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 [C_4C_1Im][NTf_2]. 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.
Catalysts are used in nearly 80% of industrial processes, demanding the maximum catalytic efficiency of rare elements, such as Au, Pt and Pd. Increasing the metal active surface area by utilising catalysts in the form of nanoclusters is one of the most powerful approaches. 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. 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 nature.

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. Crucially, this approach generates highly active cluster surfaces, since no chemical stabilizers and/or solvents are involved in the process. 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 vapor pressure allowing their use under the high vacuum conditions of magnetron sputtering. 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 may alter the environment surrounding 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 unexplored 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 ([C₄C₁Im][NTf₂]) 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 on 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). 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. 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 clusters 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 growth.

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 (99.9995%). 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 in IL 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 [C₄C₁Im]. Y as the IL cation was kept constant. For instance, Pd species deposited for 5 min (Pd concentration of 0.12 wt%) in [C₄C₁Im][NTf₂] 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. 4, 5, 9, 10 and 12), whereas in 5Pd@[NTf₂] it appears to be an array of surface Pd atoms interacting with IL and that dynamic Pd nanoclusters exist mainly at low loadings.

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 Pd 0 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 (E₀) to bulk Pd metal but with a higher white line (Hₜ) intensity in the 30Pd@[NTf₂] and a further Hₜ 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

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Fig. 2 Electronic environment of Pd and ILs system. a X-ray photoelectron spectroscopy (XPS) spectra for the Pd 3d₃/₂ 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 2-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], Pd([C₄C₁Im][NTf₂]) and Pd₃([C₄C₁Im][NTf₂]) from left to right, respectively.
Catalytic evaluation of Pd nanocluster. The catalytic performance of the Pd@[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). 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 Pd@[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 60Pd@[NTf$_2$] to 10Pd@[NTf$_2$]), with an abrupt increase for 7Pd@[NTf$_2$]. 5Pd@[NTf$_2$] and 1Pd@ [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 homogeneous catalysts, lose their catalytic activity in the presence of Hg due to the poisoning of their surface. Indeed, complete inhibition of the catalysis was observed for compact Pd nanoclusters (from 60Pd@[NTf$_2$] to 10Pd@[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 1Pd@[NTf$_2$] and only 32% decrease in yield was observed for 5Pd@[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 [C$_4$C$_1$Im][NTf$_2$] and 5Pd@[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. DCT experiments showed that 5Pd@[NTf$_2$] and 30Pd@[NTf$_2$] did not lose activity (Supplementary Table 7), confirming that dynamic Pd nanoclusters are the active sites, rather than leached single Pd atoms. These results were further reinforced by filtration tests, as both catalytic systems (5Pd@NTf$_2$ and 30Pd@NTf$_2$) did not show activity after the filtration (Supplementary Table 8). Additionally, recyclability experiments showed that 5Pd@NTf$_2$ and 30Pd@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). For pre-activation with styrene, the catalyst order was determined by the variable time normalisation analysis method introduced by Bures and co-workers, showing a catalyst order of 1.0 for 5Pd@[NTf$_2$] and 0.7 for 30Pd@[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 3Pd@[NTf2].

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 (99.9995%) was purchased from AJA International. ILS 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 × 10−3 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 CEDS 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 µL). The Pd/IL suspension was then filtered through silica gel and volatiles were removed in vacuo. High-resolution transmission electron microscopy (HR-TEM) was performed using a JEOL 2100F FEG-TEM instrument. The chamber pressure during the measurements was 0.2 mbar, 5 mA, 10 s) before the addition of Pd/IL (0.2 mbar, 5 mA, 10 s) before the addition of Pd/IL (5 mLA, 0.2 mbar, 5 mA, 10 s) before the addition of Pd/IL (10 s) before the addition of Pd/IL (10 µL).

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 CH3Cl (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 data32. 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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Author contributions
I.C., A.W. and C.M. performed part of the ionic liquid synthesis and catalytic experiments. R.W.L., J.T. 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.A. conducted the classical molecular dynamics simulations with the PARCAS code. E.I. 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.

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