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## Reduction of CO<sub>2</sub> by a masked two-coordinate cobalt(i) complex and characterization of a proposed oxodicobalt(ii) intermediate†‡

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Fixation and chemical reduction of CO<sub>2</sub> are important for utilization of this abundant resource, and understanding the detailed mechanism of C–O cleavage is needed for rational development of CO<sub>2</sub> reduction methods. Here, we describe a detailed analysis of the mechanism of the reaction of a masked two-coordinate cobalt(i) complex, L<sup>tBu</sup>Co (where L<sup>tBu</sup> = 2,2,6,6-tetramethyl-3,5-bis[(2,6-diisopropylphenyl)imino]hept-4-yl), with CO<sub>2</sub>, which yields two products of C–O cleavage, the cobalt(i) monocarbonyl complex L<sup>tBu</sup>Co(CO) and the dicobalt(ii) carbonate complex (L<sup>tBu</sup>Co)<sub>2</sub>(μ-CO<sub>3</sub>). Kinetic studies and computations show that the κN,η<sup>6</sup>-arene isomer of L<sup>tBu</sup>Co rearranges to the κ<sub>2</sub>N,N' binding mode prior to binding of CO<sub>2</sub>, which contrasts with the mechanism of binding of other substrates to L<sup>tBu</sup>Co. Density functional theory (DFT) studies show that the only low-energy pathways for cleavage of CO<sub>2</sub> proceed through bimetallic mechanisms, and DFT and highly correlated domain-based local pair natural orbital coupled cluster (DLPNO-CCSD(T)) calculations reveal the cooperative effects of the two metal centers during facile C–O bond rupture. A plausible intermediate in the reaction of CO<sub>2</sub> with L<sup>tBu</sup>Co is the oxodicobalt(ii) complex L<sup>tBu</sup>CoOCOL<sup>tBu</sup>, which has been independently synthesized through the reaction of L<sup>tBu</sup>Co with N<sub>2</sub>O. The rapid reaction of L<sup>tBu</sup>CoOCOL<sup>tBu</sup> with CO<sub>2</sub> to form the carbonate product indicates that the oxo species is kinetically competent to be an intermediate during CO<sub>2</sub> cleavage by L<sup>tBu</sup>Co. L<sup>tBu</sup>CoOCOL<sup>tBu</sup> is a novel example of a thoroughly characterized molecular cobalt–oxo complex where the cobalt ions are clearly in the +2 oxidation state. Its nucleophilic reactivity is a consequence of high charge localization on the μ-oxo ligand between two antiferromagnetically coupled high-spin cobalt(ii) centers, as characterized by DFT and multireference complete active space self-consistent field (CASSCF) calculations.

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## Introduction

Terminal oxo complexes of transition metals are often invoked as intermediates in hydrocarbon activation,<sup>1–3</sup> oxygen atom transfer<sup>4,5</sup> and water oxidation.<sup>6,7</sup> Dinuclear oxo-bridged systems have garnered less attention, even though they have numerous important roles in reactive metallocofactors,<sup>8,9</sup> materials,<sup>10</sup> catalysts,<sup>11</sup> and physiological processes.<sup>12</sup> Like other first row transition metals, cobalt has received growing interest for catalysis due to its versatility and low cost.<sup>13</sup> In particular, cobalt species containing doubly bridged oxo- or hydroxo-bridged cobalt subunits<sup>14</sup> are in focus in the context of water oxidation.<sup>15,16</sup> However, dinuclear Co–O–Co compounds with a single oxo bridge, the simplest bridging cobalt oxo species, are rare and only two examples have been reported. Stauber *et al.* presented a formally dianionic dicobalt(III) oxo species, for which XAS and computational results indicated an unusual electronic structure comprising two cobalt(II) ions, a bridging oxyl radical and an additional “hole” on the supporting ligand.<sup>17</sup> The earlier



example of a dicobalt oxo species by Zhang *et al.* does not include sufficient spectroscopic details to establish the oxidation states of the metals.<sup>18</sup> Thus, understanding the chemical behavior of dicobalt(II) oxo complexes remains elusive.

Here, we describe a new oxodicobalt(II) complex that arises in the context of CO<sub>2</sub> reduction. Carbon dioxide is a persistent environmental pollutant and a C1 feedstock for chemical industries, which has inspired a large amount of research on transition metal catalysts for CO<sub>2</sub> reduction.<sup>19–21</sup> The practical motivations for CO<sub>2</sub> transformation are accompanied by fundamental interest in the detailed mechanisms and charge localization in reduced CO<sub>2</sub> intermediates.<sup>19,22</sup> Here, we focus on cobalt complexes, which are under active study because they are adept at catalytic reductions of CO<sub>2</sub> to CO.<sup>13,23–31</sup>

In the work described here, we use the reduced, unsaturated cobalt site in L<sup>tBu</sup>Co (**1**), a “masked” two-coordinate complex supported by the β-diketiminato ligand 2,2,6,6-tetramethyl-3,5-bis[(2,6-diisopropylphenyl)imino]hept-4-yl (L<sup>tBu</sup>).<sup>32</sup> Its reaction with CO<sub>2</sub> cleaves a C–O bond in CO<sub>2</sub>, and we use a combination of experimental kinetics, density functional theory (DFT), and highly correlated domain-based local pair natural orbital coupled cluster theory with single, double, and perturbative triple excitations (DLPNO-CCSD(T)) to show cooperation of two Co ions for facilitating a bimetallic reaction pathway for CO<sub>2</sub> reduction. We demonstrate that a likely intermediate is the Co<sup>2+</sup>–O<sup>2–</sup>–Co<sup>2+</sup> complex L<sup>tBu</sup>CoOCoL<sup>tBu</sup>, which can be independently synthesized using N<sub>2</sub>O. We investigate the electronic structure and reactivity of this new oxodicobalt(II) complex in detail, including multi-reference complete active space self-consistent field (CASSCF) calculations.

## Results and discussion

### Activation of CO<sub>2</sub> by L<sup>tBu</sup>Co

The cobalt(I) source L<sup>tBu</sup>Co (**1**) was synthesized and characterized earlier,<sup>32</sup> and features a supporting β-diketiminato ligand that is bound in an unusual κ<sup>1</sup>,η<sup>6</sup> binding mode as shown at the left of Scheme 1. The reactivity of **1** includes the binding of Lewis bases like THF, CO, N<sub>2</sub>, pyridine and PPh<sub>3</sub>, and the cleavage of C–F bonds in fluoroarenes.<sup>32,33</sup> We also reported the activation of O<sub>2</sub> at **1**.<sup>34</sup> Each of these reactions causes ligand rearrangement to the more typical κ<sup>2</sup>-binding mode of the β-diketiminato ligand in the three-coordinate or four-coordinate products. This led to a description of **1** as a “masked” low-valent two-coordinate Co complex. However, in these studies, a two-coordinate isomer of **1** with κ<sup>2</sup>-binding of diketiminato was never formed; kinetic studies of the reaction of **1** with pyridine showed a first-order dependence on pyridine coordination which, when combined



Scheme 1 Reaction of L<sup>tBu</sup>Co (**1**) with CO<sub>2</sub> to form **2** and **4**.

with supporting DFT investigations, indicated that the incoming ligand coordinates *prior* to “slipping” of the arene.<sup>32</sup> Computational studies on the reaction with CO indicated a similar associative pathway, in which ligand binding precedes diketiminato rearrangement.<sup>35</sup> The high reactivity of **1** toward small molecules, in conjunction with our interest in understanding the role of low-valent cobalt complexes in CO<sub>2</sub> reduction, prompted us to investigate its reaction with CO<sub>2</sub>. We were limited to aliphatic hydrocarbon solvents, due to the reaction of **1** with arenes and ethers.<sup>32</sup>

The addition of 10 molar equivalents of CO<sub>2</sub> gas to L<sup>tBu</sup>Co in cyclohexane-*d*<sub>12</sub> causes a slow color change from brown to red-orange over several hours at 10 °C. <sup>1</sup>H NMR spectroscopy shows the appearance of new paramagnetically shifted peaks (Fig. S3†). Comparison of the resultant spectra to those of independently synthesized compounds shows that the products are a 1 : 1 mixture of previously reported L<sup>tBu</sup>Co(CO) (**2**)<sup>35</sup> and the new carbonate-bridged compound L<sup>tBu</sup>Co(μ-OCO<sub>2</sub>)CoL<sup>tBu</sup> (**4**) (Scheme 1). Comparison to an internal integration standard indicates that **2** and **4** are each formed in >86% spectroscopic yield. These two products can be separated and isolated as pure solids in 66% and 40% yields, respectively, with the lower isolated yields attributed to losses during crystallization.

### Characterization of the dicobalt(II) carbonate product

X-ray quality crystals of **4** were grown from toluene, and analysis of the diffraction data revealed two independent sites in the asymmetric unit with different carbonate binding modes to the Co centers: (site a) μ-η<sup>1</sup>:η<sup>2</sup> and (site b) μ-η<sup>2</sup>:η<sup>2</sup> (Fig. 1). Site a also had disorders in the core, with two conformations of μ-η<sup>1</sup>:η<sup>2</sup> carbonate in a 77 : 23 ratio that differ by which Co atom is η<sup>1</sup> and which is η<sup>2</sup>. The major component in site a is discussed here for simplicity. Two binding modes of a bridging carbonate were also observed in the crystal structure of an analogous Fe

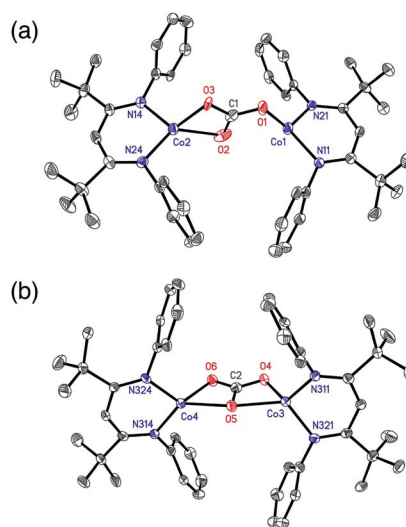


Fig. 1 The two crystallographically independent molecules in the crystal structure of **4**: (a) μ-η<sup>1</sup>:η<sup>2</sup> form, with the major disorder component shown; (b) μ-η<sup>2</sup>:η<sup>2</sup>. H atoms and <sup>i</sup>Pr groups are omitted for clarity. Thermal ellipsoids are shown at 50% probability.





coordinated cobalt(II) complex  $L^{tBu}Co(\mu-OH)_2CoL^{tBu}$ , and four-coordinate oxygen-coordinated cobalt(III) complex  $L^{tBu}Co(\mu-O)_2CoL^{tBu}$ .<sup>34</sup> The pre-edge and edge features overlapped in all of the compounds, including the previously reported cobalt(II) and cobalt(III) analogues (Fig. S10†), indicating that XAS does not unambiguously distinguish the oxidation level. This ambiguity is unfortunate, but fairly common.<sup>41</sup> Comparison of cobalt(II) and cobalt(III) species using X-ray photoelectron spectroscopy gave similarly ambiguous results.

The presence of an oxo in **3** is particularly notable, given the paucity of cobalt(II) oxo complexes.<sup>17,18</sup>  $Co^{III}_2(\mu-O)_2$  complexes have also been described.<sup>34,38</sup> Another relevant comparison is to the diiron(II) complex  $[L^{tBu}Fe]_2O$ ,<sup>42</sup> which has the same supporting ligand and connectivity as **3**. The M–O bond length is shorter for cobalt (Co–O = 1.704(4) Å) than iron (Fe–O = 1.7503(4) Å), as expected from the smaller ionic radius of cobalt(II) versus iron(II). The method of preparing **3** is also interesting, because  $N_2O$  is often kinetically inert, particularly in reactions with late transition metal complexes.<sup>43–45</sup> This serves as another demonstration of the high reactivity of the masked two-coordinate complex  $L^{tBu}Co$  toward cleaving strong bonds.<sup>32,34</sup>

### Electronic structure of the Co–O–Co core and its connection to reactivity

Previous studies have highlighted the correlation between reactivity and the electronic structure.<sup>46–49</sup> We first carried out broken symmetry density functional theory (DFT) calculations on **3** using the BP86 functional. Geometry optimization with BP86 and a mixed basis set combination, B1 (triple- $\zeta$  quality TZVP<sup>50</sup> basis set on Co, O, N, and selected C atoms, and a double  $\zeta$  quality split-valence basis set, def2-SVP,<sup>51</sup> on the rest of the atoms, in a polarizable continuum solvent model, CPCM,<sup>52</sup> using  $\epsilon = 2.3$  for benzene, and D3BJ empirical dispersion), gave a core geometry that is in good agreement with the X-ray crystal structure, including the slight Co–O–Co bending, as shown in Table 2. Using this geometry, single-point calculations with pure BP86, *meta*-GGA M06L, and hybrid B3LYP density functionals all predicted a local high-spin  $d^7$  configuration for each cobalt center. The two cobalt(II) centers are antiferromagnetically coupled, which is achieved through three pathways as indicated by three spin-coupled pairs with overlaps in the range of 0.2 to 0.5. The metal–ligand interactions are largely ionic, because doubly occupied metal d-centered orbitals and the spin coupled orbital pairs have >90% Co d-parentage. A similar electronic structure is observed for the hypothetical high spin ferromagnetically coupled species. These analyses indicated the electronic structure of **3** to be a dicobalt(II) oxo irrespective of the density functional employed (see ESI Section U†).

The unusual electronic structure of another dicobalt oxo species<sup>17</sup> encouraged us to examine the electronic structure of intermediate **3** more carefully. Therefore, we pursued CASSCF calculations (on the DFT-optimized geometry) using an active space that distributes 20 electrons into 13 orbitals (CASSCF(20,13)), the ten cobalt 3d-based orbitals and the three 2p orbitals of the oxo ligand. For the present purpose of



Fig. 3 CASSCF(20,13) natural orbitals along with occupation numbers in parentheses for the  $S = 0$  state of intermediate **3**. Atomic contributions to each orbital are also shown.

analyzing metal–ligand bonding, it is not necessary to employ an enlarged active space including double d and/or p shells of the metal center and the oxo ligand. As shown in Fig. 3, the singlet wavefunction has strong multireference character, with several competing configuration state functions having a weight in the range of 0.2–6.2% (Table S5†). Inspection of the natural orbitals obtained from the CASSCF(20,13) calculations shows that complex **3** is best described as having two high spin cobalt(II) centers bridged by an oxo ligand,  $Co^{II}-O^{2-}-Co^{II}$ . Notably, the electronic structure of **3** differs from that proposed for the previously reported Co–O–Co complex, which contains two cobalt(II) ions, an oxyl ligand, and a “hole” in the supporting ligand.<sup>17</sup> Note that due to the substantial multireference character, one cannot apply the CCSD approach to the singlet state. The septet CASSCF(20,13) solution is essentially single reference in nature, because the leading configuration accounts for 88% of the wavefunction. We further enlarged the active space to include diketiminate  $\pi$  and  $\pi^*$  orbitals (CASSCF(24,17)), which accounts for the possibility of ligand-based radicals, as discussed above for another dicobalt oxo complex found in the literature.<sup>17</sup> However, the larger CASSCF(24,17) active space predicts an identical bonding picture to CASSCF(20,13)



(for details, see ESI, Section P<sup>†</sup>). Therefore, we discuss only the CASSCF(20,13) solution.

In line with the DFT results, the Co–O interaction computed by CASSCF for **3** is very polar, with only ~20% cobalt 3d character in the oxygen-based orbitals and predominant cobalt d-parentage (>90%) in the metal-based orbitals. This is quite different from high-valent mononuclear metal–oxo species like ferryls, which feature more covalent metal–ligand interactions.<sup>53–56</sup> For example, in [Fe(O)(TMC)(NCCH<sub>3</sub>)]<sup>2+</sup> (TMC = tetramethylcyclam), there is 56% Fe(d) and 32% O(p) character in the Fe–O  $\sigma$ -bond and 54% Fe(d) and 36% O(p) in the Fe–O  $\pi$ -bond.<sup>53</sup> The different bonding picture in **3** can be attributed to the lower oxidation state of cobalt and the competitive bonding of the two metal centers with the oxo ligand. Furthermore, the calculated high electron density on the oxygen atom, as found from the population analysis (Mulliken gross atomic charges on O = –0.6195 a.u. and on Co centers = 0.6036 and 0.5989 a.u. at the BP86/def2-TZVPP level of theory; see ESI, Section U<sup>†</sup>), is consistent with facile nucleophilic attack on CO<sub>2</sub>. The bonding picture in **3** contrasts with high-valent metal–oxo intermediates, in which covalent metal–oxo interactions govern the electrophilic reactivity.<sup>46,57,58</sup>

### Kinetic studies on the CO<sub>2</sub> reduction pathway

Initial insight into the mechanism of CO<sub>2</sub> reduction by L<sup>tBu</sup>Co was gained through kinetic studies using <sup>1</sup>H NMR spectroscopy. After injection of a solution of excess CO<sub>2</sub> into a solution of **1** in C<sub>6</sub>D<sub>12</sub> at 10 °C, the reaction was monitored by NMR spectroscopy (see ESI Section C for details<sup>†</sup>). The concentrations of **1**, **2** and **4** fit to exponential decays over more than six half-lives. The first-order rate constant of  $3.7 \pm 0.5 \times 10^{-4} \text{ s}^{-1}$  was independent of the flooding concentration of CO<sub>2</sub>, indicating that the rate law has the form rate =  $k[1]$ . The zero-order dependence of the rate on [CO<sub>2</sub>] indicates that the rate-limiting step occurs prior to CO<sub>2</sub> binding and prevents the use of kinetic measurements to elucidate steps after CO<sub>2</sub> binding. Hence, we use computations to evaluate these steps below.

We considered that **3** could be formed as an intermediate that could react with CO<sub>2</sub> rapidly to give **4**; if its consumption were more rapid than its formation, it might not be observed during the reaction. With pure samples of **3** in hand, we tested this hypothesis. Treating a solution of **3** in C<sub>6</sub>D<sub>12</sub> with 1.5 equivalents of CO<sub>2</sub> for 2 min at 10 °C resulted in a high conversion (75%) of **3** to **4** (Scheme 3 and Fig. S5<sup>†</sup>). The much more rapid reaction of CO<sub>2</sub> with **3** compared to **1**, furnishing the same product, indicates that the reaction of the oxo species with CO<sub>2</sub> is kinetically competent to be a step in the formation of **2** and **4** from **1** and CO<sub>2</sub>.



Scheme 3 Reaction of CO<sub>2</sub> with **3** to form **4**.

### Computational investigations on the mechanism of CO<sub>2</sub> reduction

To gain additional insight into the reaction mechanism, we pursued computations using a model in which the bulky  $\beta$ -diketiminato ligand (L<sup>tBu</sup>) was slightly truncated to L' (2,4-bis[(2,6-diisopropylphenyl)imino]pent-3-yl, Chart 1, right) where *t*Bu substituents are replaced by methyl groups. The overlay of the crystal structure of **1** and the optimized geometry with the truncated ligand (Fig. S11 and Table S2<sup>†</sup>) shows that truncation leads to negligible differences in key metrical parameters. As above, the BP86(CPCM,D3BJ)/B1 level of theory was employed to optimize geometries and compute frequencies. We chose the local coupled-cluster approach with the DLPNO-CCSD(T) method to verify the reliability of crucial stationary points obtained with the *meta*-GGA M06L and hybrid B3LYP methods. We note that the CCSD(T) approach cannot be applied to complexes involving two antiferromagnetically coupled metal centers, such as the open-shell singlet state of complex **3**. Therefore, we performed the DLPNO-CCSD(T) calculations on the spin-aligned state of the bimetallic complexes. Although the CASSCF approach accounts for static correlation of electrons correctly (*i.e.* near-degeneracy effects), to produce reliable energies, it has to be followed by CASPT2 or NEVPT2 corrections to capture the dynamic correlation for instantaneous electron motions. Because of the exorbitant computational cost, this combined approach was not used to evaluate the energies of all intermediates and transition states. Instead, we employed open-shell DLPNO-CCSD(T), a straightforward approach, to construct reliable potential energy surfaces. For more details about the computational methods and basis sets used, see ESI, Section A.<sup>†</sup>

**Ligand isomerization and CO<sub>2</sub> binding.** We first focused on the ligand framework rearrangement and CO<sub>2</sub> binding, for which different CO<sub>2</sub> binding modes were considered as shown in Chart 1 (left). Our earlier kinetic studies demonstrated that coordination of pyridine to **1** induces isomerization of the  $\beta$ -diketiminato ligand from the  $\kappa N, \eta^6$ -arene isomer to the traditional  $\kappa^2 N, N'$  form in less than 1 s at –40 °C, with a barrier of  $\Delta G^\ddagger = 10.1 \text{ kcal mol}^{-1}$ .<sup>32</sup> The associative nature of this isomerization was supported by computational studies, which indicated that pyridine and CO each induce arene slipping.<sup>35</sup> On the other hand, we find here that CO<sub>2</sub> *does not* provide analogous assistance in arene slipping. Therefore, in contrast to the



Chart 1 (Left) Different binding modes of CO<sub>2</sub> to mononuclear and dinuclear metal sites. (Right) Actual and truncated ligand frameworks.



binding of pyridine and CO, the arene slip/imine flip isomerization must occur prior to CO<sub>2</sub> coordination, and this explains the slow rate of the reaction.

Compound **1** was previously termed a “masked two-coordinate” complex, and two-coordinate isomer **1'** with  $\kappa^2N,N'$  bonding (Fig. 4) is never observed experimentally. Our high-level computations with DLPNO-CCSD(T)/def2-TZVPP and M06L/def2-TZVPP show that two-coordinate isomer **1'** with a  $\kappa^2N,N'$  bonding mode lies at a higher energy than **1** (8.9 kcal mol<sup>-1</sup> at DLPNO-CCSD(T)/def2-TZVPP and 3.3 kcal mol<sup>-1</sup> at M06L/def2-TZVPP), consistent with the experimental results.

Furthermore, diketiminate isomerization *without* assistance is experimentally known to have a barrier of >15 kcal mol<sup>-1</sup> because no isomerization between binding modes was observed in variable-temperature <sup>1</sup>H NMR studies of L<sup>tBu</sup>Co.<sup>32</sup> Consistent with these experiments, our DLPNO-CCSD(T)/def2-TZVPP calculations predict a barrier of  $\Delta G^\ddagger = 24.5$  kcal mol<sup>-1</sup> for **TS1** (Fig. 4), which agrees well with the experimental rate that gives  $\Delta G^\ddagger = 21.0$  kcal mol<sup>-1</sup> using the Eyring equation (see Section C in the ESI<sup>†</sup>). The M06L functional with the def2-TZVPP basis set also delivers a similar value of 23.1 kcal mol<sup>-1</sup>, whereas B3LYP-D3BJ considerably underestimates the barrier with  $\Delta G^\ddagger = 13.6$  kcal mol<sup>-1</sup> (Table 3). It is notable that the B3LYP-D3BJ/def2-TZVPP calculations also erroneously predicted triplet  $\kappa^2N,N'$  isomer **1'** to be much lower in energy than  $\kappa N,\eta^6$ -arene isomer **1** by 10 kcal mol<sup>-1</sup>, in disagreement with the experimental data. Hence, for this specific case, the B3LYP-D3BJ method does not give results consistent with those of the experiment. Please note that our earlier calculations on CO<sub>2</sub> hydrogenation<sup>59,60</sup> demonstrate that even with the D3BJ corrections, the B3LYP functional sometimes cannot appropriately describe non-covalent interactions (in the present case, the metal aryl  $\pi$ -bonding), whereas M06L is designed to account for such weak interactions (see Section H in the ESI<sup>†</sup>).<sup>61</sup> Hence, in

Table 3 Selected barriers at different levels of theory, employing the def2-TZVPP basis set and CPCM(benzene) solvent model

|                         | B3LYP | M06L | DLPNO-CCSD(T) |
|-------------------------|-------|------|---------------|
| <b>TS1</b> <sup>a</sup> | 13.6  | 23.1 | 24.5          |
| <b>TS2</b> <sup>b</sup> | 30.4  | 27.2 | 26.7          |
| <b>TS4</b> <sup>c</sup> | 21.6  | 20.5 | 20.3          |
| <b>TS7</b> <sup>c</sup> | 16.0  | 15.7 | 19.4          |

<sup>a</sup> Relative to **1**. <sup>b</sup> Relative to **5**. <sup>c</sup> Relative to **10**.

the following sections we discuss the more reliable M06L energies and verify them on-the-fly by the DLPNO-CCSD(T) method and summarize the results obtained with other functionals in the ESI (Sections K and L<sup>†</sup>).

To summarize, the N1/C<sup>Ar</sup> movement of the metal center (**1** → **1'**) represents the rate-limiting transition state of the reaction with CO<sub>2</sub> (**TS1**, Fig. 4), with a calculated barrier of about 24 kcal mol<sup>-1</sup>. The magnitude of this barrier is in good agreement with the experimentally determined barrier of 21.0 kcal mol<sup>-1</sup> that is calculated from the first-order rate constant in the kinetic studies presented above (see ESI part C<sup>†</sup>), and the identification of the rate-determining transition state prior to CO<sub>2</sub> binding is also consistent with the independence of the rate on [CO<sub>2</sub>].

The closed-shell singlet  $\kappa N,\eta^6$ -arene isomer of L'Co (**1**) lies 10 kcal mol<sup>-1</sup> higher in energy than the triplet congener at the DLPNO-CCSD(T)/def2-TZVPP level of theory ( $\Delta G \sim 12$  kcal mol<sup>-1</sup> predicted with M06L/def2-TZVPP). Again, the M06L functional provides similar results to the DLPNO-CCSD(T) method and our calculations show that ligand isomerization on the closed shell singlet surface has a significantly high barrier ( $\Delta G^\ddagger = 49.6$  kcal mol<sup>-1</sup> using DLPNO-CCSD(T)/def2-TZVPP;  $\Delta G^\ddagger = 46.7$  kcal mol<sup>-1</sup> using M06L/def2-TZVPP). Therefore, the singlet surface is excluded from further consideration.

The slippage of the  $\beta$ -diketimate ligand furnishes an open site for CO<sub>2</sub> binding. In contrast to the high endergonicity found for the direct CO<sub>2</sub> coordination to **1**,  $\kappa^1O$ -bound CO<sub>2</sub>-adduct **1a'** is only 2 kcal mol<sup>-1</sup> higher in energy than **1'**. The  $\kappa^1O$  binding mode in **1a'** has been observed in the uranium complex ((<sup>Ad</sup>ArO)<sub>3</sub>tacn)U(CO<sub>2</sub>) where the CO<sub>2</sub> ligand accepts an electron from the uranium center to produce a U<sup>IV</sup>-OCO<sup>-</sup> complex.<sup>62</sup> As depicted in Fig. 4, conversion of **1a'** to a more stable  $\eta^2$ -CO<sub>2</sub> isomer (**5**) can easily occur by traversing a moderate barrier of 15 kcal mol<sup>-1</sup>. The two C–O bonds in **5** lengthen relative to those in **1a'**, and the CO<sub>2</sub> moiety becomes bent with an O–C–O angle of 142.5° (Fig. 5), indicating a substantial shift of electron density from Co to the CO<sub>2</sub>  $\pi^*$  orbital.<sup>59</sup> Intermediate **5** is related to PP<sup>Me</sup>PNi( $\eta^2$ -CO<sub>2</sub>) (PP<sup>Me</sup>P = P[Me-2-*Pi*-Pr<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>]<sub>2</sub>) and (dtbpe)Ni( $\eta^2$ -CO<sub>2</sub>) (dtbpe = 1,2-bis(di-*tert*-butylphosphino)ethane), which also feature  $\eta^2(C,O)$  coordination.<sup>63,64</sup> Earlier computational investigations have suggested that CO<sub>2</sub> coordination to metal  $\beta$ -diketimate fragments (M = Cr, Ni) favors the  $\eta^2(C,O)$  binding mode, similar to intermediate **5**.<sup>65</sup>

**Mononuclear dissociative pathway.** Two possible pathways may be envisioned for CO<sub>2</sub> reduction by **1**: associative and dissociative. The associative mechanism involves formation of



Fig. 4 Computed DLPNO-CCSD(T)/def2-TZVPP Gibbs free energy ( $\Delta G$ ) profile for isomerization of **1** (L'Co) to **1'** followed by CO<sub>2</sub> activation. Relative free energy values (in kcal mol<sup>-1</sup>) computed with M06L/def2-TZVPP are given in parentheses. The isolated species is shown in the yellow box, with L' depicted instead of L<sup>tBu</sup>.





Fig. 5 Optimized structures of **1**, **TS1**, **TS1a'** and **5** at the BP86/B1 level of theory. Important interatomic distances (Å) and angles (°) are shown. Isopropyl groups are omitted for clarity.

a CO<sub>2</sub> adduct followed by association of a second molecule of CO<sub>2</sub> prior to C–O bond cleavage/rearrangement to generate CO and carbonate. In contrast, adduct formation in the dissociative mechanism is followed by breaking a C–O bond to form CO and oxo, both of which are easily converted to products. We start by considering the mononuclear dissociative pathway (Fig. 6).

Due to the abundance of CO<sub>2</sub> in solution, an additional CO<sub>2</sub> molecule can be envisioned to add to intermediate **5** to produce a metal-bound carboxylate dimer. However, our repeated attempts to locate an intermediate resulting from association of a second CO<sub>2</sub> molecule to **5** failed (see ESI, Section S<sub>4</sub><sup>†</sup>). The lack of a mononuclear associative pathway emphasizes the need for bi-functional activation to engender sufficient nucleophilicity to bound CO<sub>2</sub>.<sup>48</sup> The rather low nucleophilicity of the bound CO<sub>2</sub> in **5** is not conducive to electron transfer to another incoming CO<sub>2</sub>. Moreover, the lack of a neighboring Lewis acid, which might help in bending of the incoming CO<sub>2</sub> prior to activation,

also inhibits facile association of a second CO<sub>2</sub> molecule to the mononuclear adduct **5**, and hence a mononuclear associative pathway is not possible (see ESI, Section S<sub>4</sub><sup>†</sup>).

As depicted in Fig. 6, our DLPNO-CCSD(T) calculations predict that the conversion of **5** to L'Co(CO)(O) (**6**, see ESI Section M for geometry<sup>†</sup>) involves a high free-energy barrier in the mononuclear dissociative pathway (**TS2**, Table 3). Similar barriers were obtained using DLPNO-CCSD(T) (26.7 kcal mol<sup>-1</sup>), M06L (27.2 kcal mol<sup>-1</sup>) and B3LYP (30.4 kcal mol<sup>-1</sup>) calculations, and the latter value is in reasonable agreement with that (35.1 kcal mol<sup>-1</sup>) calculated in earlier work on a truncated cobalt(i) complex with no substituents on the β-diketiminato (1,3-bis-imino-prop-2-yl).<sup>66</sup> Once formed, complex **6** easily transfers CO to **1'**, leading to the mononuclear carbonyl complex (**2**) and a terminal oxo-cobalt species (**7**, Fig. 6). Complex **7** may initiate nucleophilic attack of the metal-coordinated oxo on incoming CO<sub>2</sub> to form a mono-cobalt carbonate species (**9**) via a four-membered transition state (**TS3**, ΔG<sup>‡</sup> = 9.4 kcal mol<sup>-1</sup>, Fig. 6). One may envision formation of the bridging oxo **3** from **6** or **7** in the presence of **1**. The transformations were computed to have favorable driving forces; however, in the mononuclear dissociative pathway, the generation of **6** entails a high barrier for the C–O bond cleavage. Hence, these two pathways were not considered further (for details, see Section S in the ESI<sup>†</sup>). Additionally, in the mononuclear pathway, the high barrier found for the C–O bond cleavage would lead to a buildup of **5**, in contrast to the observed reaction course in which no intermediates are evident. This inconsistency with experimental evidence thus indicates that the mononuclear pathway is not feasible. We therefore explored the feasibility of the dinuclear dissociative and associative pathways, which are discussed in the following section.

**Dinuclear pathways.** We considered a carboxylate bridged bimetallic species that could lead to reductive disproportionation. As shown in Fig. 6, reaction of L'CoCO<sub>2</sub> (**5**) and L'Co (**1**) to form the dinuclear complex L'Co(CO<sub>2</sub>)CoL' (**10**) is strongly exergonic by 21.8 kcal mol<sup>-1</sup> and has a relatively low activation barrier of 10.3 kcal mol<sup>-1</sup> (**TS2'**, Fig. 6) on the quintet surface. Isomerization of the diketiminato ligand in the incoming L'Co is assisted by coordination of the bound CO<sub>2</sub> to the incoming metal, as described previously for pyridine or CO coordination; thus, the partially reduced CO<sub>2</sub> moiety in **5** is a stronger Lewis base than free CO<sub>2</sub>. In line with this reasoning, the entire process of CO<sub>2</sub> association with **1** to give **5** is energetically uphill, whereas the addition of **5** to the second molecule of **1** is downhill by more than 20 kcal mol<sup>-1</sup>. This is a key aspect of *bimetallic cooperation* that facilitates CO<sub>2</sub> activation in this system. For complex **10**, we found eight possible isomers, each containing a κ<sup>1</sup>C:κ<sup>2</sup>-O,O carboxylate bridge (CO<sub>2</sub><sup>2-</sup>) (Chart 1, left), but differing in the local spin states of the two cobalt(II) centers and exchange coupling (for details see Section J in the ESI<sup>†</sup>). The two magneto-structural isomers that are lowest in energy have high spin cobalt(II) centers with ferromagnetic (S<sub>total</sub> = 3) or antiferromagnetic (S<sub>total</sub> = 0) coupling, and these are nearly isoenergetic because of weak exchange coupling. The S<sub>total</sub> = 2 species (derived from ferromagnetically coupled high spin and low spin cobalt(II) centers) is only 3 kcal mol<sup>-1</sup> higher



Fig. 6 M06L/def2-TZVPP Gibbs free energy (ΔG) profile for the mononuclear dissociative pathway. Isolated species are shown in yellow boxes, with L' depicted instead of L<sup>tBu</sup>. \*Marked species has lower relative free energy on the S = 3 surface which has been taken to complete the profile.



in energy; thus, spin crossover from the quintet to the septet state is feasible (see Section L in the ESI<sup>†</sup>). We assume that the efficient spin orbit coupling of the metal centers is sufficient to minimize any spin-crossover barriers, for which only the local spin state of the cobalt center changes. Starting from complex **10**, we tested dissociative and associative mechanisms. In the dissociative route (toward the right of Fig. 7), **10** first undergoes C–O bond dissociation to generate CO and a  $\mu$ -oxo species, and then addition of another CO<sub>2</sub> molecule to the latter intermediate leads to formation of the carbonate-bridged bimetallic product. This reaction channel has been proposed for reductive cleavage of CO<sub>2</sub> by L<sup>tbu</sup>Fe–N<sub>2</sub>–FeL<sup>tbu</sup> and low-valent U(III) complexes with the intermediacy of an M–O–M species.<sup>62,67</sup> In the associative route (toward the left of Fig. 7), another CO<sub>2</sub> molecule first inserts into **10** yielding a (CO<sub>2</sub>)<sub>2</sub> linker, which then rearranges to CO and CO<sub>3</sub><sup>2-</sup>. The associative mechanism was described for CO<sub>2</sub> functionalization using [Re(dmb)(CO)<sub>3</sub>] (where dmb = 4,4'-dimethyl-2,2'-bipyridine) by Agarwal *et al.*<sup>68</sup>

In the dissociative pathway, C–O bond cleavage in **10** passes through **TS4** (see the geometry in Fig. 8) to afford an intermediate, L'Co(CO)–O–CoL' (**11**), where two cobalt(II) metal centers are bridged by an oxo group, and a terminal carbonyl ligand is bound to one cobalt(II) center. The transformation is thermo-neutral ( $\Delta G = -0.9$  kcal mol<sup>-1</sup> relative to **10**) and involves a moderate barrier of 20.3 kcal mol<sup>-1</sup> at the coupled-cluster level of theory (DLPNO-CCSD(T)/def2-TZVPP, Table 3) on the *S* = 3 surface while M06L/def2-TZVPP also predicts a similar result ( $\Delta G^\ddagger = 20.5$  kcal mol<sup>-1</sup>). Furthermore, M06L calculations using the broken symmetry formalism<sup>69,70</sup> also furnish a similar barrier (21.9 kcal mol<sup>-1</sup>) on the singlet surface (Fig. S17<sup>†</sup>). We note that attempts to independently synthesize **11** by addition of 1 eq or 1 atm CO to the oxo **3** instead furnished the previously reported complex L<sup>tbu</sup>Co(CO)<sub>2</sub> (see ESI Section D<sup>†</sup>).<sup>35</sup>



Fig. 8 Optimized structures of **3**, **4** and **TS2'** at the BP86/B1 level of theory and **10**, **TS4** and **TS7** at the B3LYP/B1 level of theory. (See computational details for basis set information and justification of employing the respective methods.) Important interatomic distances (Å) and angles (°) are also shown. Isopropyl groups are omitted for clarity.

As depicted in Section L in the ESI<sup>†</sup>, further conversion of complex **11** to final product **4** has similar reaction energies and barriers for the different spin coupling situations. Therefore, we focus on the septet surface only (computed using M06L(CPCM)/def2-TZVPP) and summarize the results of the singlet state in Section L in the ESI<sup>†</sup>. CO transfer from **11** to **1** to form the three-coordinate L'Co(CO) product (**2**) and an oxidicobalt(II) species (**3**) is computed to be highly thermodynamically favored with a reaction free energy of *ca.* –32 kcal mol<sup>-1</sup>.



Fig. 7 M06L/def2-TZVPP Gibbs free energy ( $\Delta G$ ) profile for dinuclear dissociative (right) and associative (left) pathways. Isolated species are shown in yellow boxes, with L' depicted instead of L<sup>tbu</sup>. \*Marked species has lower relative free energy on the *S* = 0 surface which has been taken to complete the profile.



The addition of CO<sub>2</sub> to the oxo ligand in **3** can easily occur with a low barrier of 5.8 kcal mol<sup>-1</sup>, in agreement with the observation (see above) that L<sup>tBu</sup>CoCoL<sup>tBu</sup> reacts rapidly with CO<sub>2</sub> at room temperature to yield **4**.

For the associative mechanism, our calculations show that nucleophilic addition of CO<sub>2</sub> to the electron-rich oxygen atom of the bridging carboxylate dianion in **10** can form a C(O)–O–C(O)–O<sup>2-</sup> bridge between the two cobalt(II) centers in **14** *via* a four-membered transition state (**TS6**) that lies 13.5 kcal mol<sup>-1</sup> above **10**. This is followed by rupture of a C–O bond in the original CO<sub>2</sub> through **TS7** (Fig. 8), which has ΔG<sup>‡</sup> = 15.7 kcal mol<sup>-1</sup> with M06L/def2-TZVPP and 19.4 kcal mol<sup>-1</sup> with DLPNO-CCSD(T)/def2-TZVPP, leading to intermediate **15**. This stepwise insertion/rearrangement process is thermoneutral (ΔG = -0.9 kcal mol<sup>-1</sup>) for the septet state. **15** is a dicobalt(II) carbonate complex with a CO ligand coordinated to one metal center, which is readily transferred to **1** to afford the final products. Comparison of the barriers required for C–O bond breaking as computed for the three pathways shows that the reaction of CO<sub>2</sub> with L<sup>tBu</sup>Co is not likely to follow the mononuclear dissociative pathway, while either of the dinuclear pathways is possible because the computed barriers for **TS4** and **TS7** are similar, especially those calculated by the more accurate DLPNO-CCSD(T) approach. In contrast to the mononuclear intermediate **5**, which does not undergo an associative mechanism, intermediate **10** contains two cobalt(II) centers bridged by a CO<sub>2</sub><sup>2-</sup> anion (discussed below on the basis of frontier orbitals) and facilitates further CO<sub>2</sub> addition.

For both dinuclear pathways, free energy barriers calculated with B3LYP and M06L functionals employing the def-TZVPP basis set and conducted using the CPCM solvation model were in good agreement with the DLPNO-CCSD(T) benchmark for important transformations, like C–O bond-cleavage *via* **TS2**. A similar situation is also observed for **TS4** and **TS7** (Table 3). Hence, apart from the ligand isomerization step (**TS1**), our theoretical results are analogous to those computed by the coupled-cluster approach and are evidently *not* functional dependent.

**Frontier-orbital analysis of C–O cleavage in different reaction pathways.** As shown in Fig. 9a using the M06L/def2-TZVPP technique, coordination of CO<sub>2</sub> to **1** in the mononuclear complex **5** only slightly perturbs the σ and π interactions within CO<sub>2</sub>, because the doubly occupied orbital termed σ<sub>Co–C</sub> (consisting of the Co d<sub>z<sup>2</sup></sub> and CO<sub>2</sub> π\* orbitals) contains a mere 30% carbon contribution. This indicates that there is only partial electron transfer from the metal center to CO<sub>2</sub>, consistent with the difference of only 0.06 Å between the two C–O bond distances (Fig. 5). During the transformation of **5** to **6**, both σ and π bonds undergo heterolytic cleavage, which is accompanied by two-electron transfer from the Co d<sub>z<sup>2</sup></sub> orbital to the CO<sub>2</sub> π\* orbital. As a consequence, the CO<sub>2</sub> π\* orbital eventually transforms into the CO lone pair in complex **6**, and the cobalt(I) center in **5** is oxidized to cobalt(III) (Fig. 9a). This is reminiscent of the C–F bond cleavage by **1** which was studied using kinetics<sup>33</sup> and computations<sup>71</sup> that indicated oxidative addition of the C–F bond to give a four-coordinate cobalt(III) product. However, the fluoride-containing product of C–F activation is much more stable than the oxo-containing product of C–O activation, and therefore *the mononuclear pathway is disfavored for CO<sub>2</sub> cleavage*.

In the dicobalt intermediate **10**, the CO<sub>2</sub> moiety is reduced by nearly two electrons, with each metal center donating one electron (Fig. 9b), and the Co–C σ-bonding orbital is predominantly carbon based. The greater electron transfer from the metal to CO<sub>2</sub> in **10** relative to **5** reflects the greater ease of oxidizing each metal by only one electron. Furthermore, the dinuclear dissociative pathway entails lower reorganization energy during C–O bond cleavage, because the oxidation states of the metal centers do not change. As shown in ESI, Section Q,† the conversion of **14** to **15** *via* **TS7** in the dinuclear associative pathway involves similar electronic structure changes. The target C–O σ-bond cleavage in **TS7** results in conversion of the bonding orbital to the oxygen p orbital (Fig. S19†). Concurrently the same oxygen starts interacting with one cobalt(II) center to form a new Co–O bond. This cooperation between the two metals facilitates C–O bond cleavage and makes the bimetallic pathways dominant.



Fig. 9 Important orbitals during C–O cleavage mediated by the (a) mononuclear dissociative pathway and (b) dinuclear dissociative pathway at the M06L/def2-TZVPP level of theory.



## Literature perspectives

The reaction of CO<sub>2</sub> with L<sup>tBu</sup>Co to give **2** and **4** represents the net reductive cleavage of two molecules of CO<sub>2</sub> to CO and CO<sub>3</sub><sup>2-</sup>. Chemical fixation and transformation of CO<sub>2</sub> has been reported in many earlier studies of dinuclear macrocyclic complexes where CO<sub>2</sub> is reduced to form carbonate and CO.<sup>72–75</sup> One of the earliest studies by Sorrell *et al.* included crystallographic characterization of a carbonate-bridged dinuclear copper complex from the reaction of CO<sub>2</sub> with a peroxo-dicopper(II) intermediate.<sup>72</sup> Homogeneous electrocatalytic CO<sub>2</sub> reduction to form CO was originally reported in a well-characterized cobalt system by Eisenberg<sup>76</sup> and developed in the research of DuBois<sup>77,78</sup> and Kubiak,<sup>79</sup> and many further advances in cobalt-based CO<sub>2</sub> reduction to CO have been described by Fujita and others.<sup>20,23–25,30</sup> This continues to be an active area of research by many groups, and therefore mechanistic insights are important.<sup>13,26–29,31</sup>

Stoichiometric reduction of CO<sub>2</sub> to CO using well-defined late 3d metal systems is also relevant. Peters reported that a Fe tris(phosphino)borate complex reacts with CO<sub>2</sub> to give binuclear μ-O:μ-CO and C–C coupled μ-oxalato complexes, wherein the relative yields of the two products can be controlled by the choice of solvent.<sup>80,81</sup> CO<sub>2</sub> reduction to CO using a pyridinediimine stabilized iron(II) precatalyst has also been reported.<sup>82</sup> Sadighi *et al.* identified a unique CO<sub>2</sub>/CO bridged dinuclear (NHC)Ni complex.<sup>83</sup> A particularly cogent precedent is the heterobimetallic cobalt complex Co(<sup>t</sup>Pr<sub>2</sub>PNMe)<sub>3</sub>Zr(THF), which cleaves CO<sub>2</sub> to CO and oxo (Scheme 4a).<sup>24,84</sup>

The reaction described here between CO<sub>2</sub> and the cobalt(I) source L<sup>tBu</sup>Co can be compared to the published reactions of CO<sub>2</sub> with the iron(I) source L<sup>tBu</sup>FeNNFeL<sup>tBu</sup> (Scheme 4b),<sup>37</sup> the nickel(I) source L<sup>tBu</sup>NiNNNiL<sup>tBu</sup> and the nickel(0) source K<sub>2</sub>[L<sup>tBu</sup>NiNNNiL<sup>tBu</sup>] (Scheme 4c).<sup>85</sup> The carbonate products from the Fe and Co reactions are very similar, but the CO products are different. While the iron dinitrogen complex gave the dicarbonyl complex L<sup>tBu</sup>Fe(CO)<sub>2</sub>,<sup>37</sup> the cobalt analogue gave instead

the monocarbonyl complex L<sup>tBu</sup>Co(CO). This difference can be attributed to the extent of backbonding. L<sup>tBu</sup>Fe(CO)<sub>2</sub> displays lower CO stretching frequencies ( $\nu_{\text{CO}} = 1994, 1915 \text{ cm}^{-1}$ ) as compared to L<sup>tBu</sup>Co(CO)<sub>2</sub> ( $\nu_{\text{CO}} = 2014, 1949 \text{ cm}^{-1}$ ), and thus backbonding is weaker in L<sup>tBu</sup>Co(CO)<sub>2</sub> than in the analogous Fe complex. Consistent with weaker bonding in the cobalt dicarbonyl *versus* the iron dicarbonyl, reaction of L<sup>tBu</sup>Co(CO)<sub>2</sub> with L<sup>tBu</sup>Co afforded the monocarbonyl complex, whereas L<sup>tBu</sup>Fe(CO) is unknown. Thus, we surmise that the difference in the product distribution between iron and cobalt arises from the relative M–C bond energy of the second CO. In the nickel case, Horn *et al.* reported two related reactions.<sup>85</sup> First, the reaction of the dinickel(I) complex L<sup>tBu</sup>NiNNNiL<sup>tBu</sup> with CO<sub>2</sub> resulted in reductive coupling to form an oxalate complex (Ni<sup>2+</sup><sub>2</sub>C<sub>2</sub>O<sub>4</sub>Ni<sup>2+</sup>), which was not observed in the analogous cobalt system. We speculate that the smaller nickel(I) center cannot access a L<sup>tBu</sup>NiONiL<sup>tBu</sup> intermediate and thus is diverted to C–C coupling. On the other hand, reaction of the nickel(0) complex K<sub>2</sub>[L<sup>tBu</sup>NiNNNiL<sup>tBu</sup>] with CO<sub>2</sub> gave a nickel(I)–CO complex and a nickel(II) complex of CO<sub>3</sub><sup>2-</sup>, which is more similar to the results described here.

It is also worthwhile to compare our results to the electrochemical reduction of CO<sub>2</sub> to CO catalyzed by [Ni(cyclam)]<sup>+</sup>, which we have previously examined computationally.<sup>86</sup> These studies showed that formation of an η<sup>1</sup>-CO<sub>2</sub> adduct [Ni(η<sup>1</sup>-CO<sub>2</sub>)(cyclam)]<sup>+</sup> causes partial electron transfer from the metal center to CO<sub>2</sub>, similar to complex **5** in the present work. The second electron reduction of CO<sub>2</sub> in the [Ni(cyclam)]<sup>+</sup> case requires proton donors, where Lewis acid stabilization of [Ni<sup>II</sup>(C(O)OH)(cyclam)]<sup>+</sup> enables facile C–O bond cleavage to generate [Ni(CO)(cyclam)]<sup>2+</sup> and H<sub>2</sub>O. Lewis acid stabilization facilitating CO<sub>2</sub> cleavage has also been shown clearly in iridium and iron systems by Bernskoetter and Hazari.<sup>87,88</sup> Here, the second cobalt stabilizes the growing negative charge on the bound CO<sub>2</sub> like a Lewis acid; additionally, the second cobalt in the system described here provides a second electron that is crucial for CO<sub>2</sub> reduction.

## Conclusions

The masked two-coordinate cobalt(I) complex L<sup>tBu</sup>Co cleaves the C–O bond in CO<sub>2</sub> to form a mononuclear carbonyl complex and a dinuclear carbonate complex. Both DFT and the highly correlated wavefunction-based DLPNO-CCSD(T) calculations show two feasible bimetallic pathways, and the dissociative one is supported by the independent isolation of a kinetically competent dicobalt(II) oxo complex. The electronic structure of this unusual oxo complex was analyzed using the multi-reference CASSCF approach, which showed that the oxo compound is best described as a dicobalt(II) complex, in contrast with an earlier literature compound with redox-active oxyl radical character. The negatively charged oxygen in the new dicobalt(II) oxo complex has nucleophilicity that leads to its rapid reactivity toward carbon dioxide. Overall, the cooperative interactions of the two cobalt centers facilitate charge buildup on CO<sub>2</sub>, leading to facile activation of this strong bond.



Scheme 4 Selected CO<sub>2</sub> reduction reactions from the literature.



## Conflicts of interest

The authors declare no conflicts of interest.

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