

PAPER



Cite this: DOI: 10.1039/d4cy00665h

Controlling palladium particle size and dispersion as a function of loading by chemical vapour impregnation: an investigation using propane total oxidation as a model reaction†

Liam A. Bailey, * Mark Douthwaite, Thomas E. Davies,
David J. Morgan  and Stuart H. Taylor *

A series of Pd/Al₂O₃ catalysts with metal weight loadings of 1.0 wt%, 2.5 wt%, and 5.0 wt% were synthesised by chemical vapour impregnation (CVI) and used for the total oxidation of propane. All the catalysts were highly active for propane total oxidation. Extensive characterisation showed essentially identical catalyst structural and chemical characteristics, with consistent nanoparticle size, dispersion, and metal oxidation state regardless of metal loading. The major difference between catalysts was the number of surface palladium sites which scaled directly with metal loading. Turnover frequency calculations showed that the intrinsic activity of each catalyst is the same, with conversion scaling with the number of active sites. The number of active sites was normalised experimentally with catalyst performance proving to be identical regardless of weight loading. This study shows that CVI is a technique that can produce active catalysts with high levels of control and consistency of active metal nanoparticles as a function of loading. The same level of control over dispersion and activity was not achieved when catalysts were prepared by conventional aqueous impregnation. The fundamental understanding of CVI is important for the design of highly active catalysts, which is exemplified for propane total oxidation, but has wider significance for other applications of supported metal catalysts.

Received 24th May 2024,
Accepted 21st July 2024

DOI: 10.1039/d4cy00665h

rsc.li/catalysis

Introduction

The method of catalyst synthesis can lead to significant differences in the properties and activity. It can influence many factors including nanoparticle size, distribution, shape and oxidation state.¹ Traditional impregnation and precipitation techniques are popular due to ease of synthesis but can demonstrate poor control over nanoparticle size, especially when metal loading is varied.^{2,3} Previous studies have suggested that the size distribution broadens, while dispersion decreases when metal weight loading increases. The precursor used can also introduce surface poisons, for example, chloride ions from metal chloride precursors, which remain on the surface after synthesis and have been found to be detrimental to catalytic activity. Colloidal methods like sol-immobilisation have been designed with the intention of greater particle size control through use of protecting groups.^{4,5}

Chemical vapour impregnation (CVI) is a preparation method that can negate some of the drawbacks of the more traditional impregnation techniques. CVI is a simple technique to use experimentally, metals are deposited onto a support using volatile metal precursors.⁶ Typically an acetylacetonate precursor is sublimed to deposit the complex onto the surface and then the ligands are removed *via* thermal decomposition to form metal nanoparticles. This makes it a much cleaner technique than traditional impregnation methods as there are less opportunities for impurities and poisons to be introduced. CVI is also a solventless technique allowing better compliance with the 12 principles of green chemistry through the reduction of waste production.⁷

Initial research into CVI has predominately focused on catalysts used for liquid phase reactions, demonstrating it can produce highly active catalysts for oxidation reactions due to a high level of control over nanoparticle size and distribution.^{8–10} Recently there has been an increase in study into CVI catalysts used for gas phase applications, with high performance catalysts being synthesised for CO₂ hydrogenation,^{11–18} ammonia decomposition,¹⁹ and NH₃-SCR reactions.²⁰ However, there has not been a more in-depth

Max Planck-Cardiff Centre on the Fundamentals of Heterogeneous Catalysis
FUNCAT, Cardiff Catalysis Institute, School of Chemistry, Cardiff University,
Translational Research Hub, Maindy Road, Cardiff CF24 4HQ, UK.

E-mail: baileyla2@cardiff.ac.uk, taylorsh@cardiff.ac.uk

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4cy00665h>

study into CVI as a synthesis technique investigating how factors like metal loading can influence activity.

Supported metal nanoparticle catalysts are particularly important for environmental pollution control. Volatile organic compounds (VOCs) are a major cause of air pollution, especially in urban areas,²¹ and have a significant negative impact on both the environment and human health.^{22,23} They are emitted from a wide range of anthropogenic sources, however, in urban environments they typically are emitted from automobiles due to the incomplete combustion of fuel.^{24,25} As air pollution is the leading cause of preventable death worldwide,²⁶ it is imperative that the emissions of VOCs are reduced, with wide ranging legislation enacted to achieve this aim.^{27–29} While there are several methods for removing VOCs, catalytic oxidation has proved effective due to lower operating cost, higher selectivity, and the ability to treat low concentration streams.^{30,31}

The total oxidation of propane has been extensively researched because it is a good model for VOC total oxidation.³² Many recent studies have focussed on metal oxide catalysts due to cheaper cost of production, with Co, Mn, Ce, and Fe based metal oxides all being exceptionally active.^{33–45} However, noble metal supported catalysts are favoured industrially due to greater activity and high durability.⁴⁶ Supported palladium on alumina catalysts have also been extensively studied for propane total oxidation.⁴⁷ Despite the depth of study there is still debate in the literature as to the most active oxidation state and palladium nanoparticle size. Although there is a consensus that PdO is the active form for propane oxidation.^{48–52}

This work aims to investigate in more detail CVI as a synthesis technique for supported palladium catalysts, using propane total oxidation as a model reaction. Catalyst activity is studied as a function of active metal loading and performance correlated with the structural properties of the catalysts to provide insight into the scope of CVI as an alternative preparation technique.

Experimental

Catalyst preparation

Preparation of Pd/Al₂O₃ catalysts by chemical vapour impregnation (CVI). Supported Pd/Al₂O₃ with different weight loadings (1.0, 2.5 and 5.0 wt%) were synthesised by chemical vapour impregnation (CVI) and wet impregnation (WI). The CVI procedure used for the preparation of the materials herein aligns with the procedure described previously by Forde *et al.*⁶ An appropriate amount of palladium acetylacetonate (Sigma Aldrich, 99%) was added to γ -Al₂O₃ (Sigma Aldrich, average particle size <50 nm) in a glass vial to give the desired weight loading and thoroughly mixed through shaking. The mixture was transferred to a 100 mL Schlenk flask with a magnetic stirrer bar, sealed and then evacuated to 10–3 mbar. The mixture was then heated to 140 °C for 1 hour while stirring. The resulting mixture was then calcined at 500 °C, at a ramp rate of 10 °C per minute, for 4 hours under static air, to give the desired catalyst.

Preparation of Pd/Al₂O₃ catalysts by wet impregnation (WI). Wet impregnation catalysts were prepared for comparison using a method described by Macino *et al.*⁵³ The required amount of Pd(NO₃)₂ solution (8.2 mg Pd mL⁻¹, Sigma Aldrich) was diluted to 16 mL using deionised water and heated to 60 °C while stirring. The appropriate amount of Al₂O₃ (Sigma Aldrich, average particle size <50 nm) was added and the mixture was heated to 95 °C and left for 16 hours to dry. The sample was then calcined at 500 °C at a ramp rate of 10 °C per minute, for 4 hours under static air to give the desired catalyst.

Catalyst characterisation

Microwave plasma atomic emission spectroscopy (MP-AES). MP-AES was performed on an Agilent 4100 MP-AES using Agilent MP expert software to calculate elemental concentration. A calibration plot was obtained by diluting standard metal solutions with deionised water. Samples (50 mg) were digested overnight in 1% aqua regia solution (50 mL).

Powder X-ray diffraction. (XRD) analysis was carried out on a PANalytical X'Pert Pro diffractometer using a Ge single crystal monochromated CuK α radiation source operated at 40 kV and 40 mA. Diffraction patterns were recorded with a step size of 0.0167° between 5–80° 2 θ using a backfilled sample holder over 40 minutes. Diffraction patterns were analysed by comparison with the International Centre for Diffraction Data (ICDD) standard Powder Diffraction File.

X-ray photoelectron spectroscopy (XPS). (XPS) was performed using a Kratos Axis ultra DLD photoelectron spectrometer, utilising AlK α radiation operating at an x-ray power of 144 W (12 mA \times 12 kV). Data was collected over an analysis area of approximately 700 \times 300 microns using the Hybrid spectroscopy mode. Pass energies of 160 eV and step size of 1 eV was used for survey scans and 40 eV and 0.1 eV step size for high-resolution scans. Sample charge neutralisation was achieved using low energy electrons and spectra were calibrated to the Al2p line of Al₂O₃, taken to be 74.5 eV. Analysis was performed using CasaXPS v2.3.24 (ref. 54) after removal of a Shirley type background and utilising modified Wagner sensitivity factors as supplied by the instrument manufacturer.

CO diffuse reflectance infrared spectroscopy (DRIFTS). (DRIFTS) experiments were performed on a Bruker Tensor 27 FTIR spectrometer equipped with a MCT detector cooled using liquid nitrogen. Spectra were recorded with 32 scans per spectrum with a 4 cm⁻¹ resolution. Samples were pretreated with N₂ for 30 minutes to remove surface adsorbates before being saturated with CO (2% CO/He) over 20 minutes. The system was then purged with N₂ to remove gas phase CO before data collection.

CO chemisorption. Experiments were carried out using a Micromeritics Autochem II 2920 analyser equipped with a TCD. The sample (*ca.* 100 mg) was fixed into a quartz U-tube reactor between quartz wool plugs. The sample was reduced at 400 °C with a heating ramp of 10 °C min⁻¹ under a 50 mL

min^{-1} flow of 10% H_2/Ar . Upon reaching 400 °C, the gas feed was switched to Ar and the sample cooled to the analysis temperature of 35 °C. The feed gas was subsequently changed to He (50 mL min^{-1}) with aliquots of 1% CO/He pulsed into the carrier gas at 3 minute intervals using a calibrated injection loop (5.55 mL). Pulses of 1% CO/He were continually added until the recorded peak area remained constant (three peaks with a difference in area of under 1%). For dispersion calculations a stoichiometry of 2 was used based on work by Canton *et al.*⁵⁵

BET surface area analysis. Surface area analysis was performed applying Brunauer–Emmett–Teller (BET) theory to a 20-point N_2 adsorption isotherm. Samples were initially degassed under vacuum at 120 °C for 16 hours. The isotherm was recorded using a Quantachrome Quadrasorb apparatus over a range of $0.05\text{--}0.35p/p_0$ at -196 °C .

Transmission electron microscopy (TEM). Imaging was performed on a JEOL JEM-2100 operating at 200 kV. Samples were prepared by dry dispersion on 300 mesh Cu grids coated with holey carbon film. Particle size analysis was performed through manual measurement of 200 nanoparticles.

Aberration Corrected Scanning Transmission Electron Microscopy (AC-STEM). Imaging was performed on a probe corrected Thermo Fisher Scientific Spectra 200 operating at 200 kV. Samples were prepared by dry dispersion onto 300 mesh copper grids coated with holey carbon films. A semi-angle of 30 mrad was used with a probe current between 50–100 pA.

Catalyst activity evaluation

All reactions were performed at atmospheric pressure using a fixed bed continuous flow microreactor. 0.3 g of the catalyst was packed between plugs of quartz wool in a stainless-steel tube (1 cm i.d.). Catalysts were pre-treated under a flow of N_2 (250 mL min^{-1}) at 200 °C for 1 hour. A pre-made gas mix of 1000 ppm propane/10% O_2/N_2 (BOC specialty gas) was used at a flow rate of 250 mL min^{-1} to give a GHSV of $50\,000 \text{ h}^{-1}$. Reactions were performed between 200–500 °C and were monitored using an in-line Fourier transform infrared (FTIR) process gas analyser (Gasmeter) and gas concentrations calculated using Calcmet software. Concentration readings were taken at 25 °C intervals once the temperature stabilised with readings repeated until it was certain steady state had been reached.

Results and discussion

Catalytic activity of Pd/ Al_2O_3 catalysts synthesised by CVI

To begin the investigation, blank reactions with no catalyst and the alumina support material were conducted. No propane conversion was observed with an empty reactor tube over the temperature range studied, and only 22% propane conversion was exhibited over the alumina support at 500 °C. This shows that homogeneous gas phase reactions in the reactor tube were negligible for propane conversion and that the contribution of alumina alone was relatively low compared to the Pd-containing catalysts. The performance of

the 1.0 wt%, 2.5 wt% and 5.0 wt% Pd/ Al_2O_3 catalysts synthesised by CVI for propane oxidation are shown in Fig. 1. All the CVI catalysts were active, with full conversion achieved at 450 °C for the 1.0 wt% catalyst, and 400 °C for the 2.5 wt% and 5.0 wt% catalysts. Catalyst performance increased with Pd loading, with the 5.0 wt% catalyst the most active, followed by the 2.5 wt%, then the 1 wt% Pd/ Al_2O_3 catalyst. This is demonstrated by the T_{20} and T_{50} values (Table 1). However, when the activity of the catalyst is normalised to the mass of palladium present in each reaction, very similar rates of propane conversion are seen, suggesting similar intrinsic activity of Pd sites. All catalysts showed high selectivity for carbon dioxide; in all cases over 99% conversion to CO_2 was observed, indicating that these catalysts are very selective for the deep oxidation of propane. The catalysts synthesised here were also found to be more active than analogous materials synthesised by the more traditional method of wet impregnation (Fig. S1 and Table S1†) suggesting that CVI is a superior method of making catalysts for this application.

Catalyst stability was also probed with 5 cycles of catalyst testing performed on each sample, along with 50 hours time on line at 275 °C (Fig. S2 and S3†). All cycles line up suggesting stability over multiple tests, while less than a 5% drop off in activity was noted over the 50 hours of testing. These tests show limited change in catalyst activity for all materials demonstrating high stability for catalysts synthesised by CVI.

Catalyst characterisation

MP-AES was performed on the samples and can be seen in Table 2. This demonstrated that the target loading palladium

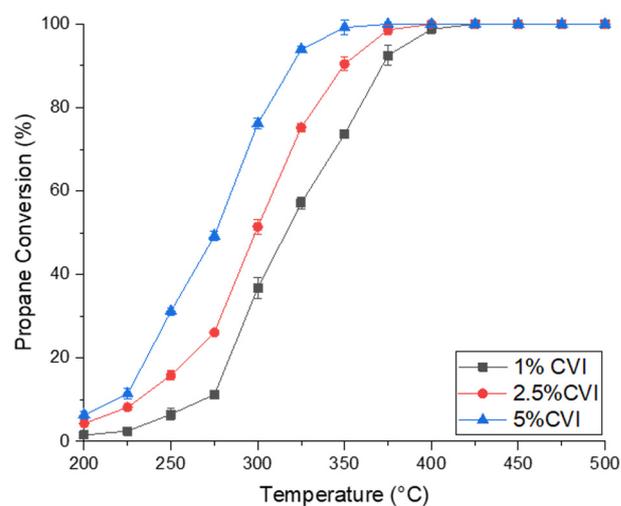


Fig. 1 Propane conversion as a function of temperature for the Pd/ Al_2O_3 catalysts synthesised by chemical vapour impregnation. (Black square) 1.0 wt% Pd/ Al_2O_3 , (red circle) 2.5 wt% Pd/ Al_2O_3 , (blue triangle) 5.0 wt% Pd/ Al_2O_3 . Reaction conditions: 1000 ppm propane/10% O_2/N_2 , gas hourly space velocity (GHSV) $50\,000 \text{ h}^{-1}$, temperature range 200–500 °C.

Table 1 Activity data for the Pd/Al₂O₃ catalysts synthesised by chemical vapour impregnation. T_{20} = temperature at which 20% conversion was achieved. T_{50} = temperature at which 50% conversion was achieved. Reaction conditions: 1000 ppm propane/10% O₂/N₂, gas hourly space velocity (GHSV) 50 000 h⁻¹, temperature range 200–500 °C

Catalyst	T_{20} (°C)	T_{50} (°C)	Activity at 250 °C (mol _{C₃H₈} g _{Pd} ⁻¹ s ⁻¹)
1.0 wt% Pd/Al ₂ O ₃	283	316	4.03×10^{-6} ($\pm 4.08 \times 10^{-8}$)
2.5 wt% Pd/Al ₂ O ₃	260	298	3.92×10^{-6} ($\pm 4.32 \times 10^{-8}$)
5.0 wt% Pd/Al ₂ O ₃	236	276	3.93×10^{-6} ($\pm 3.00 \times 10^{-8}$)

loading and actual palladium loading is very similar, identifying CVI as accurate technique for producing supported palladium catalysts.

Transmission electron microscopy was used to gain insight on the size and distribution of the supported Pd particles in the materials (Fig. 2a–c). All the catalysts had distinct palladium nanoparticles of a consistent size with an average nanoparticle size of 4.0 nm, 4.5 nm, and 4.5 nm measured for the 1.0 wt% CVI, 2.5 wt% CVI, and 5.0 wt% CVI catalysts, respectively. Consistent control over the nanoparticle size distribution was also noted with a diameter range of 2–8 nm for all CVI catalysts. Size distribution plots (Fig. S4†) shows a normal distribution of nanoparticles ranging 3–5 nm with few outliers closer to 2 nm and 8 nm. High resolution AC-STEM imaging was also performed for all three of the catalysts prepared by CVI (Fig. 2d–f). Analysis showed the alumina support was decorated with palladium nanoparticles, and also clusters under 1 nm in diameter. These clusters formed on all CVI catalysts irrespective of loading, but were more prevalent on the 1% CVI catalyst. This imaging suggests that the size and distribution of palladium nanoparticles are largely independent of weight loading and the particles formed are very consistent.

The physicochemical properties of the samples were probed by XRD and BET analysis of N₂ physisorption. All catalysts exhibited major peaks at $2\theta = 37.1^\circ$, 39.8° , 46.0° and 67.0° corresponding to (311), (222), (400), and (440) planes of γ -alumina, respectively (Fig. 3 and Table 2).⁵⁶ Additionally, a peak at $2\theta = 34^\circ$ was observed, corresponding to PdO (101)/(002) planes for all of the catalysts. This peak increases in intensity as the weight loading increases, suggesting that more PdO was present in the 5.0 wt% Pd catalyst. This observation can be important for rationalising the increase in performance with weight loading for the CVI catalysts, as PdO is considered to be the active species for propane total oxidation.⁴⁸ None of the samples show scattering for metallic

palladium, suggesting that the metal was solely in its oxide form which is expected given heat treatment under oxidising conditions. Overall, there was minimal difference in the scattering pattern besides changes in intensity between weight loadings suggesting that differences in the large-scale catalyst structure was negligible. Analysis of the surface area showed very little difference between catalysts, suggesting little impact on the wider catalyst structure with loading.

X-Ray photoelectron spectroscopy (XPS) analysis of the catalysts (Pd 3d, Fig. 4) revealed that Pd was predominantly in the 2+ oxidation state (Table S2†), which correlates with the XRD analysis. It should be noted that with the 1.0 wt% CVI sample some metallic Pd was observed, but is likely to be attributed to the photoreduction of PdO to Pd(0) or smaller particles during the analysis. These findings align with the literature, where it has been reported that PdO is more active than metallic Pd for propane oxidation, it implies that differences in the chemical state of the metal is not the reason for differences in activity. The binding energy of the 1% CVI catalyst is interesting as it increased compared to the other catalysts, with a value of 337.3 eV. This is a shift of 0.5 eV from the 336.8 eV expected from bulk PdO. Jürgensen *et al.* suggested an increase of 0.6 eV in the palladium binding energy is indicative of the presence of very small nanoparticles.⁵⁷ Thus, the XPS for the 1% catalyst indicates that it could have smaller palladium particles than the other catalysts, which corresponds to the increased number of small clusters identified in the AC-STEM imaging, an observation which is supported by the change in FWHM of the Pd 3d peaks with increasing metal loading.⁵⁸ XPS was also used to probe the surface concentration of the palladium (Table S2†). Unsurprisingly, surface concentration of palladium increases with weight loading which can explain the differences in conversion identified. The surface concentration scales in a linear fashion suggesting that the size distribution of the metal nanoparticles remains consistent, generally agreeing with the microscopy obtained (Fig. S5†).

CO DRIFTS was performed on the samples to probe the palladium sites present. The spectra recorded for the CVI catalysts are shown in Fig. 5. For all three samples there are three peaks at 1926 cm⁻¹, 1975 cm⁻¹, and 2088 cm⁻¹, corresponding to linear CO bonding of Pd⁺, Pd⁰, and bridged CO bonding on Pd⁰ respectively,⁵⁹ with the intensity of each peak increasing with weight loading. The peak for the linear Pd⁰-CO stretch is largely obscured by the linear Pd⁺-CO peak and appears as a shoulder. There are no peaks in the 2180–2160 cm⁻¹ region where a PdO-CO stretching vibration is

Table 2 Physicochemical properties of the Pd/Al₂O₃ CVI catalysts from TEM, MP-AES and BET analysis

Sample	Particle size distribution ^a (nm)	Average nanoparticle size ^a (nm)	Pd weight loading ^b (%)	BET (m ² g ⁻¹)
1.0 wt% Pd/Al ₂ O ₃	2.0–8.0	4.0	0.98	128
2.5 wt% Pd/Al ₂ O ₃	2.0–8.0	4.5	2.51	126
5.0 wt% Pd/Al ₂ O ₃	2.0–8.0	4.5	4.95	124

^a Calculated from TEM imaging. Average of 200 nanoparticles. ^b Calculated using MP-AES.

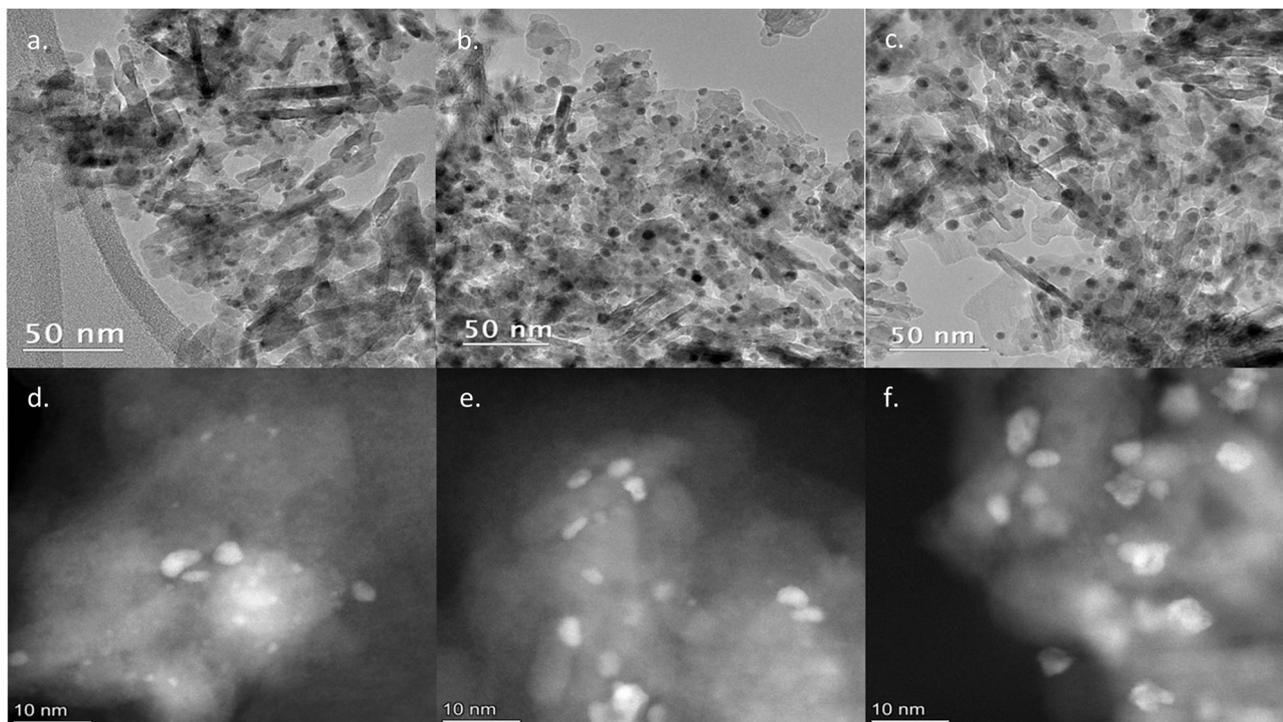


Fig. 2 Electron microscopy micrographs of the catalysts prepared by CVI. a–c: TEM images (50k \times magnification), d–f: AC-STEM images (80k \times magnification). a and d: 1.0 wt% CVI, b and e: 2.5 wt% CVI, c and f: 5.0 wt% CVI.

typically expected, showing that the surface palladium is easily reducible. This has been identified for other palladium-based catalysts in the literature as evidence of easy cycling between Pd oxidation states helping to explain improved performance.^{60,61} This is further evidence that the structure of the nanoparticles formed by CVI are very similar in nature irrespective of the metal loading across the range 1.0–5.0 wt%.

CO chemisorption was performed on the catalysts with results shown in Table 3. The concentration of CO adsorbed increases in a linear relationship with the palladium weight loading. Accordingly, the number of palladium sites present

on the catalyst surface can be measured, the number increasing linearly with weight loading. These measurements agree with the TEM analysis and help explain the increase in activity with increased Pd loading. As palladium is the active site for this reaction, having more surface sites expectedly enhances conversion. CO chemisorption was also used to calculate the dispersion of the palladium for each catalyst. The dispersion for all CVI catalysts was similar, at *ca.* 30%. This is again consistent with other complimentary characterisation data for these catalysts, suggesting that

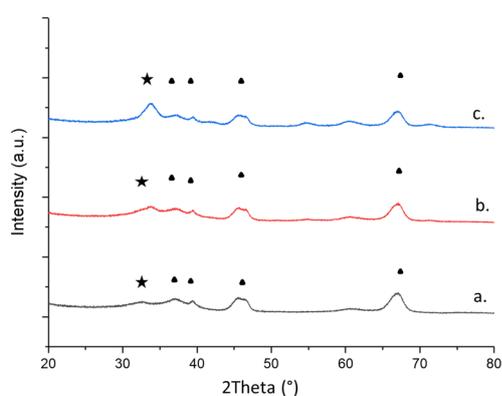


Fig. 3 XRD patterns of (a.) 1.0 wt% Pd/Al₂O₃, (b.) 2.5 wt% Pd/Al₂O₃ and (c.) 5.0 wt% Pd/Al₂O₃ catalysts. (Black star) PdO, JCPDS no. 00-006-0515. (Black triangle) γ -Al₂O₃, JCPDS no. 10-0425.

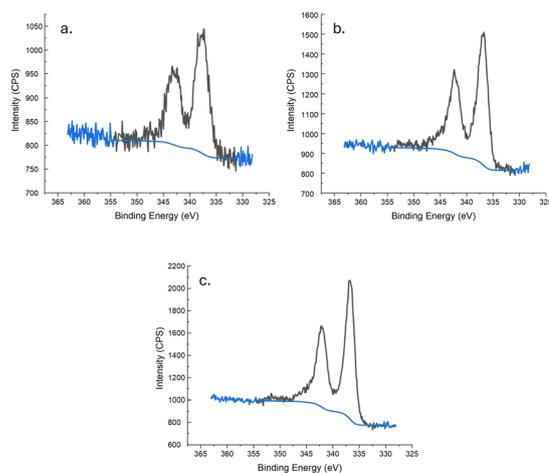


Fig. 4 Pd 3d core-level XPS spectra for (a.) 1.0 wt% Pd/Al₂O₃, (b.) 2.5 wt% Pd/Al₂O₃ and (c.) 5.0 wt% Pd/Al₂O₃ catalysts.

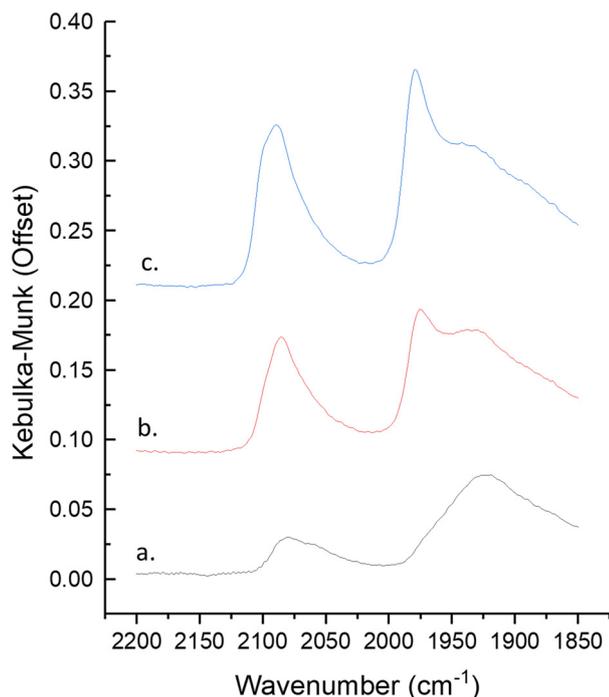


Fig. 5 DRIFT spectra of CO adsorbed at room temperature for (a.) 1.0 wt% Pd/Al₂O₃, (b.) 2.5 wt% Pd/Al₂O₃ and (c.) 5.0 wt% Pd/Al₂O₃ catalysts.

catalysts with consistent nanoparticle structure are formed *via* CVI regardless of weight loading.

Calculation of the catalytic turnover frequency (TOF) provides a measure of the intrinsic activity per active site (Table 3). The measured TOFs for the CVI catalysts showed consistent values regardless of weight loading with values of $1.43 \times 10^{-3} \text{ s}^{-1}$, $1.49 \times 10^{-3} \text{ s}^{-1}$ and $1.48 \times 10^{-3} \text{ s}^{-1}$ for the 1.0 wt%, 2.5 wt%, and 5.0 wt% CVI catalysts respectively. This agrees with the similar rates calculated (Table 1) where activity was normalised to grams of palladium present in the reaction, and shows that the intrinsic activity per site is the same regardless of Pd loading when the CVI method is employed.

Post reaction XRD and CO chemisorption (Fig. S6 and Table S3†) were performed on all samples after 50 hours time on line with little change noted from the pre-reaction samples. No change was seen on the x-ray diffraction patterns while a very small decrease in nanoparticle dispersion was noted for the 5 wt% Pd/Al₂O₃ sample which can be explained by error of the technique. This links well with the stability testing performed which demonstrated no difference in catalyst activity with

extended use suggesting that these catalysts are stable over prolonged use.

Assessing intrinsic palladium site activity

To probe this effect further, the mass, and hence volume, of catalyst used in a reaction for total propane oxidation was varied to ensure the same number of palladium sites were present in the catalyst bed, based on the number of sites measured by CO chemisorption. This was performed for the CVI catalysts and also for comparison with analogous catalysts synthesised by wet impregnation (WI) (Fig. 6). For the CVI catalysts, propane conversion was identical at each temperature point and activity curves were superimposable within experimental error. This shows that activity for these catalysts are dominated by the number of surface palladium active sites with the intrinsic activity for each site being the same. It further identifies CVI as a technique for producing highly uniform materials regardless of metal loading, allowing a scalable method to prepare catalysts with varying numbers of surface sites without changing the characteristics of the nanoparticles. This trend was not identified for the WI catalysts, where large differences in propane total oxidation activity occurs as a function of metal loading. Activity decreases with weight loading suggesting that several factors are influencing performance. Previous studies have suggested that impregnation synthesis methods have poor control over factors like dispersion and nanoparticle size distribution, with these decreasing and increasing with increased weight loading respectively.^{2,3} Characterisation performed on the wet impregnation catalysts (Tables S4 and S5, and Fig. S7–S10†) show large differences in nanoparticle structure and characteristics suggesting that the high uniformity of the CVI prepared samples is characteristic of the technique.

Conclusions

Chemical vapour impregnation is a relatively unstudied catalyst synthesis technique, but one that can be readily employed in the laboratory with simple and inexpensive equipment. Pd/Al₂O₃ catalysts with varying weight loading were prepared by this method and studied for the total oxidation of propane. In all cases the CVI catalysts were highly active with activity ascribed to high nanoparticle dispersion and the formation of many active sites. Characterisation showed control over uniformity in nanoparticle structure and composition with activity per active site equivalent, and activity scaling linearly with metal loading which scaled with the number of palladium sites.

Table 3 Physical properties and activity characteristics of Pd/Al₂O₃ catalysts extracted from and based on CO chemisorption characterisation

Sample	CO adsorbed (mmol g ⁻¹)	Palladium surface sites ^a (g ⁻¹)	Dispersion ^a (%)	TOF ^b (10 ⁻³)(s ⁻¹)
1.0 wt% Pd/Al ₂ O ₃	1.73×10^{-2}	2.08×10^{19}	30	1.43
2.5 wt% Pd/Al ₂ O ₃	3.28×10^{-2}	3.95×10^{19}	28	1.49
5.0 wt% Pd/Al ₂ O ₃	6.68×10^{-2}	8.05×10^{19}	28	1.48

^a Calculated from CO chemisorption assuming a stoichiometry of 2. ^b Calculated at 250 °C.

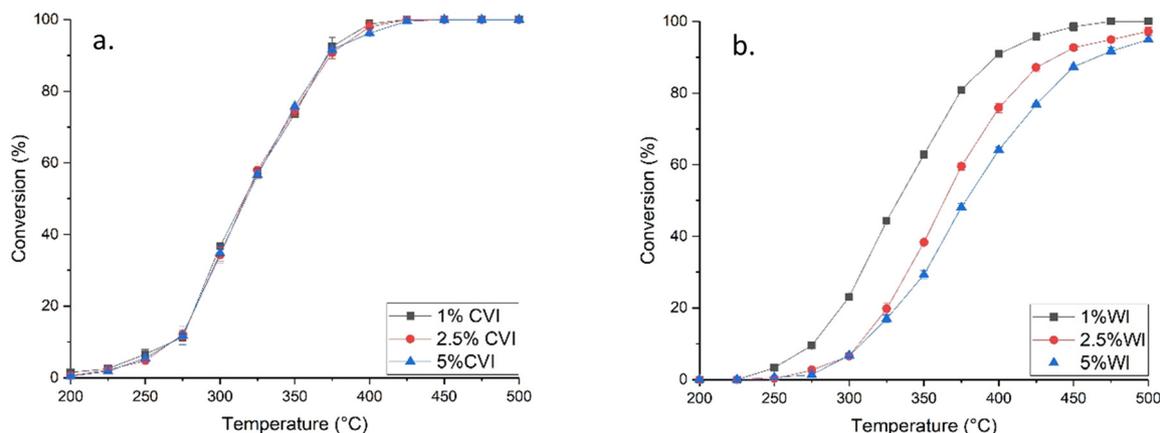


Fig. 6 Propane conversion as a function of temperature for Pd/Al₂O₃ catalysts synthesised by a. CVI and b. WI when the volume of sample was varied to normalise activity to Pd sites. (Black square) 1.0 wt% Pd/Al₂O₃, (red circle) 2.5 wt% Pd/Al₂O₃, (blue triangle) 5.0 wt% Pd/Al₂O₃. Reaction conditions: 1000 ppm propane/10% O₂/N₂, temperature range 200–500 °C.

This was studied experimentally by normalising the sample volume to ensure the same number of palladium sites were present. Near identical performance was noted, that was not seen for analogous samples prepared *via* wet impregnation exploitation of these controllable properties that are accessible by CVI provides an important tool in the process of catalyst design. The high level of control over particle size and distribution is very desirable, though additional study is needed to see whether the consistent average nanoparticle size of 4 nm is characteristic of this technique, or if the method can be optimised further to give a range of average nanoparticle size.

Data availability

All relevant data is contained within the manuscript and ESI.†

Author contributions

The manuscript was written through contributions of all authors. L. A. B. synthesised catalysts, conducted experiments and data analysis. L. A. B., M. D., T. E. D., and D. J. M. conducted material characterisation and corresponding data processing. L. A. B. and S. H. T. contributed to the design of study and provided technical advice and result interpretation.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

XPS data collection was performed at the EPSRC National Facility for XPS (“HarwellXPS”), operated by Cardiff University and UCL, under Contract No. PR16195. The authors would like to thank the CCI-Electron Microscopy Facility which has been partially funded by the European Regional Development

Fund through the Welsh Government and The Wolfson Foundation. All authors gratefully acknowledge Cardiff University and the Max Planck Centre for Fundamental Heterogeneous Catalysis (FUNCAT) for financial support.

Notes and references

- 1 J. A. Schwarz, C. Contescu and A. Contescu, *Chem. Rev.*, 1995, **95**, 477–510.
- 2 Y. J. Huang and J. A. Schwarz, *Appl. Catal.*, 1987, **30**, 239–253.
- 3 B. A. T. Mehrabadi, S. Eskandari, U. Khan, R. D. White and J. R. Regalbuto, in *Advances in Catalysis*, ed. C. Song, Academic Press, 2017, vol. 61, pp. 1–35.
- 4 A. Villa, D. Wang, G. M. Veith, F. Vindigni and L. Prati, *Catal. Sci. Technol.*, 2013, **3**, 3036–3041.
- 5 S. Campisi, D. Ferri, A. Villa, W. Wang, D. Wang, O. Kröcher and L. Prati, *J. Phys. Chem. C*, 2016, **120**, 14027–14033.
- 6 M. M. Forde, R. D. Armstrong, C. Hammond, Q. He, R. L. Jenkins, S. A. Kondrat, N. Dimitratos, J. A. Lopez-Sanchez, S. H. Taylor, D. Willock, C. J. Kiely and G. J. Hutchings, *J. Am. Chem. Soc.*, 2013, **135**, 11087–11099.
- 7 P. T. Anastas and J. C. Warner, *Green Chemistry: Theory and Practice*, Oxford University Press, 1998.
- 8 M. M. Forde, R. D. Armstrong, R. McVicker, P. P. Wells, N. Dimitratos, Q. He, L. Lu, R. L. Jenkins, C. Hammond, J. A. Lopez-Sanchez, C. J. Kiely and G. J. Hutchings, *Chem. Sci.*, 2014, **5**, 3603–3616.
- 9 M. M. Forde, L. Kesavan, M. I. bin Saiman, Q. He, N. Dimitratos, J. A. Lopez-Sanchez, R. L. Jenkins, S. H. Taylor, C. J. Kiely and G. J. Hutchings, *ACS Nano*, 2014, **8**, 957–969.
- 10 R. Su, M. M. Forde, Q. He, Y. Shen, X. Wang, N. Dimitratos, S. Wendt, Y. Huang, B. B. Iversen, C. J. Kiely, F. Besenbacher and G. J. Hutchings, *Dalton Trans.*, 2014, **43**, 14976–14982.
- 11 H. Bahruji, M. Bowker, W. Jones, J. Hayward, J. R. Esquivos, D. J. Morgan and G. J. Hutchings, *Faraday Discuss.*, 2017, **197**, 309–324.

- 12 M. Bowker, N. Lawes, I. Gow, J. Hayward, J. R. Esquiús, N. Richards, L. R. Smith, T. J. A. Slater, T. E. Davies, N. F. Dummer, L. Kabalan, A. Logsdail, R. C. Catlow, S. Taylor and G. J. Hutchings, *ACS Catal.*, 2022, **12**, 5371–5379.
- 13 N. Lawes, K. J. Aggett, L. R. Smith, T. J. A. Slater, M. Dearg, D. J. Morgan, N. F. Dummer, S. H. Taylor, G. J. Hutchings and M. Bowker, *Catal. Lett.*, 2024, **154**, 1603–1610.
- 14 N. Lawes, N. F. Dummer, S. Fagan, O. Wielgosz, I. E. Gow, L. R. Smith, T. J. A. Slater, T. E. Davies, K. J. Aggett, D. J. Morgan, S. H. Taylor, G. J. Hutchings and M. Bowker, *Appl. Catal., A*, 2024, **679**, 119735.
- 15 H. Bahruji, J. R. Esquiús, M. Bowker, G. Hutchings, R. D. Armstrong and W. Jones, *Top. Catal.*, 2018, **61**, 144–153.
- 16 H. Bahruji, R. D. Armstrong, J. Ruiz Esquiús, W. Jones, M. Bowker and G. J. Hutchings, *Ind. Eng. Chem. Res.*, 2018, **57**, 6821–6829.
- 17 J. Ruiz Esquiús, H. Bahruji, S. H. Taylor, M. Bowker and G. J. Hutchings, *ChemCatChem*, 2020, **12**, 6024–6032.
- 18 N. Lawes, I. E. Gow, L. R. Smith, K. J. Aggett, J. S. Hayward, L. Kabalan, A. J. Logsdail, T. J. A. Slater, M. Dearg, D. J. Morgan, N. F. Dummer, S. H. Taylor, M. Bowker, C. R. A. Catlow and G. J. Hutchings, *Faraday Discuss.*, 2023, **242**, 193–211.
- 19 L. A. Parker, N. Richards, L. Bailey, J. H. Carter, E. Nowicka, S. Patisson, N. F. Dummer, Q. He, L. Lu, C. J. Kiely, S. E. Golunski, A. Roldan and G. J. Hutchings, *Catal. Lett.*, 2024, **154**, 1958–1969.
- 20 A. Cooper, T. E. Davies, D. J. Morgan, S. Golunski and S. H. Taylor, *Catalysts*, 2020, **10**, 294.
- 21 C. He, J. Cheng, X. Zhang, M. Douthwaite, S. Patisson and Z. Hao, *Chem. Rev.*, 2019, **119**, 4471–4568.
- 22 Y. Guo, M. Wen, G. Li and T. An, *Appl. Catal., B*, 2021, **281**, 119447.
- 23 A. Evuti, *Int. J. Eng. Sci.*, 2013, **2**, 145–153.
- 24 B. Barletta, S. Meinardi, F. Sherwood Rowland, C.-Y. Chan, X. Wang, S. Zou, L. Yin Chan and D. R. Blake, *Atmos. Environ.*, 2005, **39**, 5979–5990.
- 25 H. Huo, Q. Zhang, K. He, Z. Yao, X. Wang, B. Zheng, D. G. Streets, Q. Wang and Y. Ding, *Environ. Pollut.*, 2011, **159**, 2954–2960.
- 26 P. J. Landrigan, R. Fuller, N. J. R. Acosta, O. Adeyi, R. Arnold, N. Nil Basu, A. B. Baldé, R. Bertollini, S. Bose-O'Reilly, J. I. Boufford, P. N. Breyse, T. Chiles, C. Mahidol, A. M. Coll-Seck, M. L. Cropper, J. Fobil, V. Fuster, M. Greenstone, A. Haines, D. Hanrahan, D. Hunter, M. Khare, A. Krupnick, B. Lanphear, B. Lohani, K. Martin, K. V. Mathiasen, M. A. McTeer, C. J. L. Murray, J. D. Ndahimananjara, F. Perera, J. Potočnik, A. S. Preker, J. Ramesh, J. Rockström, C. Salinas, L. D. Samson, K. Sandilya, P. D. Sly, K. R. Smith, A. Steiner, R. B. Stewart, W. A. Suk, O. C. P. van Schayck, G. N. Yadama, K. Yumkella and M. Zhong, *Lancet*, 2018, **391**, 462–512.
- 27 S. Scirè and L. F. Liotta, *Appl. Catal., B*, 2012, **125**, 222–246.
- 28 M. V. Twigg, *Appl. Catal., B*, 2007, **70**, 2–15.
- 29 M. S. Kamal, S. A. Razzak and M. M. Hossain, *Atmos. Environ.*, 2016, **140**, 117–134.
- 30 A. Krishnamurthy, B. Adebayo, T. Gelles, A. Rownaghi and F. Rezaei, *Catal. Today*, 2020, **350**, 100–119.
- 31 M. N. Taylor, W. Zhou, T. Garcia, B. Solsona, A. F. Carley, C. J. Kiely and S. H. Taylor, *J. Catal.*, 2012, **285**, 103–114.
- 32 B. A. Tichenor and M. A. Palazzolo, *Environ. Prog.*, 1987, **6**, 172–176.
- 33 P. M. Shah, A. N. Day, T. E. Davies, D. J. Morgan and S. H. Taylor, *Appl. Catal., B*, 2019, **253**, 331–340.
- 34 B. Solsona, T. E. Davies, T. Garcia, I. Vázquez, A. Dejoz and S. H. Taylor, *Appl. Catal., B*, 2008, **84**, 176–184.
- 35 B. Solsona, T. García, G. J. Hutchings, S. H. Taylor and M. Makkee, *Appl. Catal., A*, 2009, **365**, 222–230.
- 36 B. Solsona, T. García, R. Sanchis, M. D. Soriano, M. Moreno, E. Rodríguez-Castellón, S. Agouram, A. Dejoz and J. M. López Nieto, *Chem. Eng. J.*, 2016, **290**, 273–281.
- 37 B. Solsona, R. Sanchis, A. M. Dejoz, T. García, L. Ruiz-Rodríguez, J. M. López Nieto, J. A. Cecilia and E. Rodríguez-Castellón, *Catalysts*, 2017, **7**, 96.
- 38 B. Solsona, T. Garcia, E. Aylón, A. M. Dejoz, I. Vázquez, S. Agouram, T. E. Davies and S. H. Taylor, *Chem. Eng. J.*, 2011, **175**, 271–278.
- 39 P. M. Shah, J. W. H. Burnett, D. J. Morgan, T. E. Davies and S. H. Taylor, *Catalysts*, 2019, **9**, 475.
- 40 P. M. Shah, L. A. Bailey and S. H. Taylor, *Catalysts*, 2023, **13**, 114.
- 41 P. M. Shah, L. A. Bailey, D. J. Morgan and S. H. Taylor, *Catalysts*, 2023, **13**, 794.
- 42 G. Salek, P. Alphonse, P. Dufour, S. Guillemet-Fritsch and C. Tenailleau, *Appl. Catal., B*, 2014, **147**, 1–7.
- 43 Z. Zhu, G. Lu, Z. Zhang, Y. Guo, Y. Guo and Y. Wang, *ACS Catal.*, 2013, **3**, 1154–1164.
- 44 Y. Xie, Y. Yu, X. Gong, Y. Guo, Y. Guo, Y. Wang and G. Lu, *CrystEngComm*, 2015, **17**, 3005–3014.
- 45 K. Aggett, T. E. Davies, D. J. Morgan, D. Hewes and S. H. Taylor, *Catalysts*, 2021, **11**, 1461.
- 46 M. Taylor, E. N. Ndifor, T. Garcia, B. Solsona, A. F. Carley and S. H. Taylor, *Appl. Catal., A*, 2008, **350**, 63–70.
- 47 M. V. Twigg, *Catal. Today*, 2011, **163**, 33–41.
- 48 A. K. Khudorozhkov, I. A. Chetyrin, A. V. Bukhtiyarov, I. P. Prosvirin and V. I. Bukhtiyarov, *Top. Catal.*, 2017, **60**, 190–197.
- 49 Y. Yazawa, H. Yoshida, N. Takagi, S. Komai, A. Satsuma and T. Hattori, *Appl. Catal., B*, 1998, **19**, 261–266.
- 50 R. J. Farrauto, M. C. Hobson, T. Kennelly and E. M. Waterman, *Appl. Catal., A*, 1992, **81**, 227–237.
- 51 F. H. Ribeiro, M. Chow and R. A. Dallabetta, *J. Catal.*, 1994, **146**, 537–544.
- 52 R. F. Hicks, H. Qi, M. L. Young and R. G. Lee, *J. Catal.*, 1990, **122**, 295–306.
- 53 M. Macino, A. J. Barnes, S. M. Althabban, R. Qu, E. K. Gibson, D. J. Morgan, S. J. Freakley, N. Dimitratos, C. J. Kiely, X. Gao, A. M. Beale, D. Bethell, Q. He, M. Sankar and G. J. Hutchings, *Nat. Catal.*, 2019, **2**, 873–881.
- 54 N. Fairley, V. Fernandez, M. Richard-Plouet, C. Guillot-Deudon, J. Walton, E. Smith, D. Flahaut, M. Greiner, M.

- Biesinger, S. Tougaard, D. Morgan and J. Baltrusaitis, *Appl. Surf. Sci. Adv.*, 2021, **5**, 100112.
- 55 P. Canton, G. Fagherazzi, M. Battagliarin, F. Menegazzo, F. Pinna and N. Pernicone, *Langmuir*, 2002, **18**, 6530–6535.
- 56 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.
- 57 A. Jürgensen, N. Heutz, H. Raschke, K. Merz and R. Hergenröder, *Anal. Chem.*, 2015, **87**, 7848–7856.
- 58 I. Aruna, B. R. Mehta, L. K. Malhotra and S. M. Shivaprasad, *J. Appl. Phys.*, 2008, **104**, 064308.
- 59 Y. Zhang, Y. Cai, Y. Guo, H. Wang, L. Wang, Y. Lou, Y. Guo, G. Lu and Y. Wang, *Catal. Sci. Technol.*, 2014, **4**, 3973–3980.
- 60 E. J. Jang, J. Lee, D. G. Oh and J. H. Kwak, *ACS Catal.*, 2021, **11**, 5894–5905.
- 61 L. Meng, A.-P. Jia, J.-Q. Lu, L.-F. Luo, W.-X. Huang and M.-F. Luo, *J. Phys. Chem. C*, 2011, **115**, 19789–19796.