intensity up to 260 °C during the temperature-resolved oxidation. Above 260 °C the bands disappear abruptly, similar to that observed for the pre-reduced catalyst (Fig. 11), which coincides with the MS results that show complete conversion of CO at around 250 °C.

The logarithmic values of CO consumption rates on the pre-reduced and pre-oxidised catalysts are plotted against the inverse of temperature in Fig. 13. The apparent activation energy is derived from the CO conversion range between 7 and 30%. It is evident from Fig. 13 that the reaction rates and activation energies are of similar magnitudes for both the pre-reduced and pre-oxidised Pt/Al₂O₃ catalysts. In particular, the activation energy for the pre-reduced catalyst, determined to be *ca.* 46.6 kJ mol^{−1}, is slightly lower than that observed for the pre-oxidised catalyst (48.4 kJ mol^{−1}). In general, the activation energies are consistent with earlier studies. Different global or apparent activation energies for CO oxidation over Pt catalysts have been reported previously.^{74–77} An apparent activation energy of 56 kJ mol^{−1} was reported for the CO oxidation on a pre-reduced 5 wt% Pt/SiO₂,⁷⁷ while a value of 84 kJ mol^{−1} was reported for calcined 1–2 wt% Pt/Al₂O₃ catalysts, which was found to be independent of the platinum metal dispersion.⁷⁴

3.3.3 Spatially-resolved DRIFTS of adsorbed CO on Pt/Al₂O₃. The adsorbed CO over 1 wt% Pt/Al₂O₃ was studied using DRIFTS during CO oxidation and the corresponding

Fig. 11 Temperature-resolved CO oxidation over pre-reduced 1 wt% Pt/Al₂O₃ catalyst: (a) concentrations of the gas phase species, y_{out} , and (b) infrared spectra in the middle of the catalyst bed during the temperature-resolved CO oxidation. Gas inlet composition: 7.5% O₂ and 2.5% of CO in He, flowrate: 40 NmL min^{−1}, atmospheric pressure, catalyst amount: 29.6 mg (38–53 μm).

Fig. 12 Temperature-resolved CO oxidation over pre-oxidised 1 wt% Pt/Al₂O₃ catalyst: (a) concentrations of the gas phase species, y_{out} , and (b) infrared spectra in the middle of the catalyst bed during the temperature-resolved CO oxidation. Gas inlet composition: 7.5% O₂ and 2.5% of CO in He, flowrate: 40 NmL min⁻¹, atmospheric pressure, catalyst amount: 29.6 mg (38–53 μ m).

Fig. 13 Arrhenius plots of the CO consumption rate for (a) pre-reduced and (b) pre-oxidised 1 wt% Pt/Al₂O₃ catalyst.

DRIFTS spectra were collected from the inlet to the outlet of the catalyst bed and plotted in Fig. 14. The spectra present a prominent band at 2060 cm⁻¹, which contains a shoulder at 2075 cm⁻¹. The ratio of the intensity of the peaks is

essentially identical along the catalyst bed, indicating the same proportion of CO adsorbed species along the reactor. Earlier studies have similarly reported no change in the adsorbed CO along a fixed-bed *operando* reactor during

Fig. 14 Infrared spectra of CO adsorbed over pre-oxidised 1 wt% Pt/Al₂O₃ catalyst during CO oxidation at 310 °C. The red dot and arrow on the left indicate the DRIFTS beam and the scanning direction along the catalyst bed. Gas inlet composition: 1.25% O₂ and 2.5% of CO in He, flowrate: 40 NmL min⁻¹, atmospheric pressure, catalyst amount: 29.6 mg (38–53 μ m).

spatially-resolved DRIFTS of CO oxidation over a Pt/CeO₂-Al₂O₃ catalyst.^{51,78}

4. Summary and conclusions

We have developed a novel silicon-glass plug-flow microreactor operating under no external and internal mass transfer resistances, suitable for *operando* X-ray absorption spectroscopy (XAS), diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) and mass spectrometry (MS) studies. The applicability and versatility of the microreactor for *operando* XAS and DRIFTS studies is demonstrated in the catalytic combustion of methane and carbon monoxide oxidation. The reactor design and materials used resulted in an isothermal temperature profile along the catalyst bed. *Operando* XAS at the Pd K-edge conducted in transmission mode during methane combustion over Pd/Al₂O₃ showed that the palladium was in oxidised state at all studied temperatures at the inlet of the catalyst bed. Linear combination fitting showed a percentage of PdO larger than 90% for all the temperatures. The corresponding *operando* DRIFTS presented very weak features attributable to gas phase methane and CO₂ due to the absence of dead volume. Temperature-resolved *operando* XAS during the reduction of the palladium catalyst with methane showed that the inlet and the outlet regions of the catalyst exhibited a different reduction temperature possibly due to ageing of the catalyst after re-calcination. *Operando* DRIFTS during the temperature-resolved CO oxidation indicated the presence of linearly adsorbed CO on the pre-reduced and pre-oxidised platinum surface. Spatially-resolved DRIFTS has been performed during the CO oxidation at 310 °C, by probing 9 different axial positions along the catalyst bed. Activation energies have been calculated for both methane combustion over 2 wt% Pd/Al₂O₃ (80.7 kJ mol⁻¹) and CO oxidation over a reduced (46.6 kJ mol⁻¹) and oxidised (48.4 kJ mol⁻¹) 1 wt% Pt/Al₂O₃ surface, and were found to be in the same order of magnitude of those reported in previous studies. This novel microreactor with no dead volume facilitates experiments with high space velocities under true kinetic regimes and is particularly suited for transient response experiments.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 J. M. Thomas, C. R. A. Catlow and G. Sankar, *Chem. Commun.*, 2002, 2921–2925.
- 2 M. A. Newton and W. van Beek, *Chem. Soc. Rev.*, 2010, **39**, 4845–4863.
- 3 M. A. Bañares, *Catal. Today*, 2005, **100**, 71–77.
- 4 B. M. Weckhuysen, *Chem. Soc. Rev.*, 2010, **39**, 4557–4559.
- 5 S. Bordiga, E. Groppo, G. Agostini, J. A. van Bokhoven and C. Lamberti, *Chem. Rev.*, 2013, **113**, 1736–1850.
- 6 G. Sankar and J. M. Thomas, *Top. Catal.*, 1999, **8**, 1–21.
- 7 G. Sankar, J. M. Thomas and C. R. A. Catlow, *Top. Catal.*, 2000, **10**, 255–264.
- 8 J. M. Thomas and G. Sankar, *J. Synchrotron Radiat.*, 2001, **8**, 55–60.
- 9 J. M. Thomas and G. Sankar, *Acc. Chem. Res.*, 2001, **34**, 571–581.
- 10 N.-Y. Topsøe, *Catal. Today*, 2006, **113**, 58–64.
- 11 J. Ryczkowski, *Catal. Today*, 2001, **68**, 263–381.
- 12 T. Armaroli, T. Bécue and S. Gautier, *Oil Gas Sci. Technol.*, 2004, **59**, 215–237.
- 13 C. M. A. Parlett, C. V. Gaskell, J. N. Naughton, M. A. Newton, K. Wilson and A. F. Lee, *Catal. Today*, 2013, **205**, 76–85.
- 14 Y. Zhou, D. E. Doronkin, M. Chen, S. Wei and J.-D. Grunwaldt, *ACS Catal.*, 2016, **6**, 7799–7809.
- 15 A. M. Abdel-Mageed, G. Kučerová, J. Bansmann and R. J. Behm, *ACS Catal.*, 2017, **7**, 6471–6484.
- 16 E. K. Gibson, E. M. Crabb, D. Gianolio, A. E. Russell, D. Thompson and P. P. Wells, *Catal., Struct. React.*, 2017, **3**, 5–12.
- 17 E. K. Dann, E. K. Gibson, R. A. Catlow, P. Collier, T. Eralp Erden, D. Gianolio, C. Hardacre, A. Kroner, A. Raj, A. Goguet and P. P. Wells, *Chem. Mater.*, 2017, **29**, 7515–7523.
- 18 D. Ferri, M. S. Kumar, R. Wirz, A. Eyssler, O. Korsak, P. Hug, A. Weidenkaff and M. A. Newton, *Phys. Chem. Chem. Phys.*, 2010, **12**, 5634–5646.
- 19 A. Chakrabarti, M. E. Ford, D. Gregory, R. Hu, C. J. Keturakis, S. Lwin, Y. Tang, Z. Yang, M. Zhu, M. A. Bañares and I. E. Wachs, *Catal. Today*, 2017, **283**, 27–53.
- 20 A. M. Beale, A. M. J. van der Eerden, K. Kervinen, M. A. Newton and B. M. Weckhuysen, *Chem. Commun.*, 2005, 3015–3017.
- 21 Y. Li, D. Zakharov, S. Zhao, R. Tappero, U. Jung, A. Elsen, P. Baumann, R. G. Nuzzo, E. A. Stach and A. I. Frenkel, *Nat. Commun.*, 2015, **6**, 7583.
- 22 B. M. Weckhuysen, *Phys. Chem. Chem. Phys.*, 2003, **5**, 4351–4360.
- 23 H. Topsøe, *J. Catal.*, 2003, **216**, 155–164.
- 24 N. Al-Rifai, E. Cao, V. Dua and A. Gavrilidis, *Curr. Opin. Chem. Eng.*, 2013, **2**, 338–345.
- 25 F. C. Meunier, *Chem. Soc. Rev.*, 2010, **39**, 4602–4614.
- 26 J. D. Grunwaldt, M. Caravati, S. Hannemann and A. Baiker, *Phys. Chem. Chem. Phys.*, 2004, **6**, 3037–3047.
- 27 S. K. Matam, O. Korsak, L. Bocher, D. Logvinovich, P. Hug, A. Weidenkaff and D. Ferri, *Top. Catal.*, 2011, **54**, 1213.
- 28 C. Lamberti, C. Prestipino, S. Bordiga, G. Berlier, G. Spoto, A. Zecchina, A. Lalon, F. La Manna, F. D'Anca, R. Felici, F.

- D'Acapito and P. Roy, *Nucl. Instrum. Methods Phys. Res., Sect. B*, 2003, **200**, 196–201.
- 29 N. Weiher, E. Bus, B. Gorzolnik, M. Moller, R. Prins and J. A. van Bokhoven, *J. Synchrotron Radiat.*, 2005, **12**, 675–679.
- 30 A. Longo, A. Balerna, F. d'Acapito, F. D'Anca, F. Giannici, L. F. Liotta, G. Pantaleo and A. Martorana, *J. Synchrotron Radiat.*, 2005, **12**, 499–505.
- 31 S. B. Rasmussen, R. López-Medina, R. Portela, E. Mikolajska, M. Daturi, P. Ávila and M. A. Bañares, *Catal. Sci. Technol.*, 2015, **5**, 4942–4945.
- 32 N. S. Marinkovic, Q. Wang and A. I. Frenkel, *J. Synchrotron Radiat.*, 2011, **18**, 447–455.
- 33 K. A. Beyer, H. Zhao, O. J. Borkiewicz, M. A. Newton, P. J. Chupas and K. W. Chapman, *J. Appl. Crystallogr.*, 2014, **47**, 95–101.
- 34 S. Yao, K. Mudiysanlagel, W. Xu, A. C. Johnston-Peck, J. C. Hanson, T. Wu, D. Stacchiola, J. A. Rodriguez, H. Zhao, K. A. Beyer, K. W. Chapman, P. J. Chupas, A. Martínez-Arias, R. Si, T. B. Bolin, W. Liu and S. D. Senanayake, *ACS Catal.*, 2014, **4**, 1650–1661.
- 35 G. Agostini, D. Meira, M. Monte, H. Vitoux, A. Iglesias-Juez, M. Fernandez-Garcia, O. Mathon, F. Meunier, G. Berruyer, F. Perrin, S. Pasternak, T. Mairs, S. Pascarelli and B. Gorges, *J. Synchrotron Radiat.*, 2018, **25**, 1745–1752.
- 36 G. L. Chiarello, M. Nachtegaal, V. Marchionni, L. Quaroni and D. Ferri, *Rev. Sci. Instrum.*, 2014, **85**, 074102.
- 37 E. K. Dann, E. K. Gibson, C. R. A. Catlow, V. Celorrio, P. Collier, T. Eralp, M. Amboage, C. Hardacre, C. Stere, A. Kroner, A. Raj, S. Rogers, A. Goguet and P. P. Wells, *J. Catal.*, 2019, **373**, 201–208.
- 38 M. A. Newton, *Top. Catal.*, 2009, **52**, 1410–1424.
- 39 J. Yue, J. C. Schouten and T. A. Nijhuis, *Ind. Eng. Chem. Res.*, 2012, **51**, 14583–14609.
- 40 E. Cao, S. Firth, P. F. McMillan and A. Gavrilidis, *Catal. Today*, 2007, **126**, 119–126.
- 41 E. Cao, M. Sankar, S. Firth, K. F. Lam, D. Bethell, D. K. Knight, G. J. Hutchings, P. F. McMillan and A. Gavrilidis, *Chem. Eng. J.*, 2011, **167**, 734–743.
- 42 G. Sankar, E. Cao and A. Gavrilidis, *Catal. Today*, 2007, **125**, 24–28.
- 43 B. A. Rizkin, F. G. Popovic and R. L. Hartman, *J. Vac. Sci. Technol.*, 2019, **37**, 050801.
- 44 S. Zhao, Y. Li, E. Stavitski, R. Tappero, S. Crowley, M. J. Castaldi, D. N. Zakharov, R. G. Nuzzo, A. I. Frenkel and E. A. Stach, *ChemCatChem*, 2015, **7**, 3683–3691.
- 45 S. J. A. Figueroa, D. Gibson, T. Mairs, S. Pasternak, M. A. Newton, M. Di Michiel, J. Andrieux, K. C. Christoforidis, A. Iglesias-Juez, M. Fernandez-Garcia and C. Prestipino, *J. Appl. Crystallogr.*, 2013, **46**, 1523–1527.
- 46 D. E. Doronkin, S. Baier, T. Sheppard, F. Benzi and J. D. Grunwaldt, *J. Phys.: Conf. Ser.*, 2016, **712**, 012030.
- 47 A. Urakawa, N. Maeda and A. Baiker, *Angew. Chem., Int. Ed.*, 2008, **47**, 9256–9259.
- 48 A. Urakawa, F. Trachsel, P. R. von Rohr and A. Baiker, *Analyst*, 2008, **133**, 1352–1354.
- 49 P. Beato, R. Kraehnert, S. Engelschalt, T. Frank and R. Schlögl, *Chem. Eng. J.*, 2008, **135**, S247–S253.
- 50 C. K. C. Tan, W. N. Delgass and C. D. Baertsch, *Appl. Catal., B*, 2009, **93**, 66–74.
- 51 C. Daniel, M. O. Clarté, S. P. Teh, O. Thion, H. Provendier, A. C. Van Veen, B. J. Beccard, Y. Schuurman and C. Mirodatos, *J. Catal.*, 2010, **272**, 55–64.
- 52 I. P. Silverwood, N. Al-Rifai, E. Cao, D. J. Nelson, A. Chutia, P. P. Wells, S. P. Nolan, M. D. Frogley, G. Cinque, A. Gavrilidis and C. R. A. Catlow, *Rev. Sci. Instrum.*, 2016, **87**, 024101.
- 53 E. Gross, X.-Z. Shu, S. Alayoglu, H. A. Bechtel, M. C. Martin, F. D. Toste and G. A. Somorjai, *J. Am. Chem. Soc.*, 2014, **136**, 3624–3629.
- 54 J.-D. Grunwaldt and B. S. Clausen, *Top. Catal.*, 2002, **18**, 37–43.
- 55 S. Baier, A. Rochet, G. Hofmann, M. Kraut and J.-D. Grunwaldt, *Rev. Sci. Instrum.*, 2015, **86**, 065101.
- 56 J.-D. Grunwaldt, B. Kimmerle, S. Hannemann, A. Baiker, P. Boye and C. G. Schroer, *J. Mater. Chem.*, 2007, **17**, 2603–2606.
- 57 N. Richards, J. H. Carter, E. Nowicka, L. A. Parker, S. Patisson, Q. He, N. F. Dummer, S. Golunski and G. J. Hutchings, *Appl. Catal., B*, 2020, **264**, 118501.
- 58 A. J. Dent, G. Cibin, S. Ramos, S. A. Parry, D. Gianolio, A. D. Smith, S. M. Scott, L. Varandas, S. Patel, M. R. Pearson, L. Hudson, N. A. Krumpa, A. S. Marsch and P. E. Robbins, *J. Phys.: Conf. Ser.*, 2013, **430**, 012023.
- 59 B. Ravel and M. Newville, *J. Synchrotron Radiat.*, 2005, **12**, 537–541.
- 60 R. Burch, F. J. Urbano and P. K. Loader, *Appl. Catal., A*, 1995, **123**, 173–184.
- 61 P. Briot and M. Primet, *Appl. Catal.*, 1991, **68**, 301–314.
- 62 R. Burch and P. K. Loader, *Appl. Catal., B*, 1994, **5**, 149–164.
- 63 N. Mouaddib, C. Feumi-Jantou, E. Garbowski and M. Primet, *Appl. Catal., A*, 1992, **87**, 129–144.
- 64 S. K. Matam, G. L. Chiarello, Y. Lu, A. Weidenkaff and D. Ferri, *Top. Catal.*, 2013, **56**, 239–242.
- 65 J. C. van Giezen, F. R. van den Berg, J. L. Kleinen, A. J. van Dillen and J. W. Geus, *Catal. Today*, 1999, **47**, 287–293.
- 66 F. H. Ribeiro, M. Chow and R. A. Dallabetta, *J. Catal.*, 1994, **146**, 537–544.
- 67 W. R. Schwartz, D. Ciuparu and L. D. Pfefferle, *J. Phys. Chem.*, 2012, **116**, 8587–8593.
- 68 M. Schmal, M. M. V. M. Souza, V. V. Alegre, M. A. P. da Silva, D. V. César and C. A. C. Perez, *Catal. Today*, 2006, **118**, 392–401.
- 69 H. Na, Z. Liu and T. Zhu, *React. Kinet., Mech. Catal.*, 2014, **111**, 137–148.
- 70 O. Demoulin, M. Navez and P. Ruiz, *Appl. Catal., A*, 2005, **295**, 59–70.
- 71 C. Brieger, J. Melke, N. van der Bosch, U. Reinholz, H. Riesemeier, A. Guilherme Buzanich, M. K. Kayarkatte, I. Derr, A. Schökel and C. Roth, *J. Catal.*, 2016, **339**, 57–67.
- 72 S. K. Matam, E. V. Kondratenko, M. H. Aguirre, P. Hug, D. Rentsch, A. Winkler, A. Weidenkaff and D. Ferri, *Appl. Catal., B*, 2013, **129**, 214–224.

- 73 M. Bowker, *Chem. Soc. Rev.*, 2008, **37**, 2204–2211.
- 74 A. D. Allian, K. Takanabe, K. L. Fajdala, X. Hao, T. J. Truex, J. Cai, C. Buda, M. Neurock and E. Iglesia, *J. Am. Chem. Soc.*, 2011, **133**, 4498–4517.
- 75 R. H. Nibbelke, M. A. J. Campman, J. H. B. J. Hoebink and G. B. Marin, *J. Catal.*, 1997, **171**, 358–373.
- 76 M. Campman, *PhD thesis*, University of Eindhoven, 1996.
- 77 N. W. Cant, P. C. Hicks and B. S. Lennon, *J. Catal.*, 1978, **54**, 372–383.
- 78 C. Daniel, M.-O. Clarté, H. Provendier, A. C. Van Veen, Y. Schuurman, B. J. Beccard and C. Mirodatos, *C. R. Chim.*, 2009, **12**, 647–653.