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Blurring the boundary between homogenous and heterogeneous catalysis using palladium nanoclusters with dynamic surfaces

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Using a magnetron sputtering approach that allows size-controlled formation of nanoclusters, we have created palladium nanoclusters that combine the features of both heterogeneous and homogeneous catalysts. Here we report the atomic structures and electronic environments of a series of metal nanoclusters in ionic liquids at different stages of formation, leading to the discovery of Pd nanoclusters with a core of *ca.* 2 nm surrounded by a diffuse dynamic shell of atoms in [C₄C₁Im][NTf₂]. Comparison of the catalytic activity of Pd nanoclusters in alkene cyclopropanation reveals that the atomically dynamic surface is critically important, increasing the activity by a factor of *ca.* 2 when compared to compact nanoclusters of similar size. Catalyst poisoning tests using mercury and dibenzo[*a,e*]cyclooctene show that dynamic Pd nanoclusters maintain their catalytic activity, which demonstrate their combined features of homogeneous and heterogeneous catalysts within the same material. Additionally, kinetic studies of cyclopropanation of alkenes mediated by the dynamic Pd nanoclusters reveal an observed catalyst order of 1, underpinning the pseudo-homogeneous character of the dynamic Pd nanoclusters.

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Catalysts are used in nearly 80 % of industrial processes¹, demanding the maximum catalytic efficiency of rare elements, such as Au, Pt and Pd^{2–4}. Increasing the metal active surface area by utilising catalysts in the form of nanoclusters is one of the most powerful approaches^{5–9}. When the size of metal shrinks down to the nanoscale, the structure and dynamics and thus properties of the nanocluster cannot be interpreted as a linear function of its size, which makes it challenging to correlate them with their catalytic performance^{7,10–12}. In order to establish a structure–property relationship for nanoclusters and the link with their catalytic performance, we must explore the series of related nanoclusters at their boundaries between discrete atoms and crystalline nanoclusters, with atomic precision, and using a set of complementary analytical methods, thus shedding light on their nature¹³.

In this context, magnetron sputtering offers exciting new opportunities for the manufacture of novel catalytic materials with controlled sizes and shapes of metal nanoclusters on a variety of supports^{14–18}. Crucially, this approach generates highly active clean surfaces, since no chemical stabilisers and/or solvents are involved in the process^{16,19}. Among the supports, ionic liquids (ILs) are unique because they offer a soft, liquid environment for relatively free (in comparison to solid supports) nanoclusters to be studied in their native state. Furthermore, ILs possess low vapour pressure allowing their use under the high vacuum conditions of magnetron sputtering^{20,21}. However, atomic-scale analysis of metal nanoclusters in IL, for example, by transmission electron microscopy (TEM), remains a challenge. The current sample preparation approaches using stabilisers or solvents^{22–24} may alter the environment surrounding the nanocluster, thus potentially inducing structural changes to the nanocluster and therefore hide important information on its shape and size. To the best of our knowledge, metal nanoclusters in ILs deposited by magnetron sputtering have not been observed with atomic resolution in their native state so far. As a result, the relationship between metal nanoclusters structure and their catalytic activity in ILs remains largely underexplored and widely debated.

Herein we determine the atomic structures and electronic environments of a series of metal nanoclusters in ILs at different stages of formation, leading to a discovery of Pd nanoclusters with a core of ca. 2 nm surrounded by a diffuse dynamic shell of atoms in 1-butyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide ($[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$) that combines the features of homogeneous and heterogeneous catalysts within the same material.

Results and discussion

Synthesis of Pd nanoclusters. The magnetron sputtering metal deposition process involves the elastic collision of argon ions with a highly pure metallic target, resulting in the expulsion of atoms or clusters from the target that are deposited onto a support material (Fig. 1a and Supplementary Fig. 1)¹⁴. Previously, it was proposed that the growth of metal clusters occurs on the IL surface/near surface or in the bulk IL depending on the magnetron sputtering conditions and physical properties of the employed IL^{25–27}. However, a variety of parameters, including the applied potential, argon (Ar) working pressure and work distance (distance between the metal target and the support), can influence: (i) whether metal atoms or a few metal atom cluster are ejected from the target; (ii) the possible formation of metal clusters in the gas phase; (iii) the kinetic energy of the metal atoms/clusters landing on the IL and thus how deeply they will penetrate. Additionally, the physical properties of the IL (i.e. surface tension) or substituent functional groups also play a major role in the metal cluster growth²⁰.

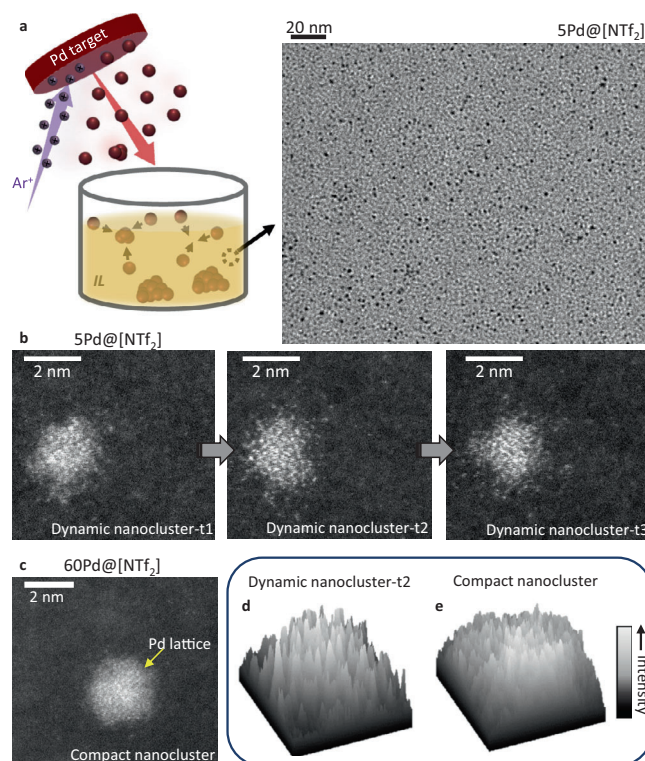


Fig. 1 The magnetron sputtering process and electron microscopy images of Pd nanoclusters in ILs. **a** Scheme of the Pd magnetron sputtering deposition in IL. Bright-field TEM image shows the homogeneous size distribution of 5Pd@[NTf₂]. **b** Aberration-corrected scanning transmission electron microscopic (AC-STEM) images acquired over time (details in Supplementary Fig. 8) to show the dynamic character of 5Pd@[NTf₂] system. **c** AC-STEM image of Pd species deposited for 60Pd@[NTf₂] (details in Supplementary Fig. 11) shows nanocluster with a more compact structure compared to 5Pd@[NTf₂], with metallic Pd lattice planes being observed. **d, e** Intensity histograms were taken from AC-STEM images to highlight the difference between compact and dynamic Pd nanoclusters surfaces, respectively. The intensity bar refers to STEM image contrast, which correlates with the number of Pd atoms in the image.

In this work, magnetron sputtering depositions of Pd species into ILs were carried out using ultraclean conditions with a background-pressure of 5.3×10^{-6} Pa, work pressure fixed at 4.0×10^{-1} Pa using Ar gas (99.9995%) and a high purity Pd target (99.995%). Theoretical simulations for different potentials were performed prior to Pd depositions in IL to optimise the ejection of single Pd atoms (see Supplementary Methods for further details). The simulations showed the highest probability for single Pd atoms to be sputtered from the target at applied potentials up to 800 eV, with two- or three-atom Pd clusters being minority species (i.e. at 320 eV, 31 single Pd atoms are ejected for each two-atom cluster—Supplementary Fig. 2). Therefore, the Pd depositions in ILs were performed using a potential applied within the range displayed in Supplementary Fig. 2. The only parameters varied during the depositions were the Pd concentration and the nature of IL anion. Further details of the magnetron sputtering parameters can be found in Supplementary Table 1 and Supplementary Methods. As expected, the Pd concentration in ILs increased linearly with the Pd deposition time (Supplementary Fig. 3). Therefore, the resulting materials were named as XPd@Y where X is Pd deposition time in min and Y is the nature of the anion in $[\text{C}_4\text{C}_1\text{Im}]$. Y as the IL cation was kept constant. For instance, Pd species deposited for 5 min (Pd concentration of 0.12 wt%) in $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ would be denoted as 5Pd@[NTf₂].

Pd nanocluster morphology and atomic structure. All TEM analyses were performed directly in IL to evaluate the true morphology and atomic structure of the Pd species. Bright-field TEM imaging revealed variation of Pd nanocluster size distribution, from sub-1 nm in 1Pd@[NTf₂] to 2–3 nm in all other ILs (Supplementary Figs. 4, 5, 9, 10 and 12).

Atomic-scale investigation using aberration-corrected scanning transmission electron microscopy (AC-STEM) for 5Pd@[NTf₂] revealed unusual features—a Pd core surrounded by a diffuse shell of satellite Pd atoms, with the overall nanocluster diameter of 2 ± 1.0 nm (Fig. 1b). The structure is dynamic with individual Pd atoms reshuffling over time, as revealed by time-series AC-STEM imaging, and henceforth is referred to as a dynamic nanocluster (Fig. 1b and Supplementary Figs. 6–8). However, the further increase of the Pd concentration in [C₄C₁Im][NTf₂] (Fig. 1c and Supplementary Fig. 11), or the use of a different IL with the anion [PF₆][−] (Supplementary Fig. 13), produces a compact Pd nanocluster with significantly lower dynamics than 5Pd@[NTf₂]. In the compact Pd nanoclusters, atomic lattice planes (111) of face-centred cubic Pd are observed (Supplementary Figs. 9, 10 and 12), whereas in 5Pd@[NTf₂] it appears to be amorphous (Fig. 1b and Supplementary Fig. 5). The overall three-dimensional disorder of the dynamic Pd nanoclusters is also apparent in the irregular intensity distribution extracted from AC-STEM images (Fig. 1d and Supplementary Fig. 8b), as compared to a more homogenous intensity pattern typical of the compact nanoclusters such as 60Pd@[NTf₂] and 5Pd@[PF₆] (Fig. 1e and Supplementary Figs. 11b and 13b).

Matrix-assisted laser desorption ionisation time of flight mass spectrometry clearly shows the presence of single Pd atoms in Pd@[NTf₂] (Supplementary Figs. 14–20). The presence of single Pd atoms appears to be strongly dependent on the type of IL, as changing the IL cation ([C₄C₁Im]⁺ to 1,2-dimethyl-3-butylimidazolium cation ([C₄C₁C₁Im]⁺) or anion ([NTf₂][−]) to hexafluorophosphate anion ([PF₆][−]) both have an impact on

the availability of Pd atoms, such that no single Pd atoms are detected in [C₄C₁C₁Im][NTf₂] and [C₄C₁Im][PF₆]. As the loading of the metal increases, the proportion of Pd existing in the atomic form decreases, consistent with the AC-STEM observations that dynamic nanoclusters exist only in [C₄C₁Im][NTf₂] and only at low loading of metal.

Pd nanocluster electronic environment. A combination of X-ray photoelectron spectroscopy (XPS), X-ray absorption spectroscopy (XAS) and Raman spectroscopy provided insight into the interactions between Pd and [C₄C₁Im][NTf₂]. The Pd 3d core XPS spectrum indicates that Pd nanoclusters in [C₄C₁Im][NTf₂] exist in two different electronic environments: the metallic Pd peak at 335.9 eV and a peak at 338.5 eV that we assign to the interaction between surface Pd atoms with [C₄C₁Im][NTf₂] (Fig. 2a, Supplementary Figs. 21–25 and Supplementary Table 2). The integral intensity of metallic Pd⁰ peak shows a relative decrease as the loading of Pd in [C₄C₁Im][NTf₂] decreases (from 60Pd@[NTf₂] to 10Pd@[NTf₂]) (Fig. 2a). These results correlate with the observation in AC-STEM and mass spectrometry, suggesting that surface Pd atoms interact with IL and that dynamic Pd nanoclusters exist mainly at low loadings.

X-ray absorption near-edge structure (XANES) measurements for 5Pd@[NTf₂] and 30Pd@[NTf₂] Pd K-edge show a similar adsorption edge (*E*₀) to bulk Pd metal but with a higher white line (*H*_w) intensity in the 30Pd@[NTf₂] and a further *H*_w intensity increases for 5Pd@[NTf₂]. This indicates *d*-electron depletion of Pd nanoclusters due to charge transfer to the more electronegative groups of the [C₄C₁Im][NTf₂] IL (Fig. 2b)²⁸, consistent with the XPS results. Extended X-ray absorption fine structure (EXAFS) analysis of 5Pd@[NTf₂] and 30Pd@[NTf₂] show a peak at ca. 2.5 Å associated with Pd–Pd bond, same as in bulk metal, as well as another peak at ca. 1.7 Å that we assigned to Pd atom interaction with [C₄C₁Im][NTf₂] (Fig. 2c). Density functional theory (DFT) modelling confirmed that the Pd coordination

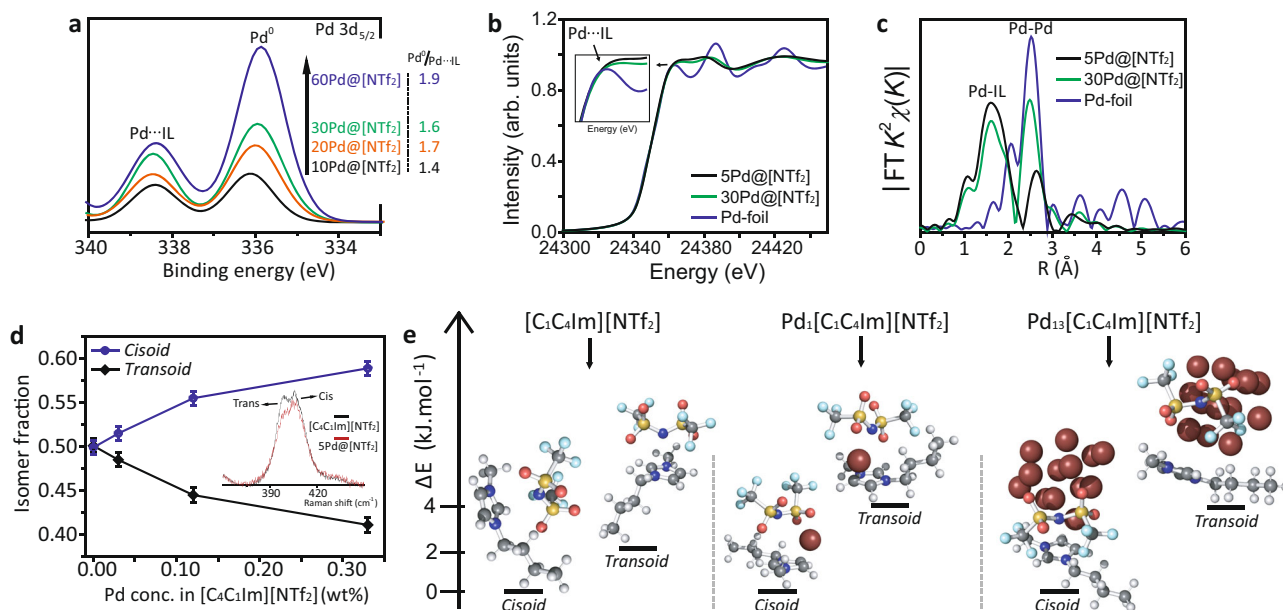


Fig. 2 Electronic environment of Pd and ILs system. **a** X-ray photoelectron spectroscopy (XPS) spectra for the Pd 3d_{5/2} of 10Pd@[NTf₂], 20Pd@[NTf₂], 30Pd@[NTf₂] and 60Pd@[NTf₂] (no Pd signal was observed at lower Pd concentration—Supplementary Fig. 22a). **b, c** Pd K-edge X-ray absorption near-edge structure (XANES) and Fourier transform of k²-weighted extended X-ray absorption fine structure (EXAFS) spectra, respectively, of 5Pd@[NTf₂], 30Pd@[NTf₂] and Pd foil (PdO spectrum can be found in Supplementary Fig. 26). Pd-IL: coordination of Pd-C from [C₄C₁Im] cation and/or Pd-O/N from [NTf₂] anion. **d** Relative *cis*-/*trans*- conformational isomer (conformer) fractions as a function of Pd concentration obtained from Raman spectroscopic analysis of pure [C₄C₁Im][NTf₂] and 1Pd@[NTf₂], 5Pd@[NTf₂] and 10Pd@[NTf₂]. **e** Energy difference between the *cis*- and *trans*- conformers for [C₄C₁Im][NTf₂], Pd₁[C₄C₁Im][NTf₂] and Pd₁₃[C₄C₁Im][NTf₂] from left to right, respectively.

occurs via N and O atoms of the $[\text{NTf}_2]^-$ anion for the $\text{Pd}_{13}[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$, whereas for the $\text{Pd}_1[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ complex the coordination occurs via C-atoms and O-atoms from the cation and anion, respectively (Fig. 2e). Moreover, the intensity and the peak corresponding to Pd–Pd distance decreases as the Pd loading decreases ($30\text{Pd}@\text{NTf}_2$ to $5\text{Pd}@\text{NTf}_2$) alongside to a relative increase of the Pd-IL peak intensity, suggesting a higher number of Pd atoms interacting with the $[\text{C}_4\text{C}_1\text{Im}]$ cation and/or $[\text{NTf}_2]$ anion at lower Pd loading in IL, in agreement with the XPS analysis. Additionally, for $5\text{Pd}@\text{NTf}_2$ EXAFS indicated slightly longer Pd–Pd bonds compared to $30\text{Pd}@\text{NTf}_2$ and Pd bulk, which can be related to the dynamic Pd nanocluster behaviour at relatively low Pd loading in IL ($5\text{Pd}@\text{NTf}_2$)²⁹. Both XPS and XAS demonstrate a strong interaction between surface Pd atoms and $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ and with the core Pd atoms remaining in metallic-like state. Raman spectroscopy allows probing the IL and shows the increase of *cis*-conformation of the $[\text{NTf}_2]$ anion, as the loading of Pd increases (Fig. 2d, Supplementary Fig. 27 and Supplementary Table 3). DFT modelling also shows that for $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ the *cis*-conformer is slightly more stable than the *trans*-conformer by 2.1 kJ mol⁻¹, the addition of a Pd atom, as in $\text{Pd}_1[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ and $\text{Pd}_{13}[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$, increases the difference in stability of the conformers to 4.1 and 3.9 kJ mol⁻¹, respectively (Fig. 2e and Supplementary Fig. 28).

Catalytic evaluation of Pd nanocluster. The catalytic performance of the $\text{Pd}@\text{NTf}_2$ system was evaluated in a cyclopropanation of alkenes where Pd facilitates carbene transfer from ethyl diazoacetate (EDA) to alkenes (Fig. 3a, b, for further details, see Supplementary Tables 4 and 5 and Supplementary Notes 1)^{30–33}.

All materials were found to be active in the cyclopropanation of styrene, giving good conversions and selectivities towards the cyclopropane product. In order to compare relative catalytic conversions of different Pd nanoclusters, a series of experiments in $\text{Pd}@\text{NTf}_2$ with a normalised metal loading in the reaction mixture were carried out (0.5 mol% Pd) (Fig. 3b; for further details see, Supplementary Table 4 and Supplementary Notes 1). While selectivities were similar, a gradual increase in the conversion was observed with a decrease in the Pd metal concentration in IL (from $60\text{Pd}@\text{NTf}_2$ to $10\text{Pd}@\text{NTf}_2$), with an abrupt increase for $7\text{Pd}@\text{NTf}_2$, $5\text{Pd}@\text{NTf}_2$ and $1\text{Pd}@\text{NTf}_2$ (Fig. 3b and Supplementary Table 4), correlating with the transition from compact to dynamic Pd nanoclusters, indicating a higher fraction of accessible Pd atoms in the latter.

In order to verify the mode of catalysis by dynamic Pd nanoclusters, mercury poisoning tests were performed (Fig. 3c). It is well established that heterogeneous catalysts, in contrast to homogenous catalysts, lose their catalytic activity in the presence of Hg due to the poisoning of their surface³⁴. Indeed, complete inhibition of the catalysis was observed for compact Pd nanoclusters (from $60\text{Pd}@\text{NTf}_2$ to $10\text{Pd}@\text{NTf}_2$; Fig. 3c), suggesting the absence and/or very low population of dynamic Pd nanoclusters in agreement with the AC-STEM observations. Remarkably, for the dynamic Pd nanoclusters, the catalytic activity was maintained for $1\text{Pd}@\text{NTf}_2$ and only 32% decrease in yield was observed for $5\text{Pd}@\text{NTf}_2$ (Fig. 3c and Supplementary Table 6), suggesting a pseudo-homogeneous mode of catalysis by the dynamic nanoclusters. Additional poisoning tests were conducted for higher Pd concentration in $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ and $5\text{Pd}@\text{PF}_6$ (Supplementary Table 6), which were completely deactivated similarly to traditional heterogeneous catalysts.

We have performed an additional poisoning tests using dibenzo[*a,e*]cyclooctene (DCT), which selectively binds to single Pd sites^{34–37}. DCT experiments showed that $5\text{Pd}@\text{NTf}_2$ and $30\text{Pd}@\text{NTf}_2$ did not lose activity (Supplementary Table 7),

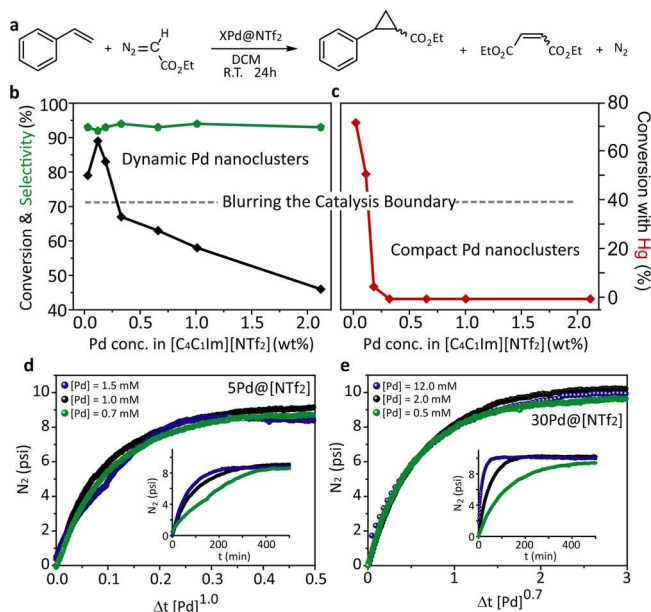


Fig. 3 Cyclopropanation of styrene. **a** Reaction scheme of the cyclopropanation of styrene as a model reaction with $\text{XPd}@\text{Y}$. A complete substrate scope can be found in Supplementary Notes 1. **b** Conversion (black) and selectivity (green) with varying Pd contents in $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ determined after 24 h; reaction conditions: diazoethyl acetate (1 mmol), styrene (5 mmol), $\text{XPd}@\text{NTf}_2$ (5 mol%, based on Pd) and DCM (5 mL) at room temperature. **c** Conversion with varying Pd contents in $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$ and the presence of Hg, reaction conditions as in **b** with Hg. **b, c** Dashed line represents the boundary in between pseudo-homogeneous and classical heterogeneous catalysts. **d, e** represent the time normalised analysis of the N_2 release (which is coupled to the overall catalytic transformation) with varying catalyst concentrations. The catalyst order is represented as the magnitude of the $[\text{Pd}]$ concentration and the original data can be found as insets in **d** and **e**, respectively. A catalyst order of 1 is found for $5\text{Pd}@\text{NTf}_2$ (**d**) and an order of 0.7 is found for $30\text{Pd}@\text{NTf}_2$ (**e**). Reaction conditions in **d, e**: 1 mmol ethyl diazoacetate, 5 mmol styrene, 5 mL DCM at 35 °C with varying concentrations of Pd but constant concentration of $[\text{C}_4\text{C}_1\text{Im}][\text{NTf}_2]$.

confirming that dynamic Pd nanocluster are the active sites, rather than leached single Pd atoms. These results were further reinforced by filtration tests, as both catalytic systems ($5\text{Pd}@\text{NTf}_2$ and $30\text{Pd}@\text{NTf}_2$) did not show activity after the filtration (Supplementary Table 8). Additionally, recyclability experiments showed that $5\text{Pd}@\text{NTf}_2$ and $30\text{Pd}@\text{NTf}_2$ are recyclable and maintain their activity over three cycles (Supplementary Table 9).

Furthermore, the N_2 release was monitored during the course of the reaction to elucidate the catalyst order (Fig. 3d, e and Supplementary Fig. 29). After pre-activation with styrene, the catalyst order was determined by the variable time normalisation analysis method introduced by Bures and co-workers^{38–40}, showing a catalyst order of 1.0 for $5\text{Pd}@\text{NTf}_2$ and 0.7 for $30\text{Pd}@\text{NTf}_2$ (Fig. 3d, e). These results further confirm the pseudo-homogeneous character of the dynamic Pd nanoclusters, while the order determined for compact Pd nanoclusters is in the expected lower range for heterogeneous catalysts due to the inherently decreased amounts of atoms accessible for catalysis. This effect relates to the condition that the reaction takes place only on the surface of the catalysts. TEM experiment where dynamic Pd nanoclusters were retrieved from the reaction mixture directly onto a TEM grid demonstrated that the dynamic clusters remain unchanged in size and structure after the reaction (Supplementary Fig. 30).

In summary, we have explored the structure, dynamics and catalytic properties of a series of palladium nanoclusters from sub-1 to 3 nm sizes by a variety of analytical and imaging techniques, providing atomic-scale information. Nanoclusters appear to behave as dynamic structures with properties controlled by the nature of IL (both cation and anion) and the loading of the metal in the liquid. Anions interacting directly with palladium atoms play a particularly important role in determining the electronic state and dynamics of surface atoms. In particular, [NTf₂] anions in [C₄C₁Im][NTf₂] have been shown to bond to surface Pd atoms and create a diffuse shell of metal atoms around the nanocluster. This phenomenon has a significant implication for the mode of catalysis, changing the heterogeneous behaviour, exhibited by Pd nanoclusters with compact structures, to a homogeneous mode exhibited by dynamic 5Pd@[NTf₂]. Our study demonstrates that the effectiveness of nanocatalysts depends not only on the surface area of the nanoclusters but also on the dynamic behaviour of their surface atoms controlled by the support environment.

Methods

Pd nanocluster preparation. Pd species were deposited into ILs (IL synthesis details are in the Supplementary Information) using a bespoke AJA magnetron sputtering system with a load-lock sample transfer facility coupled to a glovebox. The Pd target (99.995%) was purchased from AJA International. IL samples were loaded through a glovebox into the magnetron sputtering system to avoid the presence of moisture. In a typical experiment, anhydrous IL (0.750 g) is placed in a petri dish while in the glovebox. The sample was transferred to a load-lock under a N₂ environment, pumped-down to a background pressure of 5.3×10^{-5} Pa (for ca. 1 h) and then transferred to the main chamber, which reached a background pressure of 5.3×10^{-6} Pa in ca. 1 h. The power and working pressure used in all depositions were 100 W and 4.0×10^{-1} Pa (Argon—99.9999%), respectively.

Sample characterisation. Pd nanoclusters deposited by magnetron sputtering in ILs were primarily characterised by TEM, with each being analysed from at least two different deposition batches. All microscopic analyses were performed in the absence of solvents in order to evaluate the true morphology of the Pd species deposited in the ILs. TEM measurements were performed using a JEOL 2100F FEG-TEM operated with an accelerating voltage of 200 kV. AC-STEM measurements were performed using a JEOL 2100F scanning transmission electron microscope with a CEOS aberration corrector operated with an accelerating voltage of 200 kV. The samples were prepared in the following manner: copper mesh, holey carbon film TEM grids (Agar Scientific, UK) were glow discharged (Agar Turbo Coater, 0.2 mbar, 5 mA, 10 s) before the addition of Pd@IL (10 μ L). The Pd@IL suspension was kept on the grid for 2 min before the excess was removed using filter paper. This approach allowed the formation of ‘pools’ of IL within the holes of the holey carbon, providing an effective contrast-free region to observe the palladium structures.

XPS measurements were performed using a Kratos AXIS Ultra DLD instrument. The chamber pressure during the measurements was 6.7×10^{-7} Pa. Wide energy range survey scans were collected at pass energy of 80 eV in hybrid slot lens mode and a step size of 0.5 eV, for 20 min. High-resolution data on the Pd 3d, C 1s, N 1s, O 1s, S 2p and F 1s photoelectron peaks were collected at a pass energy of 20 eV over energy ranges suitable for each peak and collection times of 5 min, step sizes of 0.1 eV. The charge neutraliser filament was used to prevent the sample charging over the irradiated area. The X-ray source was a monochromated Al K α emission, run at 10 mA and 12 kV (120 W). The energy range for each ‘pass energy’ was calibrated using the Kratos Cu 2p_{3/2}, Ag 3d_{5/2} and Au 4f_{7/2} three-point calibration method. The transmission function was calibrated using a clean gold sample method for all lens modes and the Kratos transmission generator software within Vision II. The data were processed with CASAXPS (Version 2.3.17). The high-resolution data were charge corrected to the reference F 1s signal at 688.9 eV.

XAS measurements of Pd K-edge were performed at room temperature (B18 beamline) at the Diamond Synchrotron Light. XANES and EXAFS spectra of a Pd foil and PdO standards were measured and the energy calibrated by aligning the respective absorption edges. The data were calibrated and normalised by a linear pre-edge subtraction using the ATHENA software. DFT calculations were performed using version 4.1.0 of the Orca Program package⁴¹.

Micro Raman spectroscopy was performed using a Horiba Jobin Yvon LabRAM HR Raman spectrometer. Spectra were acquired using a 785 nm laser (at 24 mW power), a $\times 100$ objective lens and a 300 μ m confocal pinhole. To simultaneously scan a range of Raman shifts and control the spectral resolution, either a 600 or 1800 lines mm⁻¹ rotatable diffraction grating along a path length of 800 mm was employed. Spectra were acquired using a Synapse CCD detector (1024 pixels) thermoelectrically cooled to -60 °C.

Catalytic tests. All reactions were prepared in a glovebox under inert atmosphere using a 50 mL Schlenk flask equipped with a magnetic stir bar. EDA (1 mmol) was added into a solution containing the Pd catalyst (0.5 mol% Pd) and styrene (5 mmol) in CH₂Cl₂ (5/10 mL). The Schlenk flask was sealed with a septum cap and the reaction mixture was removed from the glovebox. After 24 h of stirring at RT, the mixture was filtered through silica gel and volatiles were removed in vacuo. Products have been previously described and their identification was straightforward from comparison with the reported data⁴². Substrate conversion and selectivity and yield of cyclopropane product were determined by ¹H nuclear magnetic resonance spectroscopy using 1,2-dibromoethane (0.25 mmol) as an internal standard.

Data availability

All experimental and simulation data in the main text and Supplementary Materials are available upon request to the authors.

Code availability

Molecular Dynamics codes used in the simulation are available upon request to harriet.ahlgren@univie.ac.at.

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References

1. Wang, A., Li, J. & Zhang, T. Heterogeneous single-atom catalysis. *Nat. Rev. Chem.* **2**, 65–81 (2018).
2. Zhang, Z. et al. Thermally stable single atom Pt/m-Al₂O₃ for selective hydrogenation and CO oxidation. *Nat. Commun.* **8**, 16100 (2017).
3. Sun, J.-K. et al. General synthetic route toward highly dispersed metal clusters enabled by poly(ionic liquid)s. *J. Am. Chem. Soc.* **139**, 8971–8976 (2017).
4. Cheng, N. et al. Platinum single-atom and cluster catalysis of the hydrogen evolution reaction. *Nat. Commun.* **7**, 13638 (2016).
5. Liu, J. Catalysis by supported single metal atoms. *ACS Catal.* **7**, 34–59 (2017).
6. Li, Y. et al. A Pt-cluster-based heterogeneous catalyst for homogeneous catalytic reactions: X-ray absorption spectroscopy and reaction kinetic studies of their activity and stability against leaching. *J. Am. Chem. Soc.* **133**, 13527–13533 (2011).
7. Papp, C. Catalysis at the limit. *Nat. Chem.* **10**, 995–996 (2018).
8. Imaoka, T. et al. Platinum clusters with precise numbers of atoms for preparative-scale catalysis. *Nat. Commun.* **8**, 688 (2017).
9. Tian, S. et al. Carbon nitride supported Fe₂ cluster catalysts with superior performance for alkene epoxidation. *Nat. Commun.* **9**, 2353 (2018).
10. Liu, L. & Corma, A. Metal catalysts for heterogeneous catalysis: from single atoms to nanoclusters and nanoparticles. *Chem. Rev.* **118**, 4981–5079 (2018).
11. Therrien, A. J. et al. An atomic-scale view of single-site Pt catalysis for low-temperature CO oxidation. *Nat. Catal.* **1**, 192–198 (2018).
12. Du, X. & Jin, R. Atomically precise metal nanoclusters for catalysis. *ACS Nano* **13**, 7383–7387 (2019).
13. Freund, H.-J. The surface science of catalysis and more, using ultrathin oxide films as templates: a perspective. *J. Am. Chem. Soc.* **138**, 8985–8996 (2016).
14. Ishida, Y., Corpuz, R. D. & Yonezawa, T. Matrix sputtering method: a novel physical approach for photoluminescent noble metal nanoclusters. *Acc. Chem. Res.* **50**, 2986–2995 (2017).
15. Luza, L. et al. Revealing hydrogenation reaction pathways on naked gold nanoparticles. *ACS Catal.* **7**, 2791–2799 (2017).
16. Torimoto, T. et al. Sputter deposition onto ionic liquids: simple and clean synthesis of highly dispersed ultrafine metal nanoparticles. *Appl. Phys. Lett.* **89**, 243117 (2006).
17. Deng, L. et al. Highly correlated size and composition of Pt/Au alloy nanoparticles via magnetron sputtering onto liquid. *Langmuir* **36**, 3004–3015 (2020).
18. Pongthawornsakun, B. et al. Deposition of Pt nanoparticles on TiO₂ by pulsed direct current magnetron sputtering for selective hydrogenation of vanillin to vanillyl alcohol. *Catal. Today* **358**, 51–59 (2020).
19. Calabria, L. et al. Confined naked gold nanoparticles in ionic liquid films. *Nanoscale* **9**, 18753–18758 (2017).
20. Wender, H., Migowski, P., Feil, A. F., Teixeira, S. R. & Dupont, J. Sputtering deposition of nanoparticles onto liquid substrates: recent advances and future trends. *Coord. Chem. Rev.* **257**, 2468–2483 (2013).
21. Eshetu, G. G., Armand, M., Ohno, H., Scrosati, B. & Passerini, S. Ionic liquids as tailored media for the synthesis and processing of energy conversion materials. *Energy Environ. Sci.* **9**, 49–61 (2016).
22. König, D., Richter, K., Siegel, A., Mudring, A.-V. & Ludwig, A. High-throughput fabrication of Au–Cu nanoparticle libraries by combinatorial sputtering in ionic liquids. *Adv. Funct. Mater.* **24**, 2049–2056 (2014).

23. Torimoto, T. et al. Ultrathin oxide shell coating of metal nanoparticles using ionic liquid/metal sputtering. *J. Mater. Chem. A* **3**, 6177–6186 (2015).
24. Wender, H. et al. Synthesis of gold nanoparticles in a biocompatible fluid from sputtering deposition onto castor oil. *Chem. Commun.* **46**, 7019–7021 (2010).
25. Sugioka, D., Kameyama, T., Kuwabata, S., Yamamoto, T. & Torimoto, T. Formation of a Pt-decorated Au nanoparticle monolayer floating on an ionic liquid by the ionic liquid/metal sputtering method and tunable electrocatalytic activities of the resulting monolayer. *ACS Appl. Mater. Interfaces* **8**, 10874–10883 (2016).
26. Wender, H. et al. Sputtering onto liquids: from thin films to nanoparticles. *J. Phys. Chem. C* **115**, 16362–16367 (2011).
27. Kuwabata, S., Tsuda, T. & Torimoto, T. Room-temperature ionic liquid. A new medium for material production and analyses under vacuum conditions. *J. Phys. Chem. Lett.* **1**, 3177–3188 (2010).
28. Cook, S. C., Padmos, J. D. & Zhang, P. Surface structural characteristics and tunable electronic properties of wet-chemically prepared Pd nanoparticles. *J. Chem. Phys.* **128**, 154705 (2008).
29. Ohba, T. et al. An origin for lattice expansion in PVP-protected small Pd metal nanoparticles. *Bull. Chem. Soc. Jpn.* **90**, 720–727 (2017).
30. Lakshmi Kantam, M. et al. Alumina supported copper nanoparticles for aziridination and cyclopropanation reactions. *Catal. Commun.* **8**, 1963–1968 (2007).
31. Chen, S., Ma, J. & Wang, J. Palladium-catalyzed cyclopropanation of electron-deficient olefins with aryldiazocarbonyl compounds. *Tetrahedron Lett.* **49**, 6781–6783 (2008).
32. Shishilov, O. N. et al. Palladium(II) carboxylates and palladium(I) carbonyl carboxylate complexes as catalysts for olefin cyclopropanation with ethyl diazoacetate. *Dalton Trans.* 6626–6633, (2009).
33. Martín, C., Molina, F., Alvarez, E. & Belderrain, T. R. Stable N-heterocyclic carbene (NHC)–palladium(0) complexes as active catalysts for olefin cyclopropanation reactions with ethyl diazoacetate. *Chem. Eur. J.* **17**, 14885–14895 (2011).
34. Gärtner, D., Welther, A., Rad, B. R., Wolf, R. & Jacobi von Wangelin, A. Heteroatom-free arene-cobalt and arene-iron catalysts for hydrogenations. *Angew. Chem. Int. Ed.* **53**, 3722–3726 (2014).
35. Hamasaka, G., Ichii, S. & Uozumi, Y. A palladium NNC-pincer complex as an efficient catalyst precursor for the Mizoroki–Heck reaction. *Adv. Synth. Catal.* **360**, 1833–1840 (2018).
36. Luo, X. L. & Crabtree, R. H. Homogeneous catalysis of silane alcoholysis via nucleophilic attack by the alcohol on an Ir(η^2 -HSiR₃) intermediate catalyzed by [IrH₂S₂(PPh₃)₂]SbF₆ (S = solvent). *J. Am. Chem. Soc.* **111**, 2527–2535 (1989).
37. Neate, P. G. N., Greenhalgh, M. D., Brennessel, W. W., Thomas, S. P. & Neidig, M. L. Mechanism of the bis(imino)pyridine-iron-catalyzed hydromagnesiation of styrene derivatives. *J. Am. Chem. Soc.* **141**, 10099–10108 (2019).
38. Burés, J. Variable time normalization analysis: general graphical elucidation of reaction orders from concentration profiles. *Angew. Chem. Int. Ed.* **55**, 16084–16087 (2016).
39. Burés, J. A simple graphical method to determine the order in catalyst. *Angew. Chem. Int. Ed.* **55**, 2028–2031 (2016).
40. Nielsen, C. D. T. & Burés, J. Visual kinetic analysis. *Chem. Sci.* **10**, 348–353 (2019).
41. Neese, F. Software update: the ORCA program system, version 4.0. *WIREs Comput. Mol. Sci.* **8**, e1327 (2018).
42. Cho, D.-J. et al. Chiral C₂-symmetric bisferrocenyldiamines as ligands for transition metal catalyzed asymmetric cyclopropanation and aziridination. *Tetrahedron* **10**, 3833–3848 (1999).

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Author contributions

I.C., A.W. and C.M. performed part of the ionic liquid synthesis and catalytic experiments. R.W.L., J.Y. and Z.Y.L. conducted TEM and AC-STEM measurements. J.P. performed the magnetron sputtering deposition of Pd species in ionic liquid and ICP-OES measurements. A.S. performed part of the ionic liquid synthesis and XPS measurements. G.A.R. conducted the Raman spectroscopy measurements. E.H.Å. conducted the classical molecular dynamics simulations with the PARCAS code. E.J. performed the DFT calculations. P.L., A.N.K. and J.A.F. designed the study, analysed the data and co-wrote the paper. All the authors discussed the results and commented on the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

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