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# Evolution of Metastable Structures at Bimetallic Surfaces from Microscopy and Machine-Learning Molecular Dynamics

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## Abstract

Restructuring of interfaces plays a crucial role in materials science and heterogeneous catalysis. Bimetallic systems, in particular, often adopt very different composition and morphology at surfaces compared to the bulk. For the first time, we reveal a detailed atomistic picture of long-timescale restructuring of Pd deposited on Ag, using microscopy, spectroscopy, and novel simulation methods. By developing and performing accelerated machine-learning molecular dynamics followed by an automated analysis method, we discover and characterize previously unidentified surface restructuring mechanisms in an unbiased fashion, including Pd-Ag place exchange and Ag pop-out, as well as step ascent and descent. Remarkably, layer-by-layer dissolution of Pd into Ag is always preceded by an encapsulation of Pd islands by Ag, resulting in a significant migration of Ag out of the surface and a formation of extensive vacancy pits within a period of microseconds. These metastable structures are of vital catalytic importance, as Ag-encapsulated Pd remains much more accessible to reactants than bulk-dissolved Pd. Our approach is broadly applicable to complex multimetallic systems and enables the previously intractable mechanistic investigation of restructuring dynamics at atomic resolution.

## Introduction

Dynamic restructuring of complex interfaces is ubiquitous in multicomponent materials and can have a significant impact on their physical and chemical properties. In particular, elemental segregation at grain boundaries, dislocations, and surfaces can dramatically alter a range of functional properties, such as fracture strength, plastic deformation, corrosion resistance, and catalytic activity.<sup>1-3</sup> In heterogeneous catalysis, bimetallic and dilute alloy catalysts have garnered much attention due to their potential in enhancing both the activity and the selectivity of hydrogenation reactions.<sup>4</sup> In dilute Pd/Ag, for example, Pd initiates the catalytic cycle as the active site for H<sub>2</sub> dissociation, and the selective hydrogenation

ensues on a more inert Ag host.<sup>5</sup> These alloy surfaces are inherently dynamic at reaction temperatures of interest: compared to the original synthesized state, the surface composition and morphology can differ dramatically after annealing or during the reaction.<sup>6–10</sup> Such restructuring has recently been shown to play a central role in catalytic performance of several alloy systems, including RhPd,<sup>6</sup> NiPt,<sup>11</sup> PdAg,<sup>12</sup> and AgAu.<sup>13</sup> Therefore, fundamental understanding of surface restructuring is crucial to enabling accurate prediction and engineering of catalyst performance.<sup>14</sup>

Mechanistic and kinetic studies of surface restructuring at the atomic level have remained scarce and challenging.<sup>15–21</sup> Experimentally, it is very difficult to probe fast atomic motions with sufficient temporal resolution, even with techniques that possess surface and chemical sensitivities. Computationally, density functional theory (DFT) enables exploration of reaction pathways on model surfaces,<sup>22</sup> but its high computational cost precludes its use for dynamical simulation of realistic systems beyond tens of picoseconds. Furthermore, exploration of large configurational space underlying restructuring processes quickly becomes intractable with brute force approaches. In this work, we overcome all of these limitations by combining microscopy, spectroscopy, and novel simulation methods (Fig. 1; see Experimental Section). Fast and large-scale machine-learning molecular dynamics (MD) is performed using our recently developed Gaussian process (GP) force field,<sup>23</sup> spanning microseconds at first-principles accuracy. Furthermore, a new automated analysis method is used to discover and characterize key surface restructuring mechanisms (Fig. 1). Our streamlined approach enables an efficient and unbiased characterization of restructuring dynamics in complex multimetallic systems.

Surface restructuring of late-transition metals can be classified into three categories: (1) relaxation involving localized atomic displacements;<sup>30</sup> (2) reconstruction involving intralayer atomic site changes;<sup>31</sup> (3) segregation in alloys involving interlayer atomic site changes.<sup>32</sup> All three processes aim to reduce the surface free energy arising from reduced surface atom coordination. In alloy systems, equilibrium distribution of surface and subsurface species

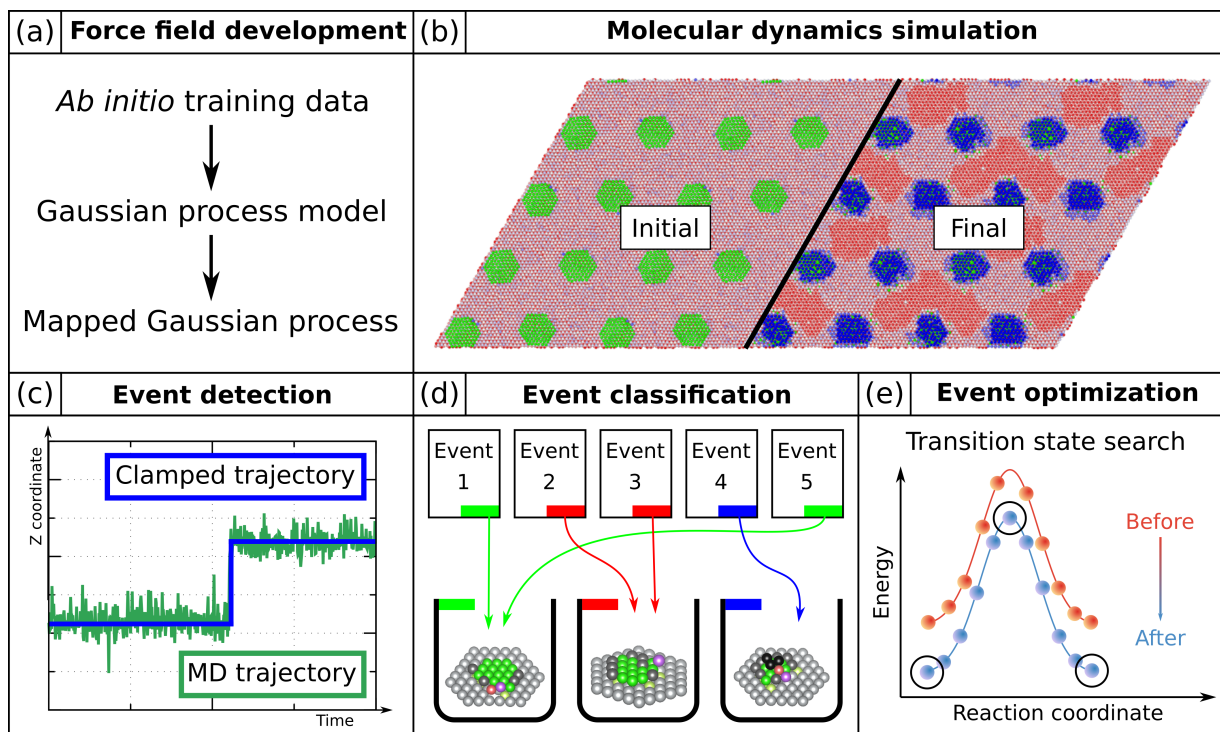


Figure 1: Automated characterization of surface restructuring events. (a) Gaussian process (GP) machine-learning force field is trained on DFT data.<sup>23</sup> (b) The GP model is used to perform fast, large-scale, and long-timescale MD simulation of surfaces. (c) The raw MD trajectory is first "clamped" by assigning all atoms onto discrete FCC lattice sites.<sup>24–26</sup> Then, an event is detected whenever the  $z$ -coordinate (normal to the surface) of any atom changes irreversibly over 1-2 ps.<sup>21</sup> (d) The events are sorted into seven main classes<sup>21</sup> based on changes in the atomic layer and the coordination number. (e) The event pathways are detailed using transition state modeling.<sup>27–29</sup>

can be modeled using the quasi-chemical approximation.<sup>33</sup> The model assumes a chemical equilibrium according to select energetic contributions, which consist of the "three effects" rule<sup>32</sup> (surface energy; strain energy; alloying energy) and the "chemical pump effect"<sup>34</sup> (adsorption energy of molecular species).

In bimetallic systems comprised of soft metals (e.g. Pt, Pd, Au, Ag), the major chemical complexity arises from elemental variations in the surface energy. In general, elements with fuller  $d$ -shells have lower surface energy and segregate to the surface under reducing conditions. For example, an inert metal of group 11 (Cu, Ag, Au) has a full  $d$ -shell and segregates to the surface, while an active metal of group 10 (Ni, Pd, Pt) migrates to the subsurface, ultimately forming e.g. core@shell nanostructures.<sup>35</sup> Based on prior studies using scanning

tunneling microscopy (STM), X-ray photoelectron spectroscopy (XPS), and *ab initio* thermodynamics, Pd dissolves into the Ag bulk after mild heating because of the higher surface energy of Pd.<sup>10</sup> However, the underlying mechanism and timescale have remained unclear.

To the best of our knowledge, the work herein provides the first direct atomistic observation and characterization of the long-timescale restructuring of a bimetallic surface. After annealing at a moderate temperature, monatomic Pd islands on Ag(111) evolve into a dramatically different state within a period of microseconds: a significant amount of Ag migrates out of the surface to encapsulate the Pd islands, resulting in extensive vacancy pits. We discover and characterize previously unidentified elementary restructuring events, including Pd-Ag place exchange and Ag pop-out, as well as step ascent and descent. In contrast to the thermodynamically favored state where Pd is dispersed in the Ag bulk, these structures are metastable and prevail at mild temperatures. They are of vital catalytic importance, as Ag-encapsulated Pd remains much more accessible to reactants than bulk-dissolved Pd. As such, our results provide an important foundation for engineering surface ensemble distributions toward optimal activity-selectivity balance.<sup>36</sup>

## Results and Discussion

Four distinct stages have been established for the restructuring of Pd islands on Ag (Fig. 3): (1) Pd-Ag place exchange; (2) formation of PdAg alloy layer; (3) Ag pop-out and vacancy pit growth; (4) layer-by-layer Pd dissolution. Surprisingly, the first three stages produce nontrivial metastable structures that result from encapsulation of the Pd deposit by Ag and the growth of extensive vacancy pits, as observed in STM images (Fig. 2). As such, these stages comprise a kinetically distinct metastable dissolution mode, previously named "surfactant" mode in the literature<sup>37</sup> due to the similarity to semiconductor growth. This mode precedes the direct layer-by-layer dissolution of Pd into the thermodynamically favored subsurface. Fig. 3 shows the evolution of a hexagonal Pd<sub>91</sub> island on Ag(111) at

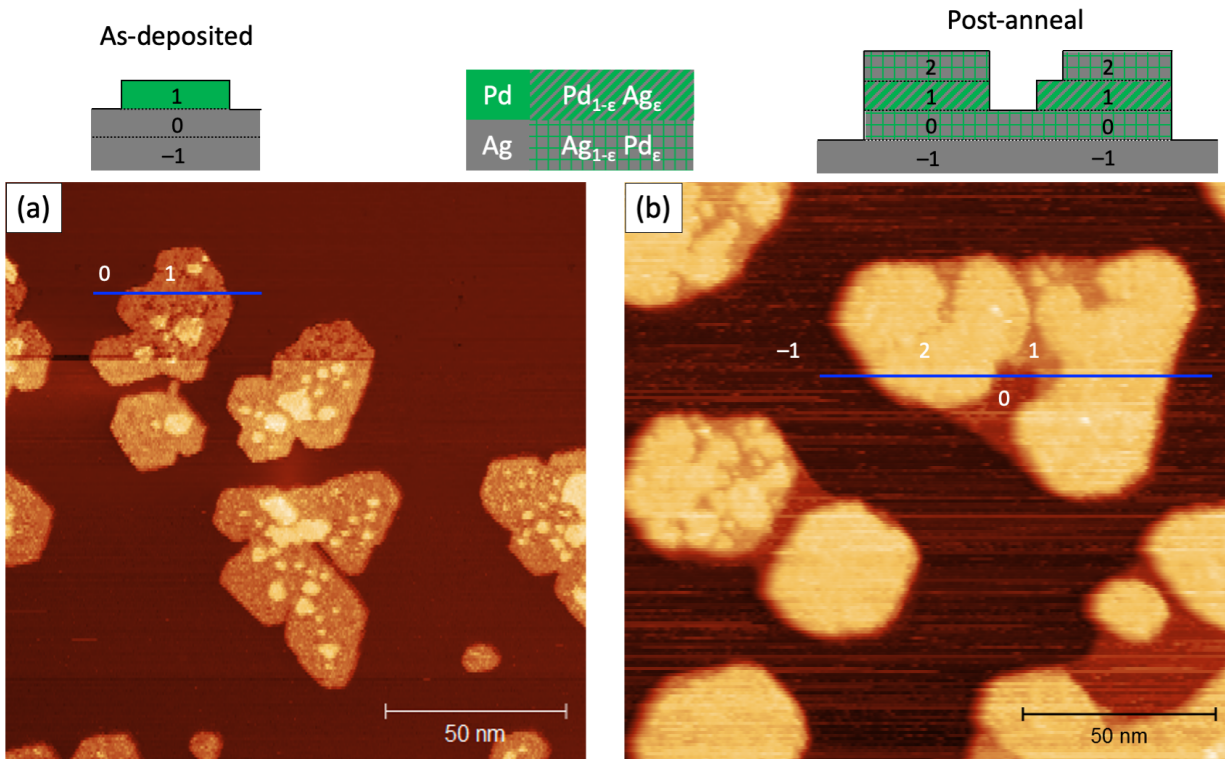


Figure 2: STM images demonstrate the restructuring of Pd islands deposited on Ag(111). Metastable structures form prior to complete dissolution of Pd into the Ag bulk. Schematic height profiles along the blue horizontal lines are depicted above the images, labeled by the layer heights: level  $-1$  (subsurface); level  $0$  (surface); level  $1$  (deposit); level  $2$  (capping). Compositions derived from simulation (Fig. 3) are indicated by colored patterns for each layer (see top center legend), with  $\epsilon \ll 1$  corresponding to the minority component. (a) Pd islands, deposited at 300 K, consist mostly of monolayers at level 1. (b) After mild annealing at 400-450 K, the difference in the elemental surface energy results in encapsulation of the Pd islands by capping layers at level 2, along with extensive vacancy pits at level  $-1$ .

500 K over  $2 \mu\text{s}$  of our simulation. In addition, Table 1 shows four main types of microscopic restructuring events discovered by our automated scheme (Sec. 2, SI).

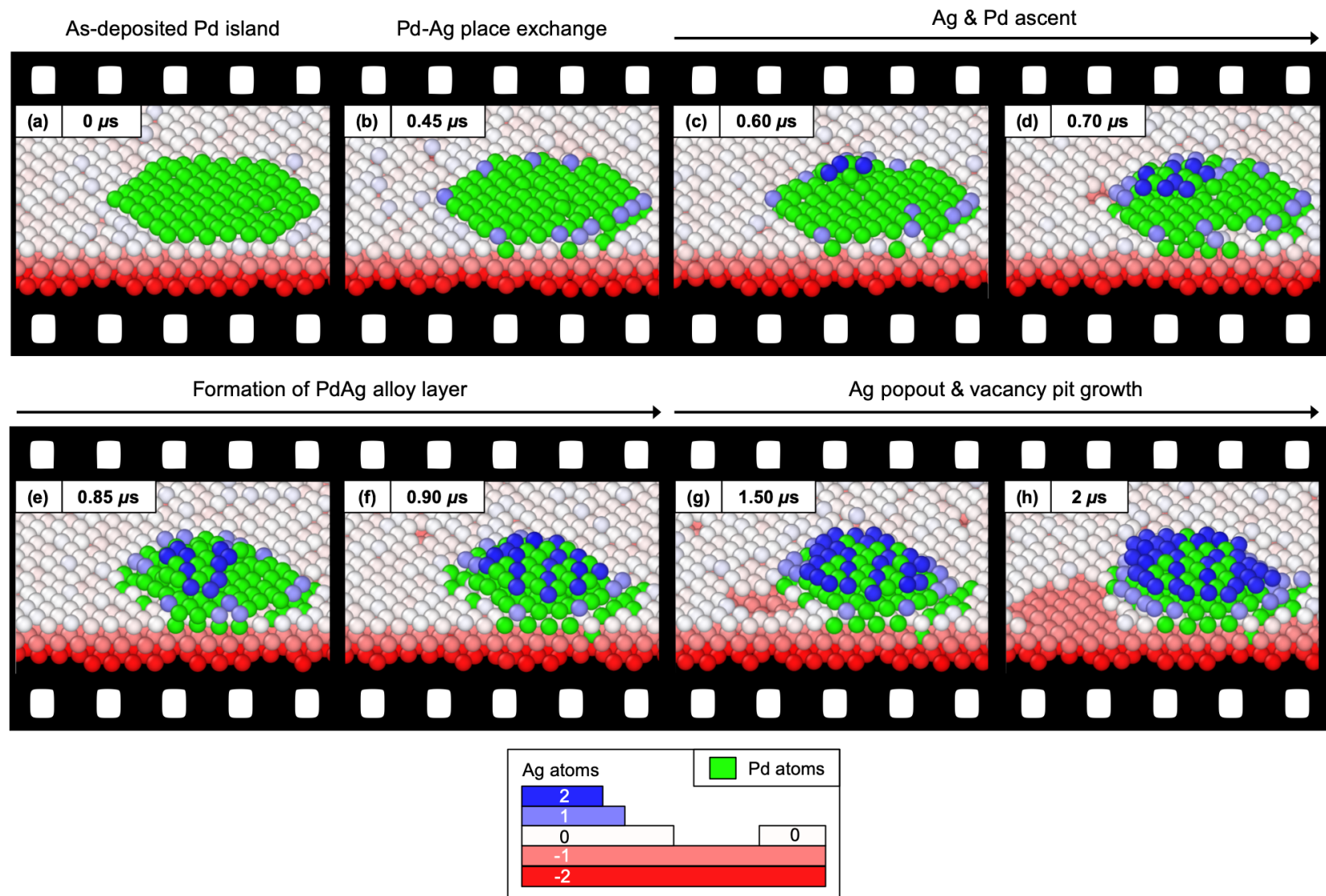
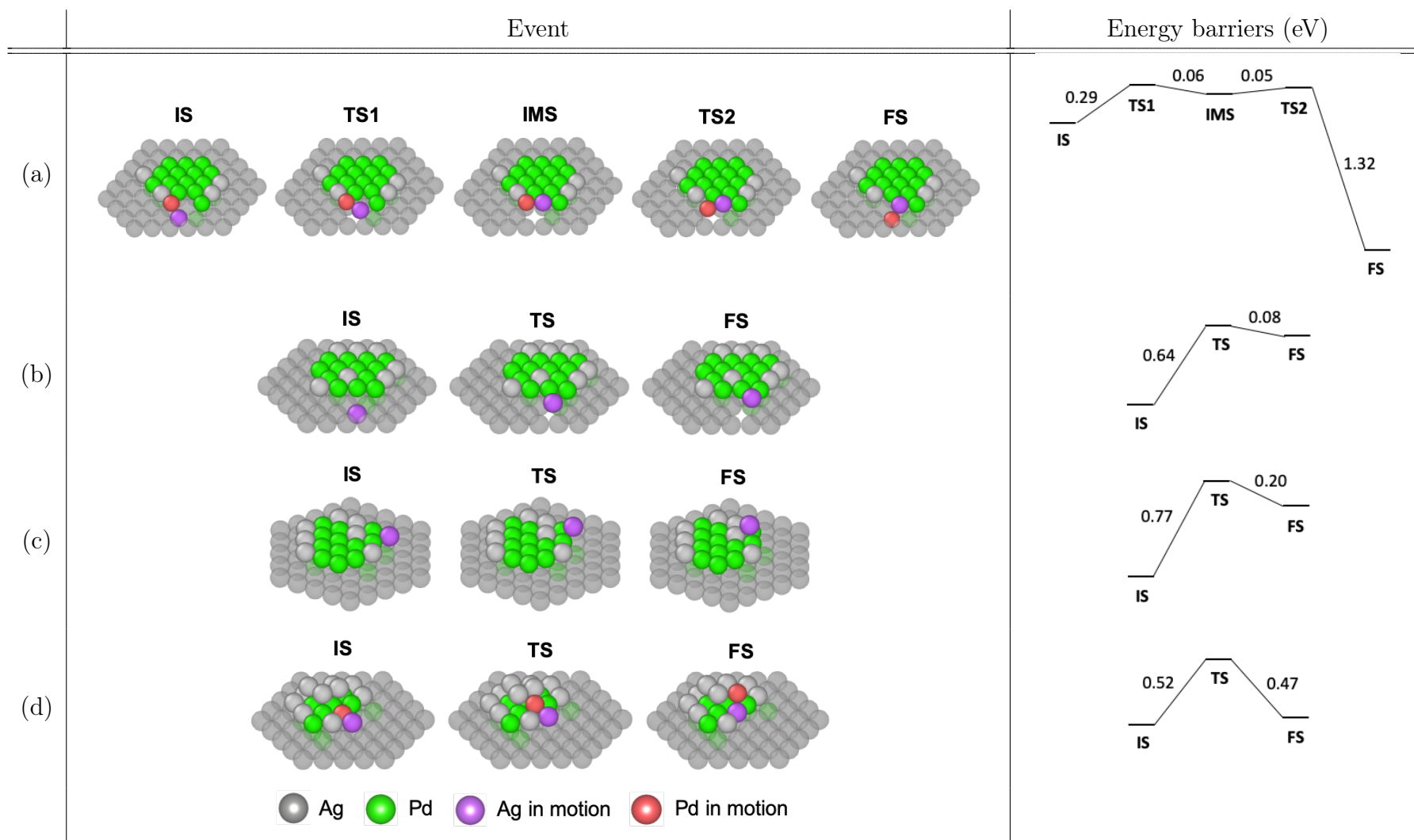


Figure 3: Frames of MD simulation showing the evolution of a Pd<sub>91</sub> island on Ag(111) at 500 K over 2  $\mu$ s. The system reaches a quasi-equilibrium after 2  $\mu$ s. As depicted in the bottom schematic, Pd atoms are green, and Ag atoms are colored by the layer height, from red (level -2) to blue (level 2). The bottommost layer (level -3) is fixed in the simulation and hence omitted. (a) As-deposited hexagonal Pd<sub>91</sub> island on Ag(111). (b) Pd undergoes a series of place exchange with the underlying Ag atoms around the island edges. (c-d) A few Ag atoms climb on top of the deposit and attempt to pull Pd atoms up to level 2. (e-f) PdAg alloy layer forms at level 2. (g-h) In the second half of the simulation, more Ag atoms migrate out of the surface to further encapsulate the cluster, resulting in a vacancy pit. See video here.



Table 1: Four main classes of restructuring events underlying the restructuring of a Pd island on Ag(111): (a) Pd-Ag direct exchange; (b) Ag pop-out; (c) Ag hopping ascent; (d) Pd exchange ascent. The structures (IS = initial state; TS = transition state; IMS = intermediate state; FS = final state) are fully optimized by the GP model, followed by DFT evaluation of the relative energies. Slab atoms are semi-transparent for clarity.



In the beginning, Pd undergoes a series of place exchange with the underlying Ag atoms around the island edges [Fig. 3(b)]. Such Pd-Ag direct exchange [Table 1(a)] is the predominant mechanism by which Pd atoms are incorporated into the Ag surface in the earlier stages. First, a Ag atom migrates out of the surface layer into a kink or a corner site present at the island edge. As seen by a modest energy barrier of 0.3 eV, such edge vacancies facilitate the otherwise constrained exchange process. The Ag pop-out creates a vacancy in the surface layer, which is quickly filled by a neighboring Pd atom. This Pd insertion has an energy barrier of only 0.05 eV, due to a large thermodynamic driving force (1.3 eV stabilization).

Next, some of the Ag atoms, now occupying the edge sites, become mobile and climb on top of the Pd deposit [Fig. 3(b)]. These Ag atoms are mobilized around kinks and corners, allowing the island edges to be rather fluxional. Adatom diffusion on close-packed surfaces has been investigated extensively in the computational literature.<sup>38</sup> For example, our previous work has shown facile edge and corner diffusion of both Ag and Pd adatom on Ag, with energy barriers below 0.4 eV in all cases.<sup>21</sup> For the simplest available benchmark of adatom self-diffusion on Ag(111), our energy barrier of 51 meV is in good agreement with the experimental values reported in the range of 50-100 meV (see Lü *et al.*<sup>39</sup> and the references therein).

Step ascent and descent are well-established in the context of island growth.<sup>40</sup> There is a coordination loss and activation energy ( $E_a$ ) associated with step descent. As such, the island growth mode is determined by the Ehrlich-Schwoebel barrier ( $E_{ES}$ ),<sup>41,42</sup> defined as the additional activation energy required for step descent, relative to the terrace diffusion barrier ( $E_d$ ):

$$E_{ES} = E_a - E_d. \quad (1)$$

There are two main mechanisms of step ascent and descent: simple hopping over the edge [Table 1(c)], or exchange involving an insertion into the edge and a concurrent displacement of an edge atom [Table 1(d)]. Consistent with our previous work,<sup>21</sup> we find that descent is more facile at threefold-terminated steps (B-step) than at fourfold-terminated steps (A-step),

and that exchange is more facile than hopping. For the simplest case of Ag adatom self-diffusion, our B-step Schwoebel barrier is 120 meV, in good agreement with the experimental value of  $120 \pm 20$  meV.<sup>43</sup> To the best of our knowledge, no experimental benchmark exists for the diffusion of Pd adatom on Ag.

In our simulation, exchange ascent and descent, followed by their hopping counterparts, comprise the majority of the observed restructuring events (Fig. S6). Interestingly, Ag atoms ascend primarily via hopping in the earlier stages, whereas exchange ascent occurs only when a sufficient number of Ag atoms are already present on the top to participate in the exchange. It is important to note that descent is easier than ascent in general, due to the larger coordination present at the lower edge than at the upper edge. However, an ascending Ag atom is capable of being stabilized by Ag atom(s) that may already be present on the top. Such extra coordination can lower the energy barrier by up to 0.3 eV (Tables S4 and S5).

Surprisingly, the top Ag atoms pull up the underlying Pd atoms, ultimately forming a PdAg alloy layer in the second level [Fig. 3(c)]. This Pd ascent occurs primarily through a series of exchange involving the Ag atoms over a relatively short period of  $0.2 \mu\text{s}$ , with energy barriers below 0.5 eV [Table 1(d)]. The process is accompanied by a distinct relaxation of the deposit layer upon becoming coordinated by this second layer (Sec. 7, SI). Mechanistically, Pd hopping ascent occurs much less frequently, in contrast to Ag hopping ascent. In general, Pd atom experiences a larger energetic cost than Ag atom undergoing a given restructuring event, unless it is driven by a significantly large thermodynamic stabilization (e.g. Pd insertion into the surface).<sup>21</sup>

In the second half of the simulation, more Ag atoms migrate out of the surface to further cover the bilayer structure around its edges, resulting in a vacancy pit [Fig. 3(d)]. In all cases, Ag mass balance is conserved, and the number of Ag atoms that migrate out of the surface is roughly proportional to the number of Pd atoms deposited, e.g. 70 Ag atoms migrate out of the surface in the case of Pd<sub>91</sub> island (Fig. 3). As such, larger Pd islands

lead to a larger amount of Ag transported out of the surface, creating larger vacancy pits. Because our Pd<sub>91</sub> island model is much smaller than the ones present in the experiment, the resulting vacancy pit is also much smaller and localized in our simulation. In contrast, the high Pd coverage (0.4 ML) and large island sizes ( $\sim 50$  nm wide) in our experiment induce almost all of the Ag atoms in the surface layer to be consumed in encapsulating and alloying into the Pd islands, leaving behind extensive and well-connected vacancy pits (Fig. 2). At a lower coverage of 0.15 ML, however, a small number of localized vacancy islands can be seen upon partial restructuring of a few as-deposited Pd islands at room temperature (Fig. S23). This observation indicates that the overall restructuring process is driven by vacancy island growth as seen in our simulation.

Our previous work found that Ag pop-out is very unfavorable on clean terraces, with energy barriers over 1.0 eV in all scenarios.<sup>21</sup> Surprisingly, a single-atom Ag pop-out next to an island edge has a relatively modest energy barrier of 0.6 eV [Table 1(b)], which is attributed to the stabilization of the transition state by the edge atoms. The resulting vacancy pits create double steps through which facile Ag migration occurs, as proposed in previous experiments of rapid vacancy island decay.<sup>44,45</sup> Remarkably, the thermodynamic driving force underlying this Ag transport becomes even more pronounced at higher Pd coverages (Sec. 6.3, SI). For example, at 0.75 ML coverage, the surface area of the exposed Ag is less than that of the deposited Pd. Consequently, additional Ag migration begins to occur out of the subsurface layers, resulting in multiple-height steps and significant surface coarsening (Fig. S18).

At a relatively moderate temperature, the system has evolved into a completely different structure from the as-deposited Pd monolayer in a period of microseconds. At 500 K, the system reaches an apparent quasi-equilibrium after 2  $\mu$ s in this metastable structure, as opposed to the thermodynamically favored state with Pd dispersed in the bulk. In particular, the exposed Pd clusters in the second layer are now much smaller and interspersed by Ag atoms. The observed structure is consistent with low-temperature CO probe experiments –

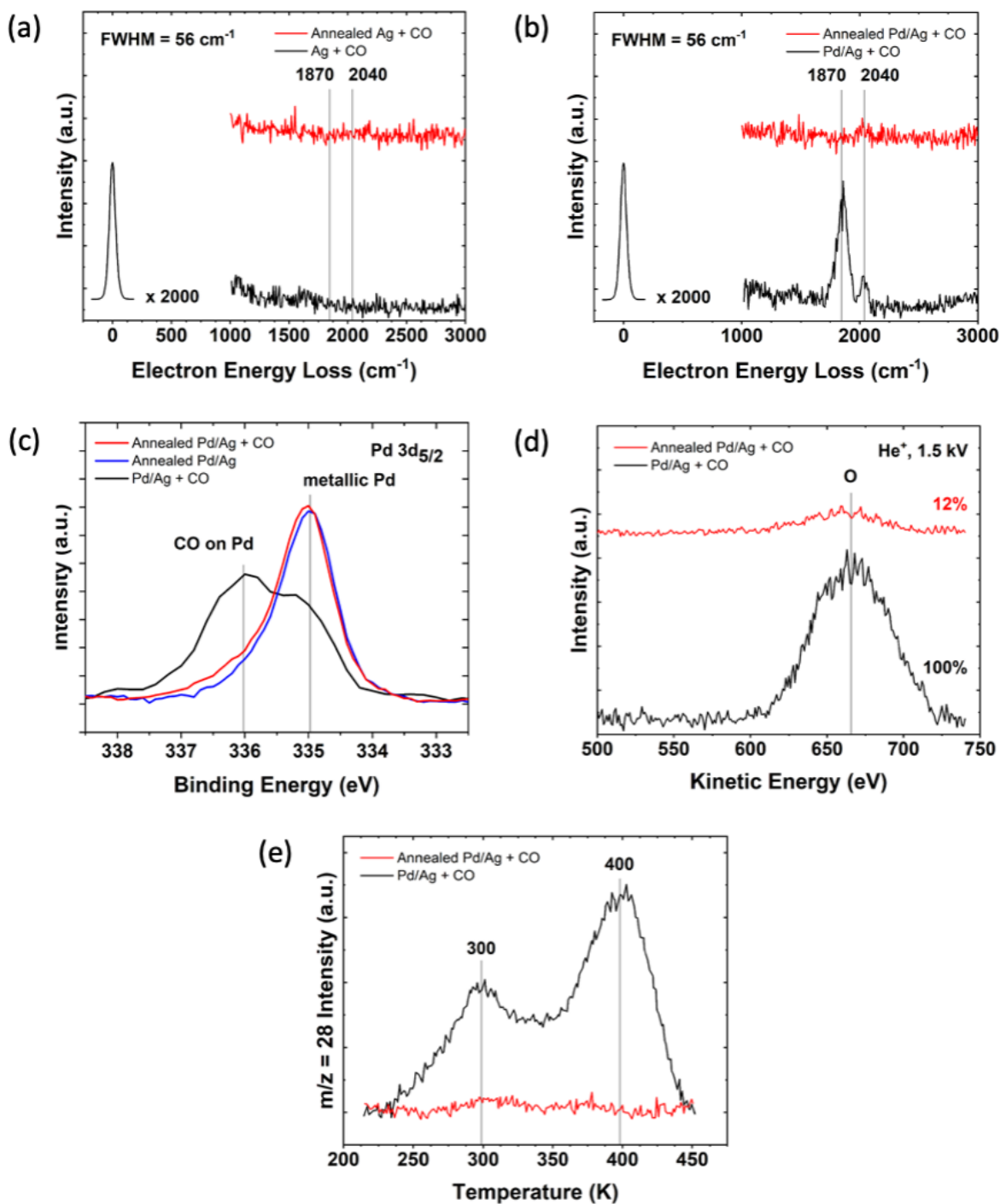


Figure 4: Adsorption of CO probes to detect the presence of surface Pd on 0.4 ML Pd/Ag(111). CO exposures are 100 L (1 Langmuir =  $1.0 \times 10^{-6}$  Torr s<sup>-1</sup>) at 100 K. Surfaces are annealed by holding the sample at 450 K for 1 min. (a) HREELS: Ag(111), as-prepared (black) and annealed (red), does not exhibit appreciable CO adsorption. (b) HREELS: Pd/Ag(111), as-deposited (black) and annealed (red), exhibits CO adsorption on Pd bridge sites (1870 cm<sup>-1</sup>) and Pd top sites (2040 cm<sup>-1</sup>). (c) XPS: Clean Pd/Ag(111) annealed (blue); CO exposure to Pd/Ag(111), as-deposited (black) and annealed (red). (d) LEISS: Quantitative CO coverage on Pd/Ag(111), as-deposited (black) and annealed (red). (e) TPD: CO desorption from Pd/Ag(111), as-deposited (black) and annealed (red).

comprised of XPS, high resolution electron energy loss spectroscopy (HREELS), low energy ion scattering spectroscopy (LEISS), and temperature programmed desorption (TPD) – that collectively demonstrate significant reduction in CO binding after annealing at 450 K (Fig. 4).

The presence of surface Pd can be determined by probing the ability of CO to bind to the surface. CO adsorption occurs up to 100 K and 450 K on Ag(111) and Pd(111), respectively.<sup>46</sup> As such, we expose 100 L (1 Langmuir =  $1.0 \times 10^{-6}$  Torr s<sup>-1</sup>) of CO to our surfaces at 100 K. The surfaces are prepared by depositing 0.4 ML of Pd on Ag(111) at 300 K, which is further annealed at 450 K for 1 min. It is important to note that the CO exposure is performed at a low temperature of 100 K to limit thermal mobility of the system and thereby prevent any CO-induced restructuring of the surface, most notably reverse segregation of Pd to the surface. Previous studies have demonstrated such restructuring upon CO exposure at room temperature and above, for bimetallic systems including Pd/Ag,<sup>10</sup> Pt/Au,<sup>47</sup> Ni/Au,<sup>48</sup> and Co/Au.<sup>49</sup>

CO does not adsorb appreciably on either as-prepared or annealed Ag(111) [Fig. 4(a)]. In contrast, CO adsorption occurs readily on as-deposited Pd/Ag(111), as evidenced by a positive core-level shift in the Pd  $3d_{5/2}$  region, from 335.0 eV of metallic Pd to 336.0 eV upon CO exposure [Fig. 4(c); blue to black].<sup>10,50</sup> This observation is further supported by the elastic peak of oxygen arising from CO binding to Pd through its carbon and exposing its oxygen to the spectrometer [Fig. 4(d); black].<sup>51</sup> Specifically, CO adsorbs on Pd bridge sites and Pd top sites with vibrational frequencies of 1870 and 2040 cm<sup>-1</sup>, respectively, indicating the presence of extended Pd on the surface [Fig. 4(b); black].<sup>46</sup> CO desorption at 300 K and 400 K further confirms the presence of CO adsorbed on Pd sites [Fig. 4(e); black].<sup>46</sup>

However, CO adsorption is minimal on annealed Pd/Ag(111), as evidenced by a very small positive core-level shift in the Pd  $3d_{5/2}$  region upon CO exposure, centered at 335.0 eV of metallic Pd [Fig. 4(c); blue to red].<sup>10,50</sup> From quantification of the elastic peak of oxygen, only 12% of CO adsorbs on the annealed surface [Fig. 4(d); red], compared to the as-deposited surface, in very good agreement with our simulation predicting  $\sim 7\%$  of exposed Pd after

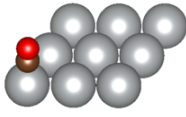
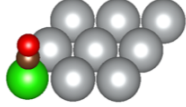
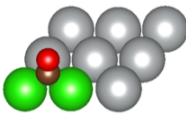
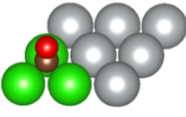
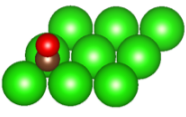
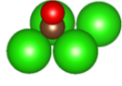
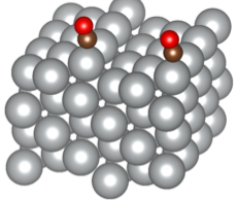
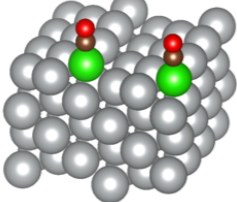
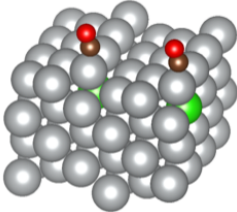
annealing. Minimal CO desorption at 300 K and 400 K confirms a significant decrease in the surface concentration of Pd [Fig. 4(e); red]. In particular, only a trace amount of CO adsorption is detected on Pd top sites ( $2040 \text{ cm}^{-1}$ ) [Fig. 4(b); red], suggesting that the annealed surface consists mostly of isolated Pd atoms and small Pd ensembles.

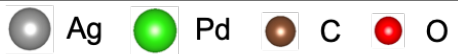
These experiments are also supported by DFT calculations showing a systematic reduction of CO adsorption energy upon decreasing the surface concentration of Pd (Table 2). Based on carefully benchmarked results (Sec. 8, SI), the adsorption is dramatically weakened by the geometric effect, from  $-1.76 \text{ eV}$  on an extended Pd(111) to  $-0.94 \text{ eV}$  on an isolated Pd monomer in Ag(111), ultimately reaching  $-0.10 \text{ eV}$  on a clean Ag(111). The same trend holds at the step edges, where the adsorption is slightly stronger by  $\sim 0.2 \text{ eV}$  than on the terraces. Comparing an extended Pd(111) with a Pd trimer in Ag(111), the electronic effect also weakens the adsorption by  $\sim 0.4 \text{ eV}$ . Overall, these validations attest to the predictive power of our modeling method, complementary to experimental techniques with surface and chemical sensitivities.

At higher temperatures (Sec. 6.1, SI), the bilayer structure becomes further encapsulated by a third Ag capping layer, followed by layer-by-layer dissolution of Pd over a longer period of time (Fig. S12; see video here). The dissolution consists of multiple interlayer transfer events (Table S6), starting at the edges and moving toward the center of the island. A subsurface Ag atom first ascends into a pre-existing surface vacancy, after which a neighboring Ag-Pd pair descends into the resulting subsurface vacancy. As Pd dissolves layer-by-layer, it is no longer thermodynamically favorable for the Ag atoms to remain in the capping layer. As such, they migrate back into the surface, resulting in a gradual decay of vacancy pits. In the end, all pits are refilled, and almost all Pd remains dissolved in the subsurface, with a small amount of Pd resurfacing transiently as isolated atoms. Such surface-subsurface equilibrium is in direct agreement with the thermodynamic model of the quasi-chemical approximation.<sup>33</sup>

The formation of metastable Ag-encapsulated Pd islands is predominant at moderate

Table 2: CO adsorption energy ( $E_{\text{ads}}$ ) on Pd/Ag model surfaces, calculated with DFT using the M06L functional.<sup>52</sup> All structures are fully optimized. The experimental benchmark for 1/4 ML CO/Pd(111) is  $-1.49$  eV.<sup>53</sup>

Surface model	CO coverage (ML)	Binding site	Structure	$E_{\text{ads}}$ (eV)
Ag(111)	1/9	Top		-0.10
Pd monomer in Ag(111)	1/9	Top		-0.94
Pd dimer in Ag(111)	1/9	Bridge		-1.16
Pd trimer in Ag(111)	1/9	Hollow		-1.37
Pd(111)	1/9	Hollow		-1.76
Pd(111)	1/4	Hollow		-1.68
Ag(211)	1/12	Edge top		-0.24
Edge Pd in Ag(211)	1/12	Edge top		-1.14
Subsurface Pd in Ag(211)	1/12	Edge top		-0.26





temperatures and always precedes layer-by-layer dissolution of Pd. Such behavior has vital implications for catalysis. Our previous work showed that Pd resurfaces at moderate temperature and pressure of CO or O<sub>2</sub>.<sup>10</sup> This reverse segregation allows the catalytically active element to be present on the surface of an inert host metal under reaction conditions. Our results suggest that such activation of the catalyst would be enabled by metastable structures, where Pd remains largely accessible to the reactants in the form of Ag-encapsulated Pd rather than bulk-dissolved Pd. Evolution of these metastable structures would play a central mediating role in engineering of catalytic activity through various pretreatment strategies.<sup>54–57</sup>

## Conclusions

For the first time, a detailed atomistic picture of the long-timescale restructuring of Pd on Ag is revealed, by combining microscopy, spectroscopy, and novel simulation methods. Because of its higher surface energy, Pd prefers to dissolve into the Ag host,<sup>10</sup> but the underlying mechanism and timescale have remained speculative at best. We directly uncover both the mechanism and the timescale of the restructuring by developing and performing accelerated machine-learning molecular dynamics, in combination with surface science experiments. Using our newly developed automated analysis method, previously unidentified restructuring events have been discovered and characterized in an unbiased fashion, including Pd-Ag place exchange and Ag pop-out, as well as step ascent and descent. Remarkably, a significant amount of Ag migrates out of the surface to cover and alloy into the Pd islands, resulting in extensive vacancy pits within a period of microseconds. The microscopic insights established in this study are broadly applicable to other bimetallic systems comprised of soft metals, where surface segregation is primarily driven by the difference in the surface energy of the individual components.

Our automated modeling strategy within machine-learning molecular dynamics, in combi-

nation with surface science experiments, has allowed complex surface restructuring dynamics to be mapped out at atomic resolution, a task that would otherwise be intractable with a brute-force static approach. In this paper, the mechanisms for the reconstruction of a pristine Pd/Ag(111) is demonstrated as a first step toward understanding the elementary events that lead to surface rearrangement. Investigation of more complex restructuring phenomena involving adsorbates is underway. In particular, the relatively low energy barriers of the discovered restructuring events suggest the possibility of catalytic processes proceeding on mobile surfaces, as seen in recent computational studies of nanocluster catalysis.<sup>58-60</sup> The reactivity and the state of the surface may evolve together within a network of competition between thermodynamics and kinetics.

## Experimental Section

*Automated characterization of surface restructuring events:* A fully automated workflow for surface restructuring event characterization has been developed (Fig. 1), building upon our previous framework.<sup>21</sup> All of our interactive Jupyter Notebook files are publicly available.<sup>61</sup> Density functional theory (DFT) data is used to train a Gaussian process (GP) machine-learning force field,<sup>23</sup> which allows us to perform large-scale and long-timescale molecular dynamics (MD) at first-principles accuracy (Sec. 1, SI). From preliminary visual inspections, we find that most restructuring events occur over 1-3 ps, with atoms transitioning between well-defined face-centered cubic (FCC) lattice sites. To better follow the events, we first "clamp" the raw MD trajectory: All atoms are assigned onto discrete FCC lattice sites, frame by frame<sup>24-26</sup> (Sec. 3, SI). The resulting clamped trajectory is automatically scanned to detect restructuring events. A frame is flagged as an event point if the  $z$ -coordinate (normal to the surface) of any atom changes irreversibly over 1-2 ps. The flagged frames within 2 ps of each other are clustered into distinct events. Then, the events are automatically sorted into seven main classes,<sup>21</sup> based on changes in the atomic layer and

coordination number (Sec. 4, SI). Finally, pathways for representative events of each class are optimized using transition state modeling<sup>27-29</sup> (Sec. 2, SI).

*Surface science experiments:* The experimental measurements are performed with three different Ag(111) crystals mounted in three separate ultrahigh vacuum (UHV) chambers, described in detail elsewhere.<sup>10,62</sup> Scanning tunneling measurements are performed in a UHV chamber using a beetle-type scanner (RHK UHV VT STM).<sup>10</sup> X-ray photoelectron and ion scattering measurements are performed in a UHV chamber using an X-ray source (Perkin Elmer 04-548/32-095 with RBD 20-042) with (non-monochromatized) Mg K $\alpha$  radiation, a 1.5 eV 4He<sup>+</sup> ion beam (SPECS IQE 12/38) with a Wien mass filter (SPECS WF-IQE), and a hemispherical analyzer (SPECS Phoibos 100 equipped with a 5-channel electron multiplier).<sup>10</sup> The high-resolution electron energy loss measurements are performed in a UHV chamber using an electron energy loss spectrometer (LK 2000) with a primary energy of 4.5 eV at 60° specular geometry.<sup>62</sup> The Ag(111) crystals are cleaned by repeated cycles of Ar<sup>+</sup> sputtering (1-2 kV) and annealing (750-800 K) in UHV. The PdAg surface alloys are formed by depositing Pd (99.99%) at room temperature on clean Ag(111) using an electron-beam evaporator. CO adsorption experiments use a mixture of 20% CO (99.9% purity) in Ar.

## Additional information

Supporting Information: Details of Gaussian process molecular dynamics; event energetics; clamping algorithm; event classification & statistics; validation of the slab model; evolution of layer-by-layer compositions (variation of temperature, Pd cluster size, and Pd coverage); evolution of layer-by-layer average interatomic distances; functional benchmark & CO adsorption energy; and additional STM data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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## Competing financial interests

The authors declare no competing financial interests.

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# Graphical TOC

