Initial stages of CO₂ adsorption on CaO: a combined experimental and computational study†

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Room temperature adsorption of carbon dioxide (CO₂) on monocrystalline CaO(001) thin films grown on a Mo(001) substrate was studied by infrared reflection–absorption spectroscopy (IRAS) and quantum chemical calculations. For comparison, CO₂ adsorption was examined on poorly ordered, nanoparticulate CaO films prepared on Ru(0001). For both systems, CO₂ readily adsorbs on the clean CaO surface. However, additional bands were observable on the CaO/Ru(0001) films compared with CaO/Mo(001), because the stricter IRAS surface selection rules do not apply to adsorption on the disordered thin films grown on Ru(0001). Spectral evolution with increasing exposure of the IRA bands suggested the presence of several adsorption sites which are consecutively populated by CO₂. Density functional calculations showed that CO₂ adsorption occurs as monodentate surface carbonate (CO₃¹⁻) species at monatomic step sites and other low-coordinated sites, followed by formation of carbonates on terraces, which dominate at increasing CO₂ exposure. To explain the coverage-dependent IRAS results, we propose CO₂ surface islanding from the onset, most likely in the form of pairs and other chain-like species, which were calculated as thermodynamically favorable. The calculated adsorption energy for isolated CO₂ on the terrace sites (184 ± 10 kJ mol⁻¹) is larger than the adsorption energy obtained by temperature programmed desorption (~120–140 kJ mol⁻¹) and heat of adsorption taken from microcalorimetry measurements at low coverage (~125 kJ mol⁻¹). However, the calculated adsorption energies become less favorable when carbonate chains intersect on CaO terraces, forming kinks. Furthermore, our assignments of the initial stages of CO₂ adsorption are consistent with the observed coverage effect on the CO₂ adsorption energy measured by microcalorimetry and the IRAS results.

1. Introduction

The interaction of carbon dioxide (CO₂) with oxide surfaces is important in a variety of industrial and environmental applications such as methanol synthesis, exhaust cleaning, and CO₂ capturing.1–4 Among alkaline-earth oxides used in such processes, calcium oxide (CaO) often exhibits a higher activity due to a more delocalized electron distribution across surface oxygen atoms.5

Adsorption and reactivity studies on CaO were primarily performed on powders. It was demonstrated that CO₂ adsorption capacities depend upon particle size and thus surface area.5,7 Complete adsorption resulted in formation of calcium carbonate (CaCO₃). However, the reversibility of the carbonate formation falls off with increasing carbonation/calcination cycles, accompanied by rapid decay of CO₂ adsorption.8 Therefore, many studies addressed the stability of CaO particulates, with the aim of improved catalyst regeneration.3,4,9 CaCO₃ formation was found to occur in two kinetic regimes,6 whereby the initial reaction was fast and closely related to the partial pressure of CO₂,6 and a slower second stage that was diffusion-controlled.11

This early work raised great interest in understanding the initial stages of CO₂ adsorption. In a pioneering infrared (IR) spectroscopy study,12 the formation of surface carbonate species (CO₃²⁻) was proposed as monodentate at room temperature, with the appearance of bidentate carbonates at higher temperatures. Substantial effort has been spent to understand the IR spectra employing computational quantum chemistry. Adopting successively larger cluster models, the formation of...
carbonates were also predicted to form on edge 15 and step 18 above 10 showing formation of a surface carbonate species at pressures vacuum-cleaved synthetic single crystal CaO(001) surfaces. Monodentate carbonates were also predicted to form on corner sites. Of the latter, pseudo-tridentate structures can also form, in which all three oxygen atoms of CO$_2$ have elongated bonds to Ca$^{2+}$ surface ions.

Experimental studies of CO$_2$ adsorption on well-defined, single crystal surfaces of CaO are scarce. A synchrotron-based photoemission spectroscopy study of CO$_2$ adsorption on vacuum-cleaved synthetic single crystal CaO(001) surfaces showed formation of a surface carbonate species at pressures above 10$^{-6}$ Torr. However, traces of water were discovered in the ultrahigh vacuum (UHV) system, which partially hydroxylated the CaO(001) surface. Molecular beam scattering and temperature programmed desorption have been employed to study the adsorption kinetics of CO$_2$ on a CaO(001) single crystal. The absence of a clear diffraction pattern indicated a large density of defects; these defects were assigned to oxygen vacancies, which were assumed to cause carbonate decomposition to CO. Few adsorption studies were performed on CaO thin films grown on Si(111) and Si(100) surfaces. In both cases, the resulting films were polycrystalline. The results obtained by various electron spectroscopy techniques suggested the formation of a closed CaCO$_3$ top layer upon CO$_2$ adsorption.

Only recently, the preparation of well-ordered CaO(001) films has been reported using a Mo(001) single crystal as a substrate. Low-energy electron diffraction (LEED), Auger electron spectroscopy (AES), and scanning tunneling microscopy (STM) results showed that well-ordered closed films above 5 nm in nominal thickness with high quality and low defect concentration can be fabricated, with the band gap and optical properties being virtually identical to that reported for bulk CaO. However, depending on the film thickness and annealing temperature, a considerable diffusion of Mo ions from the support into the oxide film may occur that, in turn, alter the surface reactivity. As shown by high-resolution STM in combination with DFT calculations, the presence of Mo in the sub-surface region was responsible for the formation of superoxo O$_2^-$ species and tip-induced oxygen dissociation.

For CO$_2$ on MgO(001), it has been shown that electron transfer from subsurface Mo can affect the preferred binding of CO$_2$ as surface carboxylate rather than carbonate.

In this combined experimental and theoretical study, we report on CO$_2$ adsorption on well-characterized CaO thin films grown on metal substrates, focusing on the initial stages of adsorption in the low coverage regime. Using infrared reflection–absorption spectroscopy (IRAS), temperature programmed desorption (TPD), microcalorimetry, and quantum chemistry calculations, we show that CO$_2$ strongly adsorbs as monodentate carbonates first on step edges, followed by terrace sites.

2. Experimental methods and results

2.1 Experimental methods

The experiments were carried out in three UHV chambers. The first chamber is equipped with LEED (Omicron), STM (Omicron), X-ray photoelectron spectroscopy (XPS) (Scienta 200 analyzer), and an IR spectrometer (Bruker IFS 66v) for the IRAS measurements. The Mo(001) and Ru(0001) crystals (both from MaTeck GmbH) were mounted on Omicron sample holders, with the temperature measured by a thermocouple spot-welded to the edge of the crystal. Clean metal surfaces were prepared by cycles of Ar$^+$ sputtering and UHV annealing at high temperatures. Carbon contaminations were removed by mild annealing in oxygen followed by thermal flash in UHV to 1600 K. The cleanness of metal substrates was inspected by XPS prior to the film preparation. The crystalline CaO(001) films were grown on Mo(001) as described elsewhere. The same preparation was adopted for a Ru(0001) substrate. On both supports, Ca was deposited in $6 \times 10^{-7}$ mbar O$_2$ at 300 K, followed by UHV annealing at $\sim 1200$ K for 5 min. The nominal film thickness (> 5 nm) was determined by attenuation of the substrate signals in XPS.

The IRA-spectra were recorded using p-polarized light at an 84$^\circ$ grazing angle of incidence (spectral resolution 4 cm$^{-1}$). CO$_2$ was dosed via back filling the chamber to 10$^{-8}$ mbar and then pumped before the IRAS measurements. Dosage and measurements were performed at room temperature. CO$_2$ exposures are given in Langmuirs (L), 1 L = 10$^{-6}$ Torr s.

TPD spectra of CO$_2$ on CaO(001)/Mo(001) films, prepared in the same way as described above, were measured in the second UHV chamber equipped with LEED, AES and quadrupole mass spectrometer (QMS). To minimize re-adsorption from the UHV background, CO$_2$ was dosed on CaO(001) using the calibrated directional gas doser.

Finally, the microcalorimetry experiments were performed in the third UHV setup equipped with LEED, AES, QMS and a single-crystal adsorption calorimeter. CO$_2$ was dosed using molecular beam to determine the sticking coefficient and CO$_2$ coverage. Since a crystal substrate for such experiments must have a thickness of only several micrometers, in these studies the CaO(001) films were grown on Pt(001) using the same preparation conditions as used for Mo(001). More detailed characterization of the films on Pt(001) will be presented elsewhere. The resulting CaO(001) films showed good crystallinity as judged by LEED, and showed the same TPD features as those grown on Mo(001), both measured in the second chamber, for direct comparison.

2.2 Experimental results

The typical morphology of well-ordered CaO(001) films grown on Mo(001) is shown in the STM image in Fig. 1a. The films annealed at 1200 K expose relatively wide terraces, although with considerable amounts of screw dislocations. In addition, few point-like defects may be observed on atomically flat terraces as depressions, of which some are outlined by circles in Fig. 1b. For comparison, Fig. 1c displays the STM image of the film prepared under similar conditions on a Ru(0001) substrate, where only
poorly-defined nanoparticles are observed. Such a granular-like morphology may be explained by the epicatxial film growth on metal close-packed (111) surfaces that drives oxide layers to grow in the (111) orientation, which is thermodynamically unfavorable for oxides with the rock salt crystal structure like CaO. As a result, the Ru-supported films did not show any LEED pattern. Based on the XPS results (not shown here), the film stoichiometry on Ru(0001) is close to that observed for CaO(001) on Mo(001), although it might deviate at the surface of highly dispersed nanoparticles due to the abundance of low-coordinated atoms. However, for simplicity, we will use stoichiometric CaO notation for the films grown on Ru(0001).

CO₂ adsorption on the CaO films was monitored by IRAS at 300 K at increasing dosage (Fig. 2). Bearing in mind high sensitivity of CaO towards traces of water in the vacuum background, we note that no surface hydroxyls were detected in all IRA spectra discussed below. At the lowest exposure (~0.1 L) of CO₂ on CaO(001)/Mo(001), a sharp signal appears at 1298 cm⁻¹ together with a broad band centered at ~985 cm⁻¹ (Fig. 2a, top).

At increasing exposure, an additional peak emerges at 1311 cm⁻¹. The latter peak gains intensity while a prominent shoulder appears at 1319 cm⁻¹, which is more clearly observed upon spectral deconvolution (see below). The low frequency band (~990 cm⁻¹) considerably broadens, but qualitatively follows the evolution of the principal bands (or their envelope), slightly blue-shifting with increasing CO₂ exposure.

For comparison, Fig. 2b collects IRA spectra recorded on a nanoparticulate CaO overlayer prepared on Ru(0001). In addition to the 1298 and 986 cm⁻¹ bands, observed on a Mo(001)-supported film at the lowest exposure, a broad feature is observed in the 1630 cm⁻¹ region, which gains considerable intensity with increasing CO₂ exposure. Concomitantly, a sharp band develops at 875 cm⁻¹. The intensities of these two additional bands do not correlate with the 1298 cm⁻¹ band, but rather with the bands that appear at slightly higher frequencies.

Both the 875 and ~1640 cm⁻¹ bands are missing in the spectra of the CaO(001)/Mo(001) films (Fig. 2a). In principle, the observed differences might be due to CO₂ adsorbing in different coordination (e.g., carbonates, carboxylates) on these two films. However, the similar bands (i.e. in the ~1300 and ~980 cm⁻¹ regions) on both systems develop with increasing coverage almost identically. This suggests the adsorbed CO₂ species on these surfaces are, in fact, very similar in nature. In addition, we conclude that the CO₂ interaction with the CaO(001) films is not affected by the presence of Mo in the sub-surface region, if any.

The ~1640, ~1310, ~980, and 875 cm⁻¹ bands on the CaO/Ru(0001) films all gain intensity with increasing exposure,
suggestions they also belong to the same species. The two unobserved bands on the CaO(001)/Mo(001) films, i.e. at \( \sim 1300 \) cm\(^{-1}\) and \( 875 \) cm\(^{-1}\), are likely absent due to the surface selection rules applied to IRAS on metal supported systems, which state that only vibrations with net dipole changes normal to the surface are detected. In contrast, CO\(_2\) ad-species on CaO nanoparticles may be present in random orientations with respect to the metal (Ru) substrate.

Focusing solely on the coverage dependent evolution of the bands in the \( \sim 1300 \) cm\(^{-1}\) region on both systems, it appears that increasing CO\(_2\) coverage causes new peaks to emerge at higher frequencies. This observation contrasts with a single band exhibiting blue shifting, which would be expected in the case of increasing dipole–dipole interaction. The appearance of new high-frequency peaks is more clearly seen in the deconvoluted spectra presented in Fig. 3.

We first address the spectral evolution observed on CaO(001)/Mo(001) (left panel in Fig. 3). At the lowest coverage, the signal at 1298 cm\(^{-1}\) dominates the spectrum. At increasing coverage, this band approximately doubles intensity, but then attenuates by a factor of two. Concomitantly, the band at 1310 cm\(^{-1}\) emerges and then stays fairly constant as if it saturates, while another band at 1322 cm\(^{-1}\) gains intensity. Finally, the new band at 1345 cm\(^{-1}\) appears at the highest coverage studied here. Such a spectral evolution can, in principle, be rationalized in terms of consecutive population of the spatially separated (isolated) adsorption sites.

Interestingly, the respective bands on a Ru-supported film (right panel in Fig. 3) are only slightly broader than on a monocrystalline Mo-supported film, which is consistent with the formation of isolated adsorbates. More surprising is that the intensities and positions of the bands, as well as the coverage dependence, are very similar on these two systems, despite the very different morphology of the films. This finding would indicate that the CO\(_2\) adsorption on CaO is structure insensitive. Alternatively, one may suggest that the film on Ru(0001) is composed of cuboid-shaped particles primarily exposing the (001) surface, and the IRAS intensities are strongly affected by orientation of adsorbed CO\(_2\) with respect to metal surface underneath the film.

In addition, Fig. 4 shows deconvoluted IRA-spectra of CO\(_2\) adsorbed on CaO(001)/Mo(001) at low temperature (\( \sim 110 \) K). This allows for the deconvolution of the \( \sim 985 \) cm\(^{-1}\) band, thereby providing a correlation between the \( \sim 1300 \) cm\(^{-1}\) and \( 985 \) cm\(^{-1}\) bands. At the lowest CO\(_2\) exposure, the \( \sim 1300 \) cm\(^{-1}\) peak is the only prominent signal, regardless of temperature. At increasing coverage, the signal intensifies and does not attenuate. Additionally, new isolated peaks emerge at higher frequencies in a similar fashion to the room temperature results. The emergence of high-frequency signals at 1315 cm\(^{-1}\) and 1330 cm\(^{-1}\) occurs concomitantly with blue-shifted signals at 1005 cm\(^{-1}\) and 1030 cm\(^{-1}\), respectively. These results confirm our suspicion that unique adsorption structures are responsible for the growth of both observed IRA bands.

On the basis of the IRAS results presented above, we may conclude that CO\(_2\) readily adsorbs on the CaO surface in several structurally different adsorption sites. The site corresponding to the 1298 cm\(^{-1}\) band populates first, at both 300 and 110 K, implying that it exhibits the highest sticking coefficient. At increasing exposure, CO\(_2\) will adsorb in the next energetically favorable adsorption sites, giving rise to additional IRA bands at 1310 cm\(^{-1}\) and above.

Comparison of spectra on a well-ordered CaO(001) and nanoparticulate CaO films shows that newly adsorbed species are characterized by the bands at \( \sim 1640, \sim 1310, \sim 980, \) and 875 cm\(^{-1}\). The two out of four bands (i.e. at 1640 and 875 cm\(^{-1}\)) are not observable on Mo(001) supported films, possibly due to the surface selection rules applied to IRAS. According to the
normal mode analysis of Fujita et al.\textsuperscript{31} and the correlation diagram of Davydov\textsuperscript{32} commonly used for the interpretation of IR spectra of CO$_2$ adsorption, these two bands can be assigned to the $\nu$(C–O) and $\pi$ modes of monodentate carbonate species, designated $\nu_6$ and $\nu_3$, respectively, in Fig. 5. These modes have negligible nuclear displacements normal to the surface, consistent with the IRAS surface selection rules. The frequencies of the bending modes ($\nu_1$ and $\nu_2$) are close to the cut-off frequency of our setup and cannot be addressed experimentally. Therefore, only the $\nu_4$ and $\nu_5$ modes will be seen on the CaO(001)-oriented films. Among these two, the observed $\nu_5$ mode exhibits a higher intensity (Fig. 2a), suggesting a larger dipole moment change associated with this mode. Also, the frequencies reported in the literature for these modes are fairly consistent with those measured in our experiments.

To obtain information about the CO$_2$ adsorption energy, we performed TPD and microcalorimetry experiments on well-ordered CaO(001) films in other experimental setups using the same preparation conditions. Fig. 6 shows a typical desorption trace of CO$_2$ (44 amu) recorded after adsorption of 0.5 L at 220 K on the CaO(001)/Mo(001) surface, with the LEED pattern shown in the inset. Note that at the lower exposures, the spectra scale in intensity with small deviations in maximum desorption temperature $T_{\text{max}}$ (not shown). The TPD spectra were reproducible (upon five consecutive TPD runs up to 1000 K), thus suggesting that CO$_2$ desorbs intact. Applying the well-known Redhead formalism\textsuperscript{34} and using the pre-exponential factor $10^{13}$ s$^{-1}$, $T_{\text{max}} = 470$ K, and heating rate 3 K s$^{-1}$, we calculated the desorption energy to be $\sim 120$ kJ mol$^{-1}$. With an increased pre-exponential of $10^{15}$ s$^{-1}$, determined experimentally for physisorbed CO on MgO(001),\textsuperscript{35} this desorption energy increases to $\sim 140$ kJ mol$^{-1}$.

The CO$_2$ adsorption enthalpy on the CaO(001) surface was directly measured by microcalorimetry. The measurement was combined with a sticking coefficient measurement using the King and Wells method,\textsuperscript{36} allowing for a quantitative determination of the absolute number of adsorbed CO$_2$ molecules. The results for adsorption at 300 K are shown in Fig. 7. The heat of adsorption measured at the lowest CO$_2$ coverage, $\sim 125$ kJ mol$^{-1}$, agrees well with the result from TPD. The adsorption enthalpy decreases with increasing coverage and levels off at high exposures.
3. Computational methods and results

3.1 Computational models and methods

Neutral CaₙOₙ clusters were employed to model different morphological sites of CaO(001) for CO₂ adsorption, using the previously used lattice parameter a = 483.2 pm.²⁵ The periodic electrostatic embedded cluster method (PEECM)³⁷ was employed in the DFT calculations using the computational package TURBOMOLE.³⁸

We considered three types of morphological sites: terraces, step-edges, and corners. These sites are represented by twelve clusters, shown in Fig. S1 (ESI†), which are labeled according to the morphological site they represent: T1, T2, T3, T4, and T5 (terrace); MS1 and MS2 (monatomic step); DS (diatomic step); E (edge); RC (reverse corner); and C1 and C2 (corner). The models were constructed such that Ca atoms comprise the outermost shell of the cluster. The presence of the much more polarizable oxygen anions in the outermost shell has to be avoided to minimize nonphysical polarization effects due to point charges in the immediate vicinity.³⁷

Structure optimizations were carried out with the B3LYP³⁹,⁴⁰ and PBE⁴¹,⁴² functionals at a triple-z valence plus polarization (TZVP) basis set ("def2" in the TURBOMOLE library) for Ca, O, and C atoms, with additional Ca atoms containing all-electron effective core potentials (ECPs).⁴⁴ The quadruple-z valence plus polarization (QZVP) basis set was used for basis set extrapolation. We performed MP2 and CCSD(T) calculations with the resolution of identity method, where the auxiliary TZVP and QZVP basis sets were taken from the TURBOMOLE library.⁴⁵ Contributions to the B3LYP and PBE energies resulting from dispersion were calculated according to the procedure outlined by Grimme.⁴⁶ The Cₙ and Rₙ dispersion parameters of Ca²⁺ were assigned to the isoelectric Ar atom, following earlier work that substituted Ne parameters for the Mg²⁺ ions in MgO.⁴⁷

Periodic calculations were performed on terrace and monatomic step models with PBE using the Vienna ab initio simulation package (VASP).⁴⁸,⁴⁹ Additional details can be found in the ESI. The morphological site they represent: T1, T2, T3, T4, and T5 (terrace); MS1 and MS2 (monatomic step); DS (diatomic step); E (edge); RC (reverse corner); and C1 and C2 (corner). The models were constructed such that Ca atoms comprise the outermost shell of the cluster. The presence of the much more polarizable oxygen anions in the outermost shell has to be avoided to minimize nonphysical polarization effects due to point charges in the immediate vicinity.³⁷

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Vibrational frequencies of adsorbed CO₂ were calculated numerically in the harmonic approximation. B3LYP frequencies were scaled by a factor λ = 0.9800, the average of the ratios of observed anharmonic frequencies (fundamentals) and the calculated (TZVP basis set) harmonic frequencies for gas phase CO₂ and CO₂⁻ [see Table S2, ESI†]. PBE frequencies were not scaled because the calculated scale factors for CO₂ and CO₂⁻ at the TZVP level are very close to unity. The global frequency scaling factors for B3LYP and PBE with a comparable 6-311+G(2df,p) basis set are λ = 0.9686 and λ = 0.9948, respectively.⁵⁰ Intensities were calculated with the x- and y-directions (parallel to the surface) of the dipole gradient projected out, consistent with the surface selection rules for IRAS. Additional computational details can be found in Section S1 of the ESI†.

3.2 Computational results

Structures and energies. Upon adsorption on CaO(001), CO₂ preferentially forms a monodentate surface carbonate species at nearly all morphological sites. Optimized structures for CO₂ adsorption at terrace, monatomic step, and corner sites are shown in Fig. 8. On the terrace, the oxygen atoms of CO₂ are positioned along the surface lattice structure, forming a 0° torsional angle with the surface. On step-edges, the carbonate leans away from the (001) face, retaining the 0° torsional angle between the CO₂ oxygen atoms and the rim. For CO₂ adsorption onto a corner site, the carbon atom embeds into the cluster by replacing the corner lattice oxygen atom, forming a tridentate-bound carbonate species. Calculated adsorption energies are listed in Table 1. B3LYP and PBE give the same ranking from

<table>
<thead>
<tr>
<th>Model</th>
<th>Composition</th>
<th>ΔE_ads (kJ mol⁻¹)</th>
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<tbody>
<tr>
<td>T1</td>
<td>Ca₉O₁₄Ca⁺⁰⁺Ca⁻⁰⁻</td>
<td>-151.9</td>
</tr>
<tr>
<td>T2</td>
<td>Ca₉O₁₅Ca⁺⁰⁺Ca⁺⁰⁺</td>
<td>-151.0</td>
</tr>
<tr>
<td>T3</td>
<td>Ca₉O₁₆Ca⁺²⁺Ca⁻⁰⁻</td>
<td>-145.7</td>
</tr>
<tr>
<td>T4</td>
<td>Ca₉O₁₇Ca⁺⁰⁺Ca⁻²⁺</td>
<td>-135.1</td>
</tr>
<tr>
<td>T5</td>
<td>Ca₉O₁₈Ca⁺²⁺Ca⁺²⁺</td>
<td>-134.6</td>
</tr>
<tr>
<td>MS1</td>
<td>Ca₉O₁₃Ca⁺⁰⁺Ca⁺⁰⁺</td>
<td>-287.1</td>
</tr>
<tr>
<td>MS2</td>
<td>Ca₉O₁₄Ca⁺⁰⁺Ca⁺⁰⁺</td>
<td>-276.0</td>
</tr>
<tr>
<td>DS</td>
<td>Ca₉O₁₅Ca⁺⁰⁺Ca⁻⁰⁻</td>
<td>-244.7</td>
</tr>
<tr>
<td>E</td>
<td>Ca₉O₁₆Ca⁺⁰⁺Ca⁻⁰⁻</td>
<td>-249.3</td>
</tr>
</tbody>
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The subscript "pc" indicates the clusters are embedded in a 2D-periodic array of point charges extending six layers deep. Ca atoms in the outermost shell of the clusters are fixed at their lattice positions during structure optimizations, denoted “fix” for atoms described by basis functions and “ecp” for atoms described with ECPs. DFT-D single point calculation at the DFT structure. The surface carbonate is rotated 45° with respect to the surface lattice structure. See Section S2.1 of the ESI for details.
most to least stable for the CO₂ adsorption energy: monatomic step, corner, edge, diatomic step, reverse corner, and terrace.

The adsorption energy calculated with the terrace models converge to about $-155$ and $-151 \text{ kJ mol}^{-1}$ with B3LYP+D and PBE+D, respectively, at very large cluster sizes. Similar convergence was achieved by using successively larger supercells in calculations with periodic boundary conditions, yielding $-146 \text{ kJ mol}^{-1}$ with PBE+D (see Table S12 of the ESI†).

To more accurately determine the adsorption energy of isolated CO₂ on terraces, we performed single-point CCSD(T) calculations on a non-embedded $3 \times 3 \times 2 \text{ Ca}_9\text{O}_9$ cluster. Its structure was taken from the embedded T1 model optimized with B3LYP/TZVP. Subtracting the DFT+D adsorption energies of the Ca₉O₉ cluster from the CCSD(T) results extrapolated to the complete basis set limit gives a coupled cluster correction, $\Delta_{\text{CC}} \text{DFT+D}$, of $-30.3 \text{ kJ mol}^{-1}$ (B3LYP+D) and $-31.9 \text{ kJ mol}^{-1}$ (PBE+D). The final calculated coupled cluster-corrected adsorption energy is $-185.1 \text{ kJ mol}^{-1}$ for B3LYP+D and $-182.5 \text{ kJ mol}^{-1}$ for PBE+D, or $-184 \text{ kJ mol}^{-1}$ on average. Similar hybrid schemes⁵¹ have been used previously to accurately calculate adsorption energies of small molecules on oxide surfaces.⁵⁷

To estimate the uncertainty of the coupled cluster corrections, we also calculated the adsorption energy of CO₂ on the entire B3LYP-optimized T1 cluster at the TZVP level within the PEECM formalism, giving a $\Delta_{\text{CC}} \text{B3LYP+D/TZVP}$ of $-22.8 \text{ kJ mol}^{-1}$. This differs by $+10 \text{ kJ mol}^{-1}$ from the corresponding value for the Ca₉O₉ cluster ($-32.9 \text{ kJ mol}^{-1}$, see Table S18 of the ESI†). The differences between CCSD(T) and B3LYP+D for the analogous gas-phase reaction CaO(g) + CO₂(g) → CaCO₃(g), $-40.5 \text{ kJ mol}^{-1}$ (see Table S19 of the ESI†), deviates by about $+10 \text{ kJ mol}^{-1}$. We therefore conclude that the uncertainty of our CCSD(T) correction, and hence of the final adsorption energy due to the applied methods, is $\pm 10 \text{ kJ mol}^{-1}$, whereas convergence has been achieved for type (periodic vs. embedded cluster) and size of the models. Therefore, our best estimate for the energy of adsorption of CO₂ on terrace sites is $184 \pm 10 \text{ kJ mol}^{-1}$.

The effect of increasing the CO₂ coverage was explored by optimizing the structure of two or more CO₂ molecules on several CaO(001) morphological sites. On terrace sites, adsorption of CO₂ in close proximity to a monodentate carbonate occurs at the nearest oxygen atom, causing both adsorbates to form a 45° torsional angle with respect to the surface lattice structure (see “Terrace 2 CO₂” in Fig. 8 and the top of Fig. 9). This structure is similar to a previously calculated carbonate pair on CaO(001) with an interatomic potential model.⁵² The present calculations indicate that formation of such a pair is thermodynamically favorable. Subsequent CO₂ molecules will add to the length of a carbonate chain stretching diagonally across the terrace surface, as shown in Fig. 9. Details are given in Section S2 of the ESI†.

Structural changes also occur when CO₂ binds to step and edge sites in close proximity to other populated 4-coordinate sites. As shown in Fig. 8 (“Monatomic Step 2 CO₂”) and the bottom of Fig. 9, adsorption of CO₂ at the nearest neighbor oxygen atom on the rim of the step results in an asymmetric structure due to steric interactions of the terminal oxygen atoms. While the pre-adsorbed CO₂ binds nearly in the plane of the missing surface layer, the adjacent CO₂ more closely resembles a carbonate bound to a 5-coordinated site. On adsorption of the second CO₂ molecule in the nearest neighbor position, 60 kJ mol⁻¹ less energy is released compared to adsorption of isolated CO₂ on a monatomic step site. Adsorption in the next nearest neighbor position, however, only incurred a 9 kJ mol⁻¹ energy penalty, and results in a structure similar to that of an isolated CO₂. CO₂ binding in between two pre-adsorbed carbonates is $\sim 100 \text{ kJ mol}^{-1}$ less favorable than in isolation. In spite of these energy penalties, adsorption of CO₂ on steps is more favorable than on a clean terrace, even at the most crowded step adsorption site, by $\sim 10 \text{ kJ mol}^{-1}$.

We calculated structures and energies for CO₂ adsorption at 4- and 5-coordinated sites in proximity to CO₂ pre-adsorbed to low-coordinated sites to study the interactions of adsorbates bound to different morphological sites. Adsorption on the rim

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Fig. 9 Structures of adsorbed CO₂ on terrace sites (top view) and monatomic step sites (side view of the step-edge) from low (left) to high (right) coverage optimized with PBE using periodic models. The step models have been truncated for clarity. Adsorption of CO₂ results in an infinite diagonal stripe across the CaO(001) terrace. The monatomic step sites become completely filled by monodentate carbonate species in a zigzag pattern, whereby every other adsorbate leans toward the step surface layer. Color code: red, O; gray, Ca; cyan, C.
(4-coordinated) nearest a pre-adsorbed corner site (3-coordinated) is predicted to be the most stable, with a similar adsorption energy to a clean edge site. Adsorption at a terrace site (5-coordinated) in proximity to filled 4-coordinated (steps and edges) and 3-coordinated (corner) sites were found to be the least favorable among the tested sites. Details on the adsorbate–adsorbate interactions can be found in Section S2 of the ESI. In addition, a physisorbed paired structure was considered, owing to the stability of the (CO₂)₂ dimer in the gas phase. However, the physisorbed pair, although energetically stable, was calculated to be the least favorable, ~140 kJ mol⁻¹ less favorable than chemisorption on the clean terrace.

After the low-coordinated sites have become filled, adsorption at terraces commences. While the calculations suggest that infinite chains of surface carbonates will form diagonally across the CaO(001) terrace, we sought to calculate different terrace-adsorption structures to simulate kinks in such adsorption chains that are likely to occur upon room temperature adsorption. Such kinks also represent the interaction of two chains intersecting on the terrace surface. Due to the available terrace adsorption sites on T1, we only considered structural patterns up to a total of five carbonates. Optimized local minima for adsorption of three, four, and five CO₂ molecules on cluster T1, which simulates 0, 1, and 2 kinks in the CO₂ adsorption chain, are shown in Fig. 10. The relative energies of the local minimum structures are labeled with letters, “A” being the calculated global minimum, and subsequent letters denoting structures with higher energy. The presence of a single kink in the CO₂ adsorption chain causes a decrease in adsorption strength of ~13 kJ per mol per adsorbed CO₂ as calculated with the T1 cluster; an additional ~10 kJ per mol penalty per adsorbate is calculated for the presence of two kinks in the carbonate chain. However, due to the interaction of the adsorbates on the periphery of the cluster with the surrounding point charges, these calculated energy differences are likely exaggerated. See Section S2 of the ESI for an analysis of computational artifacts due to CO₂ cluster adsorption sites.

Vibrational frequencies. As shown in Fig. 5, monodentate surface carbonates exhibit six IR-active bending and stretching modes: ν₁, asymmetric bending δ(O–CO); ν₂, symmetric bending δ(O–CO); ν₃, out of plane deformation π; ν₄, symmetric stretch ν(C–O); ν₅, symmetric stretch ν(C–O) + ν(C–O); and ν₆, asymmetric stretch ν(C–O), where Oₘ denotes the surface oxygen. The B3LYP simulated IRA spectra for the adsorption of a single CO₂ molecule as a surface carbonate on CaO(001) cluster models are depicted in Fig. 11. Note that the calculated intensities are corrected for surface dipole selection rules. With the exception of the corner site, where the CO₂ embeds into the corner, the carbonate formation yields two significant surface IR-active modes, both symmetric stretchings: a ν₁ mode between 1271 and 1293 cm⁻¹, and a ν₄ mode at ~922 cm⁻¹ for the terrace site and between 987 and 1014 cm⁻¹ for the other sites. These calculated vibrational frequencies qualitatively explain the peaks in the experimental IRA spectra at low CO₂ coverage. For the embedded carbonate at the corner site, the peak calculated at ~1420 cm⁻¹ is better described as a degenerate asymmetric mode similar to ν₆. The ν₅ bands (~842–875 cm⁻¹ with B3LYP and ~805–849 cm⁻¹ with PBE) and ν₆ bands (~1584–1615 cm⁻¹ with B3LYP and ~1528–1617 cm⁻¹ with PBE) are lacking intensity due to the applied surface selection rules (see Table S26 of the ESI). These bands, however, should not be ignored when discussing the experimental spectra of CO₂ adsorption on CaO/Ru(0001) in Section 4.

![Fig. 10](image-url) Structures of three, four, and five adsorbed CO₂ molecules on cluster T1 optimized with B3LYP from top views, simulating 0, 1, and 2 kinks in the chain of adsorbed carbonates, respectively. Color code: red, O; gray, Ca; cyan, C.

![Fig. 11](image-url) Calculated (scaled) IRA spectra of CO₂ adsorption on cluster models T1, MS2, DS, E, RC, and C1 with B3LYP. The experimental peaks are indicated by dashed lines for the lowest CO₂ exposure. Frequencies and intensities tabulated in Table S26 (ESI). All calculated spectra are Lorentzian broadened.
IRA spectra simulated with B3LYP for the adsorption of two CO₂ molecules as surface carbonates on CaO(001) cluster models are shown in Fig. 12. On the terrace site, adsorption of CO₂ at the nearest neighbor site to a pre-adsorbed species causes a 17 cm⁻¹ blue shift in the ν₂ peak and a 41 cm⁻¹ blue shift in the ν₅ peak. On the monatomic step site, CO₂ adsorption at the nearest neighbor site to a pre-adsorbed species causes a red shift of 30 cm⁻¹ in the ν₄ peak with approximately six times increased IRA intensities for ν₅ and ν₄. The orientation of the adsorbed CO₂, which does not lean toward the step surface layer as much as the pre-adsorbed species, results in dipole changes perpendicular to the surface and observation by IRAS more likely. The calculated frequencies of two CO₂ molecules to the rim of the edge cluster model are similar to those on the monatomic step. On a corner site, the calculated IRA spectrum of adsorption of two CO₂ molecules shows three peaks in the vicinity of ν₂: one at ~1300 cm⁻¹ that is attributed to the CO₂ binding on the nearby edge, and two additional peaks at ~1390 and ~1440 cm⁻¹ that are attributed to the pre-adsorbed CO₂ embedded into the corner. The appearance of a second ν₅ peak due to the embedded CO₂ is due to the broken symmetry caused by the additional nearby CO₂ adsorbate. As with the isolated CO₂ adsorption, calculated ν₃ bands (~836–879 cm⁻¹ with B3LYP and ~795–854 cm⁻¹ with PBE) and ν₆ bands (~1577–1664 cm⁻¹ with B3LYP and ~1541–1655 cm⁻¹ with PBE) can be compared to the experimental CaO/Ru(0001) spectra (see Section 4 for discussion and Tables S27 and S28 of the ESI† for data).

Calculated IRA spectra for the adsorption of three, four, and five CO₂ molecules as surface carbonates on CaO(001) terraces are shown in Fig. 13. Adsorption of CO₂ in the nearest neighbor position to a pre-adsorbed carbonate pair begins the formation of a carbonate chain across the terrace. The ν₅ frequency is blue-shifted 11 cm⁻¹ compared to the paired structure, while the ν₄ frequency splits into two peaks that correspond to atomic motions of the outer carbonates in the chain (blue-shifted peak) and inner carbonates of the chain (red-shifted peak). Details regarding the changes in the ν₄ and ν₅ frequencies upon growth of the carbonate chain on terraces are given in Section S2 of the ESL†. A blue shift of the calculated IRA frequencies of the carbonate chain is expected due to the increased dipole–dipole interaction of the adsorbates. However, this effect remains constant in chains ranging from three to infinite carbonates in length. Because of the limited size of the T1 model, addition of a fourth CO₂ molecule leads to a kink in the adsorption chain of carbonates, causing the simulated IRA bands to split. An increased blue-shift in the ν₅ frequency is also observed. The effect of adding a fifth CO₂ molecule to the surface, which simulates two kinks in the carbonate chain, results in additional blue-shifting of the ν₅ frequency.

4. Discussion

Adsorption of CO₂ on CaO(001) was found to be most favorable on low-coordinated sites such as steps, edges, and corners (−257 to −306 kJ mol⁻¹). Despite an energy penalty due to crowding, the calculations showed that adsorption on an isolated terrace site (−184 ± 10 kJ mol⁻¹) was ~10 kJ mol⁻¹ less favorable than to completely fill available step-edge sites. The adsorption energies obtained by TPD (~120–140 kJ mol⁻¹) and heats of adsorption from microcalorimetry measurements (~125 kJ mol⁻¹) at low coverage are inconsistent with the very strong adsorption energies calculated for CO₂ at any isolated morphological site. However, when carbonate chains on the
terrace surface intersect, forming kinks, the average adsorption strength decreases by ~10–20 kJ per mol per kink. These data suggest that the TPD and microcalorimetry results correspond to aggregates of intersecting CO₂ chains, even at low surface coverage. Presuming the CO₂ sticking coefficient is not too large, we expect that CO₂ adsorption at the lower-coordinated CaO(001) sites should manifest themselves in the experimental IRA spectra obtained at the lowest exposures, followed by adsorption on terraces at higher exposures.

Comparison of the calculated IRA spectra (surface selection rules) of single CO₂ molecule adsorption (Fig. 11) with the experimental spectra at the lowest CO₂ coverage (Fig. 2, top curves) shows that terrace and corner sites do not fit the experimental data. Also for adsorption at the reverse corner site, a peak at 1585 cm⁻¹ is predicted, which is absent in the experimental spectra for the Mo substrate. The calculated spectra at monatomic step, diatomic step, and edge sites more closely agree for the 1298 cm⁻¹ (ν₁) band, while the diatomic step and edge sites agree best with the 985 cm⁻¹ (ν₄) band. Considering the calculated energies for adsorption of carbonate pairs on low coordinated sites (−192 to −245 kJ mol⁻¹), it is likely that pairs and higher oligomers have already formed at the lowest CO₂ exposure. Even in UHV conditions, CO₂ traces in the background readily bind to low-coordinated CaO(001) sites. Therefore, we must consider the calculated IRA spectra of carbonate pairs to assign the experimental low-coverage CO₂ spectra.

Comparison of the calculated IRA spectra of adsorption of two CO₂ molecules (Fig. 12) with the low-coverage experimental spectra (Fig. 2) indicates that adsorption at terrace, monatomic step, and edge sites agree well with the experimental bands. Because the CO₂ adsorption energies are much stronger on step and edge sites than terrace sites, the low-exposure IRA spectra are likely due to adsorption of two or more CO₂ molecules at step and edge sites. From the calculations on periodic step models, we can compare the ν₁ frequency of the completely filled monatomic step (1260 cm⁻¹ with PBE) with that of a pair of carbonates adsorbed at nearest neighbor positions (1266 cm⁻¹). The latter structure corresponds to the calculated carbonate pair on the cluster model (simulated IRA spectrum in red in Fig. 12). From these calculations, we estimate a 6 cm⁻¹ deviation of the calculated ν₁ frequency when using the step-adsorbed CO₂ pair as a model for completely filled step sites. The calculated ν₁ frequency of the nearest-neighbor step-adsorbed pair on the MS2 cluster with PBE is 1266 cm⁻¹, in perfect agreement with the periodic model. Applying the 6 cm⁻¹ correction to the calculated ν₁ frequencies on cluster MS2 (1266 cm⁻¹ for PBE and 1290 cm⁻¹ for B3LYP) gives estimated ν₁ frequencies of completely filled step sites of 1260 cm⁻¹ (PBE) and 1284 cm⁻¹ (B3LYP). These frequencies are assigned to the first observable peak at 1298 cm⁻¹ in the experimental spectrum, given in Table 2.

The experimental IRA spectra at higher CO₂ exposures (Fig. 2, middle curves) show the appearance of new blue-shifted ν₁ peaks. Spectral deconvolution (Fig. 3 and 4) indicates that new peaks sequentially appear without blue-shifting, suggesting that each new peak is attributable to CO₂ adsorption at a different site. The experimentally determined shifts in the ν₁ frequencies are 12 cm⁻¹, 24 cm⁻¹, and 47 cm⁻¹ from the original 1298 cm⁻¹ peak, summarized in Table 2. According to the calculations, these new adsorption peaks can be assigned to various configurations of adsorption on terraces, corresponding to long chains of carbonates and kinks. The calculated frequencies of a single CO₂ adsorbed on a terrace do not qualitatively match the experimental spectrum. However, it is thermodynamically favorable to form carbonate chains, rendering isolated carbonates on the terrace unlikely to be observed via IRAS. The calculated ν₁ frequencies of two or three adsorbed CO₂ molecules on a terrace are 1291 cm⁻¹ and 1302 cm⁻¹ with B3LYP, respectively, which is 7 cm⁻¹ or 18 cm⁻¹ blue-shifted compared to the estimated ν₁ frequency of completely filled monatomic steps, 1284 cm⁻¹. With PBE, the calculated shifts in the ν₁ frequencies from the completely filled steps to a carbonate pair on a terrace are 3 cm⁻¹ with the cluster model and 12 cm⁻¹ with the periodic model; the analogous shifts for carbonate chains are 15 cm⁻¹ (cluster model) and 19–23 cm⁻¹ (periodic model). These values are summarized in Table 2.

The kinks in the carbonate chain simulated by adsorption of four and five CO₂ molecules onto cluster T₁ cause a split in the ν₁ frequencies. The high-frequency ν₁ bands range from 1311–1336 cm⁻¹ with B3LYP and from 1283–1295 cm⁻¹ with PBE, which correspond to blue-shifts of 27–52 cm⁻¹ with B3LYP and 23–35 cm⁻¹ with PBE from the completely filled step ν₁ frequency estimate. The low-frequency ν₁ bands range from 1283–1294 cm⁻¹ with B3LYP and from 1261–1272 cm⁻¹ with

Table 2: Calculated ν₁ frequencies and shifts [cm⁻¹] of (CO₂)₅-CaO for cluster and periodic models

<table>
<thead>
<tr>
<th>CO₂ adsorption</th>
<th>Cluster models</th>
<th>Periodic model</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>B3LYP</td>
<td>PBE</td>
</tr>
<tr>
<td>Filled steps</td>
<td>1284</td>
<td>1260</td>
</tr>
<tr>
<td>Terrace pair</td>
<td>1291 (+7)</td>
<td>1263 (+3)</td>
</tr>
<tr>
<td>Terrace chain</td>
<td>1302 (+18)</td>
<td>1275 (+15)</td>
</tr>
</tbody>
</table>

a The frequency shifts are given in parentheses, defined by subtracting the ν₁ frequency of the completely filled step sites. b Calculated for cluster models MS2 (filled steps), T₂ (terrace pair), and T₁ (terrace chain and kinks). The rationale for utilization of different clusters is discussed in Section S2 of the ESL. c The frequencies are calculated for the largest periodic models. The range of frequencies for the chain are calculated from adsorption of three or more CO₂ molecules in a diagonal stripe. d B3LYP frequencies are scaled by λ = 0.9800. e Experimental frequencies are taken from the room temperature spectral deconvolution in Fig. 3. f The ν₁ frequency ranges given are only for the calculated high-frequency bands.
PBE. These low-frequency $\nu_5$ bands overlap with the calculated $\nu_5$ frequencies of filled step sites and carbonate pairs and chains on terraces. The high-frequency $\nu_5$ bands, however, exhibit additional blue-shifting indicative of the experimental results at higher CO$_2$ exposures.

Our assignments of the initial stages of CO$_2$ adsorption on CaO(001) are outlined in Table 2. The initial peak observed at 1298 cm$^{-1}$ is likely due to completely filled step and edge sites, which are predicted to be completely filled prior to adsorption on terraces. Corner and reverse corner sites are also expected to be populated at the lowest exposure, but at a small enough density as to remain undetected by IRAS. The next observable peak at $\sim$1310 cm$^{-1}$ is assigned to carbonate pairs and short chains on terraces. The subsequent peaks at $\sim$1322 cm$^{-1}$ and $\sim$1345 cm$^{-1}$ are assigned to long carbonate chains on the terrace that intersect, forming various 90° kinks. We note that our assignments are consistent with the positions and shifts of the $\nu_5$ bands. Experimentally, the $\nu_4$ band is especially broad and asymmetric at higher CO$_2$ exposures. The length and kinks of the carbonate chains cause the $\nu_5$ bands to split into numerous peaks that are blue- and red-shifted compared to the original 985 cm$^{-1}$ peak.

We can further assign peaks that are only visible upon adsorption of CO$_2$ onto CaO/Ru(0001). The asymmetric stretches ($\nu_6$) of monodentate carbonates on terraces correspond to the broad peak at $\sim$1635 cm$^{-1}$ in Fig. 2b. The calculated $\nu_6$ frequencies for adsorption of two CO$_2$ molecules on a monatomic step are 1577 cm$^{-1}$ with B3LYP and 1564 cm$^{-1}$ with PBE. The analogous calculated $\nu_6$ frequencies of a carbonate pair on a terrace are 1629 cm$^{-1}$ with B3LYP and 1618 cm$^{-1}$ with PBE. Finally, the calculated $\nu_6$ frequencies of a carbonate chain are 1683 cm$^{-1}$ with B3LYP and 1674 cm$^{-1}$ with PBE. Accordingly, the peak at 875 cm$^{-1}$ in Fig. 2b can be attributed to the $\pi$ mode ($\nu_7$) of CO$_2$ pairs and chains on terraces. The calculated $\nu_7$ frequencies on a terrace are $855$ cm$^{-1}$ for a pair and 858 cm$^{-1}$ for a chain with B3LYP, and 828 cm$^{-1}$ for a pair and 832 cm$^{-1}$ for a chain with PBE. Given our assignments that the initial IRA peaks are due to populated step and edge sites, while subsequent peaks are attributed to various configurations of surface carbonates on terraces, the qualitative agreement with the $\nu_6$ and $\nu_5$ modes on these morphological sites provides additional validation.

5. Conclusions

Well-ordered CaO(001) films grown on a Mo(001) substrate were used as a model system to study initial stages of CO$_2$ adsorption on CaO by IRAS. Comparison of the low-coverage IRA spectra on CaO/Mo(001) with CaO/Ru(0001), which is characterized by a mixed nanoparticulate morphology by STM, indicates similar peak position and intensities of two bands at $\sim$1300 and $\sim$985 cm$^{-1}$. Other peaks present in the IRA spectra on the Ru(0001) substrate are missing in the spectra of CaO(001)/Mo(001) due to surface selection rules, in which only dipole changes normal to the surface are observed.

The calculated adsorption energies for isolated CO$_2$ on terrace sites (184 ± 10 kJ mol$^{-1}$) are larger than adsorption energies obtained by TPD ($\sim$120–140 kJ mol$^{-1}$) and heats of adsorption from microcalorimetry measurements ($\sim$125 kJ mol$^{-1}$). However, the calculations indicate that formation of pairs and chains are thermodynamically favorable and therefore present from the onset of the adsorption experiments. A decrease in adsorption energy is observed when such terrace chains of adsorbed CO$_2$ intersect, forming kinks. This is consistent with experimental evidence from microcalorimetry that suggests CO$_2$ aggregates on the CaO surface.

The present DFT calculations indicate that the initial IRA peaks correspond to the symmetric stretching of monodentate surface carbonates bound to low-coordinated sites, primarily monotonic steps. Such sites are the most thermodynamically favorably adsorption sites ($\sim$295 kJ mol$^{-1}$) and will be completely filled prior to adsorption at terraces. At increasing CO$_2$ dosing, additional IRA peaks are observed in the range $\sim$1310–1345 cm$^{-1}$. These peaks can be assigned to symmetric stretching modes of adsorbed carbonate pairs and chains on terraces. At higher exposure, the carbonate chains on terraces intersect, forming kinks. The additional blue-shifting in the calculated frequencies of these kinks agrees well with the observed shifts in the experimental IRA spectra.

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