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Subsurface Stabilization of Interstitial Pt Atoms on CeO₂(111): Rethinking Single-Atom Catalyst Architectures

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ABSTRACT

Single-atom catalysts (SACs) offer maximal efficiency by stabilizing isolated metal atoms on oxidic supports. Platinum on cerium oxide (CeO₂) is a key SAC system, where adsorbed CO typically exhibits red-shifted vibrational modes due to Pt back-donation. Using polarization-resolved IR spectroscopy on Pt-deposited CeO₂(111) single-crystal surfaces, we do not observe CO vibrational bands below 2140 cm⁻¹ at low coverages, indicating that the surface-bound Pt atoms are not present in detectable amounts. DFT calculations demonstrate that this unexpected observation is consistent with Pt atoms occupying buried interstitial sites. Such subsurface single-atom sites, not considered in previous studies, are thermodynamically favored at low coverages, adopt an unusual oxidation state, and are inaccessible to direct CO binding. Our findings challenge the prevailing assumption that single atoms remain surface-bound and highlight the critical role of subsurface interstitial species, prompting a rethinking of how active sites in single-atom catalysts are stabilized on reducible oxides.

1 | Introduction

Single-atom catalysis (SAC) has emerged as a powerful paradigm in heterogeneous catalysis, offering both maximal atom efficiency and access to unique reaction pathways through isolated metal sites [1–8]. These systems often surpass traditional nanoparticle-based catalysts in activity, selectivity, and thermal stability, making SAC a promising route toward more efficient and sustainable catalytic technologies.

For certain oxide substrates, such as Fe₃O₄, the presence of single, surface-bound metal atoms has been clearly demonstrated

[2]. However, for many other systems, including cerium dioxide (CeO₂), the nature and stability of isolated metal sites remain under debate. Among these, platinum supported on CeO₂ (Pt–CeO₂) stands out due to the strong electronic and structural interplay between Pt atoms and the redox-active oxide support [9–13].

Over the past decade, Pt–CeO₂ single-atom catalysts have been the focus of intense research, particularly in key reactions such as CO oxidation, the water–gas shift reaction, and selective hydrogenations [14–17]. However, despite this progress, fundamental questions about structural dynamics and reaction mechanisms at

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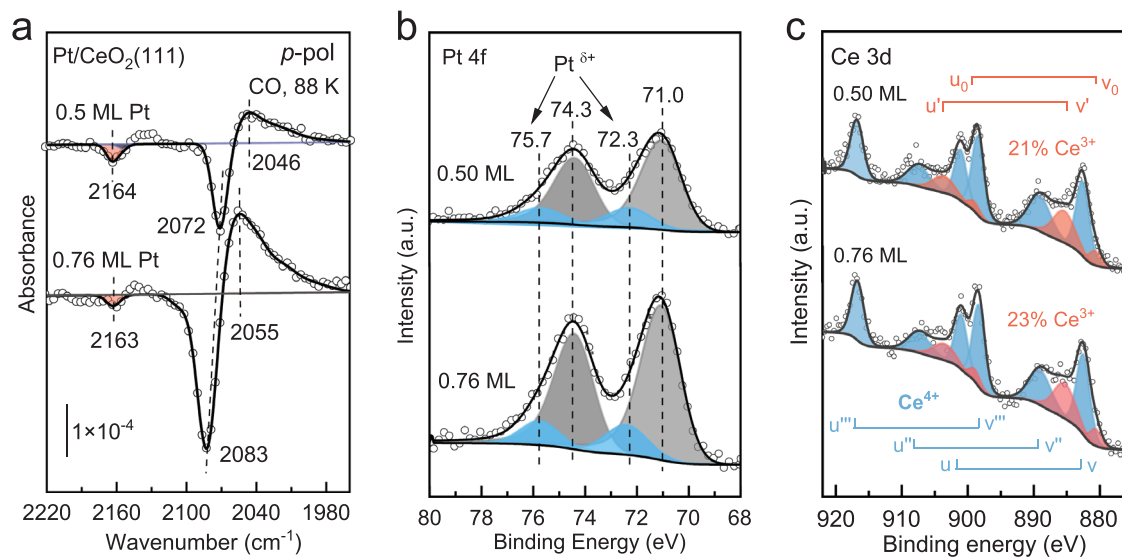


FIGURE 1 | Polarization-resolved IRRAS and grazing-emission XPS characterization of Pt clusters/particles deposited on CeO₂(111) single crystal surfaces. (a) p-polarized IRRAS data recorded after CO saturation adsorption at 88 K on the Pt-deposited CeO₂(111) surfaces at high Pt coverages (0.50 and 0.76 ML); (b) deconvoluted Pt 4f and (c) Ce 3d spectra of 0.50 ML and 0.76 ML Pt/CeO₂(111).

the atomic scale remain unresolved. The identity of the active site at low Pt coverages, in particular, is highly controversial: while some studies attribute the activity to isolated Pt atoms on the CeO₂ surface [9, 17–19], others present evidence that contradicts this view [20–24]. To address these challenges, infrared (IR) spectroscopy using carbon monoxide (CO) as a vibrational probe has proven invaluable, offering detailed insight into the electronic structure and coordination environment of surface species [25–31]. Among the techniques available to bridge the pressure and materials gaps in surface science, CO-based IR spectroscopy is especially versatile. Vibrational bands observed under operando conditions in diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) on powders can be rigorously assigned using Infrared reflection–absorption spectroscopy (IRRAS) data obtained from model substrates and validated by theoretical calculations [32, 33].

The assignment of CO stretching frequencies in Pt–ceria catalysts remains debated, as red-shifted bands attributed to Pt single atoms often overlap with signals from Pt clusters. Polarization-resolved IRRAS on oxide single crystals provides the spectral precision needed to benchmark theoretical predictions [33, 34], but its application has been limited by the inherently low reflectivity of dielectric substrates. To date, no such measurements have been reported for Pt on CeO₂, leaving the vibrational signature of CO bound to isolated Pt atoms experimentally unresolved.

Here we present a comprehensive analysis of CO-IRRAS data for Pt deposited on CeO₂(111) single-crystal surfaces recorded using both p- and s-polarized light. Pt coverages were precisely quantified by quartz crystal microbalance (QCM) and x-ray photoelectron spectroscopy (XPS), while the fraction of uncovered ceria was tracked via the CO vibrational band characteristic of pristine CeO₂(111). Particular focus is placed on Pt single atoms, with interpretation of the IRRAS data supported by CO-SLIR (surface-ligand IR) measurements [31, 32] and density functional theory (DFT) calculations.

2 | Results and Discussion

2.1 | Pt Clusters/Particles Deposited on CeO₂(111)

Figure 1a shows polarization-resolved IRRAS data collected after CO saturation at 88 K on Pt-deposited CeO₂(111) surfaces with high Pt coverages of 0.50 and 0.76 monolayer (ML). Pt/CeO₂(111) samples were prepared by well-controlled deposition of Pt at 300 K, followed by annealing in UHV at 700 K for 10 min (for details, see Supporting information). Here, 1 ML of Pt is defined as one Pt atom per surface unit cell of CeO₂(111), corresponding to 7.9×10^{14} atoms/cm². The clean CeO₂(111) surface exhibits a sharp band at 2154 cm⁻¹, blue-shifted relative to the gas-phase CO value at 2143 cm⁻¹, which is assigned to CO bound to sevenfold-coordinated surface Ce⁴⁺ sites (Ce_{7c}⁴⁺) [35]. Following Pt deposition at 0.50 and 0.76 ML, broad red-shifted CO bands appear at 2100–2000 cm⁻¹, assigned to CO bound to Pt clusters or particles, consistent with previous reports. [36–39] As these Pt-related IR signals grow, the 2154 cm⁻¹ feature of the pristine CeO₂(111) surface diminishes, indicating progressive Pt cluster formation and surface coverage. XPS analysis (Figure 1b,c) confirms that at 0.75 ML, Pt is predominantly metallic (4f_{7/2}/4f_{5/2} doublet at 71.0/74.3 eV), with only a minor contribution from positively charged Pt^{δ+} species (72.3/75.7 eV) (Figure 1b).

Interestingly, the p-polarized IRRAS data show a splitting of the Pt-related CO bands with an unusual sign reversal (see Figures 1a and 2a). The negative features at 2072–2083 cm⁻¹ are due to upright CO on high-coordinated Pt atop sites on terrace planes, excited by the normal component of the p-polarized IR radiation ($E_{p,n}$, see Figure 2b,c). In contrast, the positive bands centered at 2048–2055 cm⁻¹, not reported previously [29], are characteristic of tilted CO on undercoordinated Pt atop sites at particle edges and corners, excited by the tangential component ($E_{p,t}$, see Figure 2b,c). These assignments are further supported by additional IRRAS spectra recorded in s-polarization (see Figure 2a)—an approach not feasible to oxide thin films on

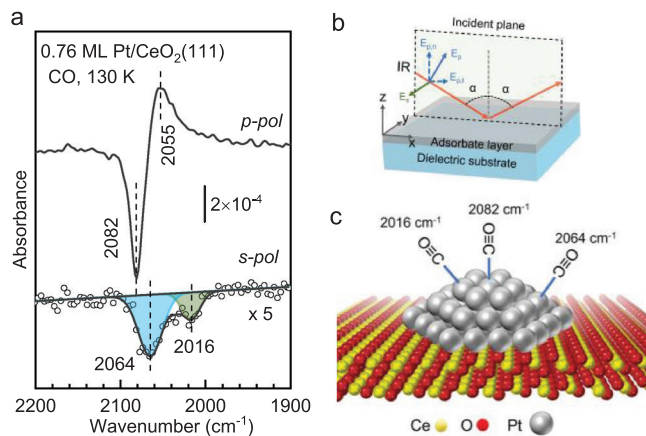


FIGURE 2 | Pt clusters/particles deposited on $\text{CeO}_2(111)$ single-crystal surfaces characterized by polarization-resolved IRRAS. (a) p- and s-polarized IRRAS data recorded after CO saturation adsorption at 130 K on the Pt-deposited $\text{CeO}_2(111)$ surfaces with a high Pt coverage of 0.76 ML. (b) Schematic representation of the interaction between incident IR light and molecular adsorbates on dielectric substrate surfaces. (c) Schematic illustration of distinct CO species adopting site-specific adsorption geometries: upright adsorption on high-coordinated terrace Pt sites and tilted adsorption on undercoordinated edge and corner sites.

metallic substrates due to the surface selection rule [40, 41]. The s-polarized spectrum reveals two well-resolved negative bands at 2064 and 2016 cm^{-1} , which originate from CO bound to edge and corner Pt atoms, respectively, with the molecular axis oriented predominantly parallel to the substrate and characterized by reduced coordination numbers (Figure 2c).

Temperature-dependent IRRAS measurements allowed to estimate the binding energy of these red-shifted CO species, which amount to about 1.55 eV (Supporting Information, Figure S1). The

strong interaction between CO and Pt clusters/particles arises from the significant electronic back-donation from Pt 5d states into the CO $2\pi^*$ antibonding orbital, in line with the substantial red-shift of the CO stretching vibrations.

2.2 | Pt Single Atoms Deposited on $\text{CeO}_2(111)$

While the results at higher coverages (>0.5 ML) are consistent with previously reported Pt cluster/particle formation, the low-coverage regime reveals unexpected behavior. As shown in Figure 3a, the red-shifted Pt-related CO bands ($2100\text{--}2000$ cm^{-1}) vanish entirely below 0.12 ML, even though QCM and XPS unambiguously confirm the presence of Pt (Figure 3b). The effect of Pt deposition is further demonstrated by attenuation of the pristine 2154 cm^{-1} peak and the emergence of a previously unreported band at 2169 cm^{-1} (Figure 3a). The new 2169 cm^{-1} CO species is only weakly bound (< 0.3 eV from temperature-dependent IRRAS), in pronounced contrast to CO adsorption on Pt clusters. Importantly, no additional CO bands indicative of surface defect sites are observed. XPS (Figure 3b,c) reveals Pt^{d+} species ($d < 2$) together with $\sim 15\%$ Ce^{3+} , leading us to hypothesize that, in this low-coverage regime, isolated Pt atoms do not remain on the surface but migrate into subsurface sites.

Upon increasing the Pt coverage to 0.25 ML (Figure 3a), the 2169 cm^{-1} band shifts slightly to 2165 cm^{-1} and becomes predominant, while a weak red-shifted, negative CO band appears at 2062 cm^{-1} , characteristic of CO bound to low-coordinated Pt sites in small clusters. The corresponding Pt 4f (Figure 3b) and Ce 3d (Figure 3c) XPS data show that Pt remains predominantly in the Pt^{d+} state, with the fraction of reduced Ce^{3+} species increasing slightly to 19%.

Figure 4 summarizes the integrated intensity evolution of various CO bands as a function of Pt coverage, determined by QCM.

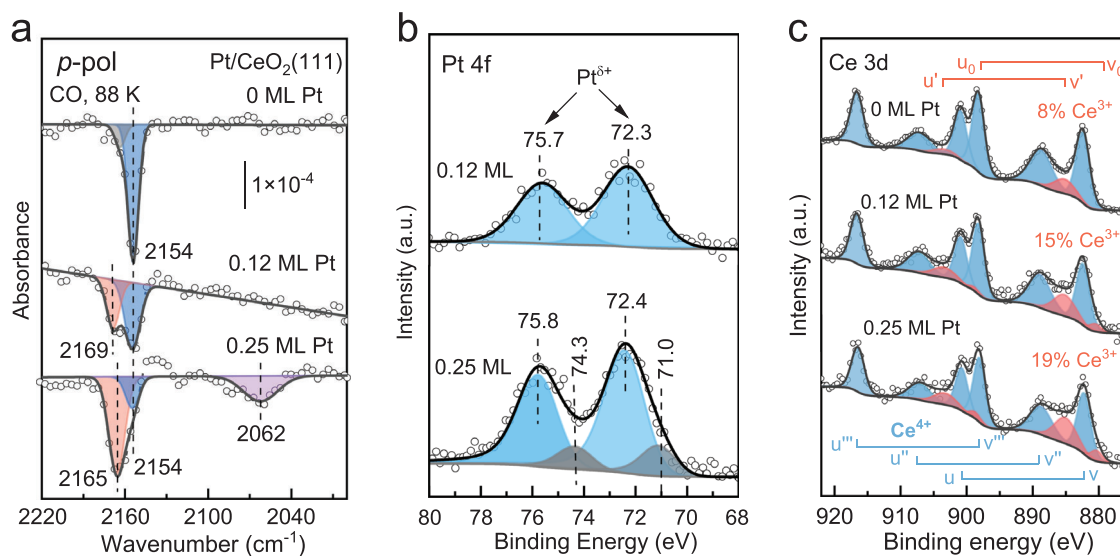


FIGURE 3 | Polarization-resolved IRRAS and grazing-emission XPS characterization of Pt single atoms and small clusters deposited on $\text{CeO}_2(111)$ single-crystal surfaces. (a) p-polarized IRRAS data recorded after CO saturation adsorption at 88 K on the pristine and Pt-deposited $\text{CeO}_2(111)$ surfaces at low Pt coverages (0.12 and 0.25 ML); (b) deconvoluted Pt 4f spectra of 0.12 and 0.25 ML Pt/ $\text{CeO}_2(111)$; (c) deconvoluted Ce 3d spectra of pristine and Pt (0.12, 0.25 ML) deposited $\text{CeO}_2(111)$.

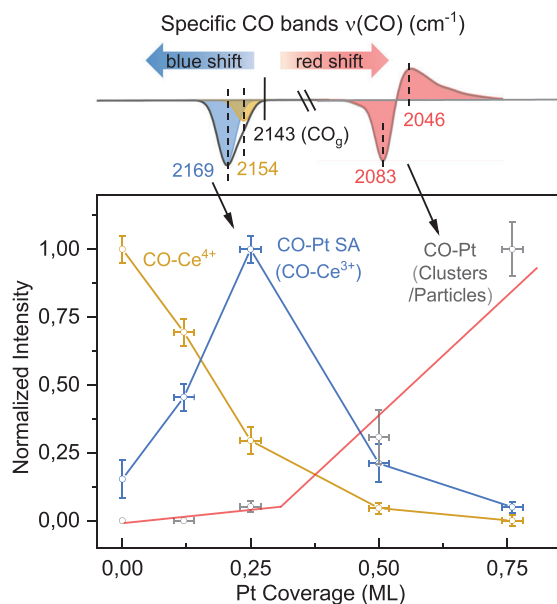


FIGURE 4 | Quantitative analysis of distinct CO vibrations. Coverage-dependent intensity evolution of specific CO bands characteristic of distinct adsorption sites: Pt single atoms (CO-Pt SA, $\sim 2169\text{ cm}^{-1}$); Pt small clusters and particles (CO-Pt, $2100\text{--}2000\text{ cm}^{-1}$) surface Ce cations (CO-Ce⁴⁺, 2154 cm^{-1}). The Pt coverage was determined by QCM and XPS (see text).

(i) The 2154 cm^{-1} band, reflecting the fraction of the CeO₂(111) surface not covered by Pt, decreases rapidly with increasing coverage. (ii) CO bands attributed to Pt clusters and particles ($2100\text{--}2000\text{ cm}^{-1}$) vanish almost completely at coverages below 0.25 ML. (iii) The CO band associated with Pt single atoms at 2169 cm^{-1} reaches its maximum intensity at 0.25 ML and then declines sharply at higher coverages. Spectroscopic analysis indicates the coexistence of Pt single atoms and small Pt clusters at intermediate coverages. Since this blue-shifted frequency has not been reported previously, the location of the deposited Pt atoms cannot be inferred directly from the band position alone, necessitating a thorough theoretical analysis.

2.3 | Theoretical Results

Initial calculations for a bare (2×2) CeO₂(111) unit cell show CO binding to Ce_{7c}⁴⁺ sites with $E_{\text{ads, CO}} = -0.21\text{ eV}$ and $\nu(\text{CO}) = 2150\text{ cm}^{-1}$ (Figure S2), consistent with previous reports [34]. Geometry optimization of a surface-bound Pt atom on CeO₂(111) at 0.25 ML coverage (one Pt per (2×2) unit cell) identifies a bridge site above two surface O atoms as the most stable configuration (Figures S2 and S3), also consistent with earlier studies [29]. In this state, Pt carries a formal charge of +1.3 (Table 1) and binds CO with $E_{\text{ads, CO}} = -2.28\text{ eV}$ and $\nu(\text{CO}) = 2047\text{ cm}^{-1}$ (Table 2). While this red-shift matches previous experimental IR data, it does not account for the newly observed blue-shifted, weakly bound CO species.

To resolve this discrepancy with the experimental data, subsurface Pt configurations were explored. Since in earlier work [20] we reported that replacing Ce atoms in the ceria bulk with Pt carries a high energy penalty (1.62 eV, see also Refs

TABLE 1 | Calculated stabilities and Pt oxidation state of different Pt/CeO₂(111) structures relative to Pt_{bulk} and CeO₂(111) in eV.

	Pt coverage	Pt oxidation state ^a	ΔE
Pt@CeO ₂ (111) in subsurface	1/4 ML	+1.9	+1.61
Pt@CeO ₂ (111) in subsurface	1/9 ML	+1.9	+1.23
Pt@CeO ₂ (111) on top surface	1/4 ML	+1.3	+1.84
Pt@CeO ₂ (111) on top surface	1/9 ML	+1.3	+1.77

^aDerived from Bader charge analysis (Figure S4 and Table S1a,b).

[11, 42, 43].), we focused on interstitial rather than substitutional sites. Although such configurations have rarely been considered in previous theoretical work, we found—surprisingly—that a Pt atom occupying the empty space between O²⁻ ions in the bulk interstitial is energetically more favorable by -0.23 eV (see Table 1) than on-surface sites. Using a larger (3×3) unit cell (1/9 ML coverage), representative of the experimental regime (0.12 ML), this stabilization increases to -0.54 eV , with Pt adopting a formal charge of +1.9 (Figure S4 and Table S1a,b).

In this unusual interstitial geometry (see Figure 5), the Pt atom is buried beneath the CeO₂(111) surface in a fourfold coordination with surrounding O²⁻ ions, rendering it inaccessible to CO. As a consequence, a nearby surface Ce⁴⁺ cation is displaced upwards by 1.3 \AA (see Figure 5c), becoming reduced to Ce³⁺ together with an adjacent Ce atom (Figure S5). This redox process facilitates the oxidation of Pt⁰ to Pt^{1.8+}. CO binds to the protruding Ce³⁺ site with a slightly tilted geometry (74°), showing an adsorption energy of $E_{\text{ads, CO}} = -0.29\text{ eV}$ and a vibrational frequency of $\nu(\text{CO}) = 2174\text{ cm}^{-1}$ (Table 2), closely matching our experimental data. Notably, the calculated oxidation state of subsurface Pt aligns well with XPS measurements, while surface-bound Pt remains less oxidized. Furthermore, we find that CO adsorption on distinct positively charged surface Pt single-atom sites at coverages of 0.12 and 0.25 ML—including configurations with one or two CO molecules per Pt site—leads to a red shift in the calculated stretching frequencies, in marked contrast to our experimental observations (see Table 2).

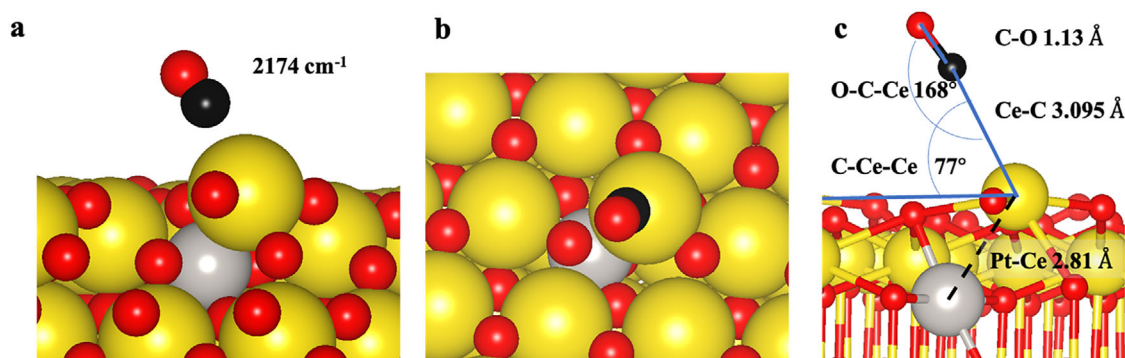
The 0.38 eV difference in subsurface interstitial Pt stability between 1/4 and 1/9 ML (Table 1) arises from strain and relaxation. In the smaller (2×2) cell, overlapping displacement fields hinder subsurface insertion, whereas the larger (3×3) cell better accommodates lattice distortions, stabilizing subsurface Pt at low coverage.

To assess the thermal stability of Pt single atoms on CeO₂, it is essential to compare their energies to bulk Pt as a reference state (see Table 1). On this basis, all configurations studied—both surface and subsurface—are thermodynamically unstable with respect to bulk Pt. However, subsurface Pt is consistently more stable than surface-bound Pt atoms, and this relative stability increases further at lower coverages.

The computed activation barrier for Pt diffusion into the subsurface interstitial site (2.03 eV, see Supporting Information, Figures S6 and S7) is consistent with the mild 700 K annealing applied

TABLE 2 | Calculated CO binding energies (in eV), vibrational frequencies (scaled and in cm^{-1}), and vibration intensities for Pt/CeO₂(111) with Pt single atoms on the surface and in subsurface interstitial sites. CO has been scaled with a factor of 1.008.

	ΔH_{CO}	Vibration frequency (cm^{-1})	Intensity ($(\text{D}/\text{\AA})^2/\text{amu}$)
1/4 ML Pt—in subsurface	-0.30	2156	3.6426
1/4 ML Pt—on top surface	-2.28	2047	32.6469
1/4 ML Pt—on top surface ^a	-1.46 ^b	2120/2050 ^c	19.6347/0.0764 ^d
1/9 ML Pt—in subsurface	-0.29	2174	3.9740
1/9 ML Pt—on top surface	-2.35	2031	38.0939
1/9 ML Pt—on top surface ^a	-1.29 ^b	2118/2067 ^c	24.2305/0.0007 ^d

^aAdsorption of 2 CO molecules^bDifferential heat of adsorption of second CO molecule^cSymmetric and asymmetric CO vibrations^dIntensities of symmetric and asymmetric CO vibration modes.**FIGURE 5** | DFT-optimized structure of single-atom Pt/CeO₂(111) model catalyst. (a) Side view, (b) top view, and (c) specific bond distances and angles of the optimized geometry of CO on Ce atoms on top interstitial Pt stabilized in the subsurface of CeO₂(111).

after deposition, comparable to temperatures used in ceria-based catalysts. Assuming a standard attempt frequency of 10^{13} s^{-1} , Pt incorporation at 700 K is expected to occur within 1 min, well below the experimental annealing time.

3 | Conclusion

Our combined IR spectroscopic and DFT study reveals that, contrary to prevailing assumptions, single Pt atoms deposited on CeO₂(111) surfaces are not present in detectable amounts at the surface at low coverages, consistent with migration into subsurface sites. This previously unrecognized configuration gives rise to a distinct, blue-shifted CO vibrational band, reflecting a weakly bound CO species and minimal back-donation from the embedded Pt atom. Thermodynamically, buried interstitial Pt is predicted to be more stable than surface-bound Pt, particularly at low coverages, a trend supported by both experiment and theory. These findings redefine the structural landscape of single-atom catalysts on reducible oxides and underscore the important role of considering subsurface incorporation, in particular at interstitial sites, as an intrinsic design parameter. Given the generality of strain, redox, and coordination effects across oxide supports, similar subsurface interstitial stabilization may occur for other metal-oxide combinations [44], suggesting a broader

need to reassess surface-bound assumptions in the design of the next-generation SACs. The catalytic activity of such subsurface interstitial sites will be addressed in future work on powder samples, as direct catalytic measurements are not feasible for single-crystal model systems.

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Conflicts of Interest

The authors declare no conflicts of interest.

Data Availability Statement

The data that support the findings of this study are available in the Supporting information of this article.

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Supporting Information

Additional supporting information can be found online in the Supporting Information section.

Supporting File: anie72230-sup-0001-SuppMat.pdf.