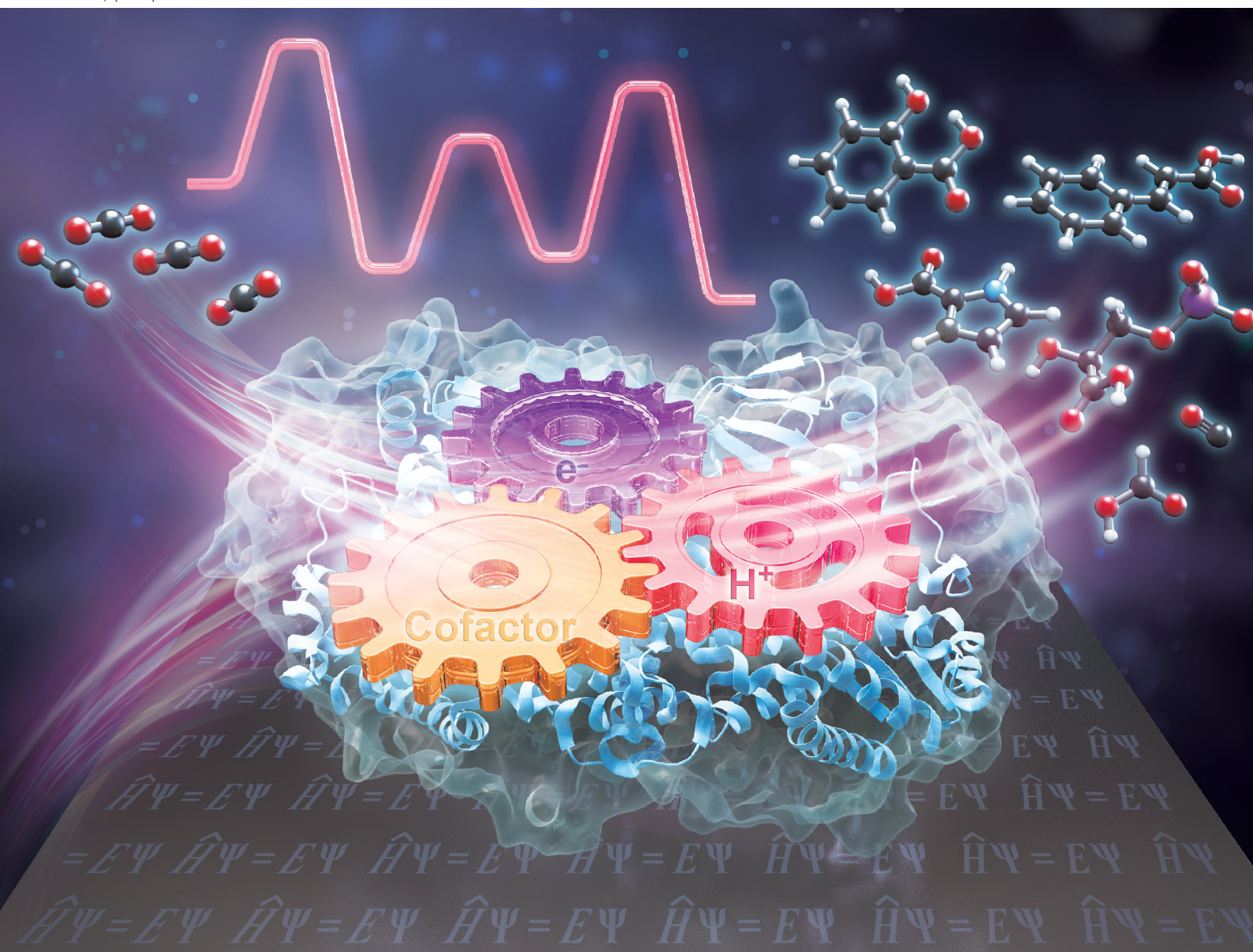


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Quantum chemical studies of the reaction mechanisms of enzymatic CO₂ conversion

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Enzymatic capture and conversion of carbon dioxide (CO₂) into value-added chemicals are of great interest in the field of biocatalysis and have a positive impact on climate change. The quantum chemical methods, recognized as valuable tools for studying reaction mechanisms, have been widely employed in investigating the reaction mechanisms of the enzymes involved in CO₂ utilization. In this perspective, we review the mechanistic studies of representative enzymes that are either currently used or have the potential for converting CO₂, utilizing the quantum chemical cluster approach and the quantum mechanical/molecular mechanical (QM/MM) method. We begin by summarizing current trends in enzymatic CO₂ conversion, followed by a brief description of the computational details of quantum chemical methods. Then, a series of representative examples of the computational modeling of biocatalytic CO₂ conversion are presented, including the reduction of CO₂ to C1 species (carbon monoxide and formate), and the fixation of CO₂ to form aliphatic and aromatic carboxylic acids. The microscopic views of reaction mechanisms obtained from these studies are helpful in guiding the rational design of current enzymes and the discovery of novel enzymes with enhanced performance in converting CO₂. Additionally, they provide key information for the *de novo* design of new-to-nature enzymes. To conclude, we present a perspective on the potential combination of machine learning with quantum description in the study of enzymatic conversion of CO₂.

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1. Introduction

As a trace gas in Earth's atmosphere, the concentration of CO₂ has risen by more than 50% from pre-industrial levels of 280 parts per million (ppm) to 423.65 ppm as of April 2024,¹ largely due to anthropogenic excess emissions. Atmospheric CO₂ levels have been consistently increasing on an annual average, according to data measured by NOAA's Global Monitoring Laboratory.² This persistent rise has made CO₂ capture and utilization an urgent global challenge and a booming area of research. Enzymatic CO₂ conversion, involved in both primary and secondary metabolic pathways, is considered a green and sustainable option.^{3–7} Natural CO₂ fixation pathways include the CBB cycle, the reductive glycine pathway, the Wood–Ljungdahl pathway, the reverse TCA cycle, the 3HP/4HB cycle, the 3HP bicycle, and the dicarboxylate/4-hydroxybutyrate cycle. Modifying these natural pathways or *de novo* designing new synthetic pathways provides efficient solutions for CO₂ capture

and utilization. Examples include ASAP,⁸ CETCH,^{9,10} HOPAC,¹¹ rGPS-MCG,¹² POAP,¹³ and GED.¹⁴

The central carbon atom in CO₂ is in its highest oxidation state, rendering its formation energetically favored. Therefore, the utilization of CO₂ requires the provision of electrons or energy to pull it from the '*bottom of the potential energy well*'.¹⁵ Some enzymes have evolved to rely on cofactors, such as NADPH and NADH, which can transfer a hydride ion, thereby donating two electrons along with energy.^{14,16–18} Others utilize electron transfer agents known as ferredoxins, which are capable of producing a reduction potential of around –500 mV when the ferredoxin pool is 95% reduced.^{19,20} Another strategy involves using the direct energy from the hydrolysis of ATP.^{21,22} Additionally, certain enzymes coordinate CO₂ binding with bivalent metal ions in the active site, thereby activating it.^{23,24} Furthermore, enzymes such as RuBisCO²⁵ and phosphoenolpyruvate carboxylase (PEPC)²⁶ utilize their high-energy substrates, which are ribulose-1,5-bisphosphate (RuBP) and phosphoenolpyruvate (PEP), respectively, to drive their respective reactions.

Enzymatic CO₂ conversion consists of mainly two types of reactions: the reduction of CO₂ to carbon monoxide or formate by dehydrogenases,^{27,28} and the carboxylation, which introduces an additional carbon atom to a substrate, utilizing CO₂ as the carbon source.^{23,24} The carboxylation mechanism

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generally comprises two key steps: the generation of an enol or enolate nucleophile, and the nucleophilic attack of the enol (or enolate) on the carbon of CO₂ to form a C–C bond.³ However, it is important to note that certain enzymes, such as prenylated FMN-dependent ferulic acid decarboxylases²⁹ and thiamin pyrophosphate (TPP)-dependent ferredoxin oxidoreductase,^{30,31} follow different pathways. Understanding the reaction mechanisms of enzymatic CO₂ conversion is crucial for developing new strategies for CO₂ capture and utilization.

Quantum chemical (QC) methods have become essential tools for studying different aspects of enzymatic reactions, including substrate specificity, reaction mechanism and various selectivities.^{32–39} While biocatalytic CO₂ conversion follows some common rules, each reaction exhibits unique features due to the differences in substrates and enzyme properties. In-depth quantum chemical illustrations of enzymatic CO₂ reduction and carboxylation can elucidate specific reaction steps and identify key factors that shift the equilibrium toward the energetically less favorable direction. Moreover, these depictions can provide critical information for the enzyme design of new catalysts with enhanced efficiency or tailored properties, thereby improving their potentials in industrial applications.

In this perspective, we present the utilization of quantum chemical calculations, specifically through the cluster approach and the hybrid quantum mechanical/molecular mechanical (QM/MM) method, in investigating the reaction mechanisms of enzymatic CO₂ conversion. The technical aspects of the two methods are first briefly discussed. Then, we illustrate case studies of enzymatic transformation of CO₂ into various compounds, including C1 species, aliphatic acids, and aromatic carboxylic acids. Furthermore, we highlight the promising integration of machine learning with quantum chemical techniques, which represents a significant advancement that could open up new possibilities for accelerating the discovery and optimization of CO₂-converting enzymes with enhanced catalytic performance.

2. Methodology

2.1 Quantum chemical cluster approach

The quantum chemical cluster approach, also known as the all-QM approach, has been extensively applied to elucidate various aspects of enzyme catalysis.^{34–37} It allows a direct comparison of energy profiles for all possible reaction pathways, which is invaluable in identifying the most favorable substrate binding mode to the enzyme, as well as in elucidating the most plausible reaction mechanism that the enzyme may adopt. Moreover, the cluster approach has recently been expanded to model various selectivities of enzymatic reactions, and to provide insights into the pivotal roles of metal in the metal-dependent enzymes. Some of these studies identified the roles of key residues in catalysis, which are crucial for enzyme engineering and rational design.

The basic idea of the cluster approach is to focus on a small, chemically relevant portion of the enzyme's active site, along

with its proximate surroundings. The entire model is treated using the quantum mechanical method. Historically, it has been demonstrated that even a small active site model consisting of less than 100 atoms can effectively elucidate the reaction mechanism. Nowadays, a cluster model can contain more than 400 atoms, including substrates, organic cofactors, metal ions, and neighboring residues that constitute the active site. This expansion enhances the ability of the cluster approach to model selectivities and strategically design mutants. However, it is suggested that one should begin mechanistic investigations with a relatively small model, then gradually increase the model size to detect multiple minima errors and gain chemical insights into different functional groups. To accurately simulate the enzyme systems, a selection of atoms at the periphery of the model are fixed to mimic the steric influence of the actual binding pocket, and an implicit homogeneous polarizable solvation medium, generally with a dielectric constant $\epsilon = 4$, is used to provide an approximate description of solvation effect.

Among all the quantum chemical methods, density functional theory (DFT), particularly the B3LYP hybrid functional, which balances speed and accuracy well, has been broadly used. When describing dispersion interactions, one can add an empirical dispersion correction to the B3LYP calculations, for example, Grimme's D3 dispersion correction. Alternatively, the Minnesota M06 suite, which includes weak interactions in its training set, can also be used. Geometries are often optimized using the 6-31G(d,p) basis set, while single-point energies are typically calculated with a larger basis set, such as 6-311++G(2d,2p). For metals, the SDD⁴⁰ pseudopotential basis set is commonly employed. It is important to note that different theoretical methods may be adopted depending on the specific requirements of each study. For a detailed methodology, readers are referred to the original paper.

The entropy effect is usually not considered in the cluster approach, because the constraints at the periphery of the model render the entropy calculation inaccurate and the entropy effect is deemed to contribute very little to the barriers. However, when it comes to reactions involving the gas molecule entering or exiting an enzyme pocket, the translational entropy of the gas molecule is suggested to be factored into the calculations. Alternatively, harmonic confining potentials can be introduced to the computational model to achieve unconstrained geometry relaxations.^{41,42}

2.2 Quantum mechanical/molecular mechanical (QM/MM) method

The quantum mechanical/molecular mechanical (QM/MM) method is a robust computational method designed to investigate complex systems with high efficiency.^{38,39} It combines accuracy with speed by calculating the key region using a high level of QM theory, similar to the QC cluster approach discussed above, and its environment with a low-cost level of QM theory or a classical molecular mechanics (MM) simulation. This dual-level strategy allows for a comprehensive analysis of reaction mechanisms and other properties within complex biological systems.

Depending on the way of dealing with the interface between the two types of systems, there are subtractive and additive

schemes to calculate the overall potential energy by QM/MM method. In these methods, the real catalytic system is artificially divided into two parts, and the overall energy is composed of particle interactions within the quantum mechanical (QM) region, the atomic interactions within the molecular mechanical (MM) region, and the interactions between QM and MM particles. In the subtractive scheme, the potential energy of the total system is calculated by summing the MM-level energy of the whole system and the QM-level energy of the isolated QM system, then subtracting the MM energy of the QM system. In the additive schemes, which are considered more accurate, the potential energy of the total system is calculated as the sum of QM region energy terms, classical MM region energy terms, and QM/MM boundary energy terms.

As the QM/MM method provides an approximation of the potential energies of embedded quantum-chemical systems in a larger molecular environment, the proper treatment of the interaction between the QM and MM regions is important for the accuracy of the method. The QM/MM coupling boundary is composed of both non-bonded and bonded parts. For the non-bonded part, van der Waals interactions and electrostatic interactions demand careful consideration. The latter can be calculated using one of three methods: mechanical embedding, electronic embedding, or polarized embedding. For the covalently bonded part, it is necessary to saturate the dangling bond of the QM system to prevent the appearance of bond cleavage. To this end, and also to mitigate the potential over-polarization, three schemes have been developed: link-atom schemes, boundary atom schemes and localized-orbital schemes. In practice, link-atom schemes are more commonly used than the others because of its simplicity and robustness.

In the QM/MM calculations on enzymatic reactions, numerous minima and transition states are expected to be identified due to the extensive configuration space of studied large molecules. Therefore, multiple snapshots from classical MD simulations should be chosen as starting structures for the mechanistic investigation to ensure that the most plausible pathway is obtained. Alternatively, one can employ the computationally demanding QM/MM molecular dynamics calculations to fulfill the requirement for thorough sampling of the configuration space.

In the following section, we will present examples that demonstrate how quantum chemical computational research can aid in elucidating the reaction mechanism of CO₂-converting enzymes and provide relevant information. The discussed enzymes are those catalyzing the reduction of CO₂ to form C1 compounds (Section 3.1), and the carboxylation to generate aliphatic acids (Section 3.2) and aromatic acids (Section 3.2). This perspective paper ends with an outlook on the promising integration of machine learning with quantum chemical techniques, highlighting their potential application in CO₂ fixation.

3. Emerging practices in application

Quantum chemical calculations allow us to investigate mechanistic proposals involving the enzyme-substrate complex, intermediates, transition states, and the enzyme-product complex.

By assessing the calculated energies, we can evaluate the feasibility of these proposals. In the following sections, case studies of the reaction mechanisms of CO₂ conversion into various compounds, including C1 species, aliphatic acids, and aromatic carboxylic acids, are presented. For the reduction of CO₂ to C1 species, the discussed enzymes are the molybdenum-containing formate dehydrogenase (Mo-FDH), the Ni, Fe-containing carbon monoxide dehydrogenase (CODH), and the artificial human carbonic anhydrase (hCAII). In the section on the fixation of CO₂ to form aliphatic acids, the ribulose 1,5-bisphosphate carboxylase (RuBisCO), the crotonyl-CoA carboxylases/reductase (CCR), the biotin-dependent pyruvate carboxylase (PC) and two ThDP-dependant enzymes are discussed. For the carboxylation to generate aromatic carboxylic acids, the prFMN-dependent UbiD family and the (de)carboxylases from the amidohydrolase superfamily are presented. All the enzymes and their catalytic reactions illustrated in this paper are listed in Table 1.

3.1 Reduction of carbon dioxide to C1 compounds

3.1.1 Molybdenum-containing formate dehydrogenase (Mo-FDH). Formate dehydrogenases (FDHs) are a group of enzymes that catalyze the reduction of CO₂ to formate, consuming one proton and two electrons. Among the identified formate dehydrogenases, the molybdenum-containing formate dehydrogenases (Mo-FDHs) from *Escherichia coli* achieve net CO₂ reduction at potentials that are more negative than −0.4 V, compared to other FDHs, displaying thus higher potential for practical applications in CO₂ fixation.⁴³

The active sites of Mo-FDHs are highly conserved, and their reaction mechanisms have been extensively studied using both quantum calculations and laboratory experiments.^{44–47} In a mechanistic study by Dong *et al.*, previously proposed mechanisms were evaluated employing combined QM/MM and large-scale QM calculations.⁴⁶ It was suggested that the CO₂ or formate molecule binds in the second coordination sphere of molybdenum, rather than binding directly to the metal, and the CO₂ reduction involves a hydride transfer *via* the sulfido group, yielding Mo⁶⁺=S from Mo⁴⁺-SH (Fig. 1a).⁴⁶ However, the full energy diagram was not reported in this study.

Siegbahn conducted a thorough theoretical study of the CO₂/formate interconversion catalyzed by Mo-FDH using the quantum chemical cluster approach.⁴⁷ In the starting point of the reduction direction before the binding of CO₂, the electronic structure of Mo(vi) was confirmed to be in the closed-shell singlet state. A (H⁺, e[−]) couple was then added before the CO₂ was bound, and the localized spin on Mo indicated a Mo(v) state. Subsequently, CO₂ bound to the sulfur of Cys196. This led to the spin on Mo being delocalized over the ligands, indicating a Mo(iv) state. Then, an electron was added and therefore a closed-shell Mo(iv) was obtained. During this step, the structure of the Mo complex hardly changed, while the spin distribution changed significantly. Finally, the calculations showed that the hydride transfer to CO₂ involves a proton from sulfide ligand and two electrons from Mo(iv) (Fig. 1b), giving the formate product and Mo(vi). Additionally, it was proposed that the

Table 1 Overview of the selected examples of enzymatic CO₂ conversion investigated using quantum chemical calculations

Enzyme	Reaction scheme ^a
Molybdenum-containing formate dehydrogenase (Mo-FDH)	$\text{CO}_2 + \text{H}^+ + 2\text{e}^- \xrightleftharpoons{\text{Mo}^{6+}, \text{pyranopterin}} \text{HCOO}^-$
Ni,Fe-containing carbon monoxide dehydrogenase (CODH)	$\text{CO}_2 + 2\text{H}^+ + 2\text{e}^- \xrightleftharpoons{\text{NiFe}_3\text{S}_4} \text{CO} + \text{H}_2\text{O}$
Artificial human carbonic anhydrase (hCAII)	$\text{CO}_2 + \text{H}_2 \xrightleftharpoons{\text{Rh(I)}} \text{HCOOH}$
Ribulose 1,5-bisphosphate carboxylase/oxygenase (RuBisCO)	
Crotonyl-CoA carboxylases/reductases (CCR)	$\text{H}_3\text{C}-\text{CH}=\text{CH}-\text{C}(=\text{O})-\text{S}-\text{CoA} + \text{NADPH} + \text{CO}_2 \longrightarrow \text{H}_3\text{C}-\text{CH}(\text{COO}^-)-\text{C}(=\text{O})-\text{S}-\text{CoA} + \text{NADP}^+$
Pyruvate carboxylase (PC)	$\text{CH}_3\text{C}(=\text{O})\text{COO}^- + \text{HCO}_3^- + \text{ATP} \xrightleftharpoons{\text{Mg}^{2+}, \text{biotin}} \text{CH}_3\text{C}(\text{O}^-)(\text{COO}^-)\text{COO}^- + \text{ADP} + \text{Pi}$
Pyruvate decarboxylase (PDC)	$\text{CH}_3\text{C}(=\text{O})\text{COO}^- + \text{CO}_2 \xrightleftharpoons{\text{Mg}^{2+}, \text{ThDP}} \text{CH}_3\text{C}(=\text{O})\text{COOH} + \text{CO}_2$
Pyruvate dehydrogenase (PDHc) E1 subunit	$\text{CO}_2 + \text{CH}_3\text{C}(=\text{O})\text{SH} \xrightleftharpoons{\text{Mg}^{2+}, \text{ThDP}} \text{CH}_3\text{C}(=\text{O})\text{COOH} + \text{CH}_3\text{C}(=\text{O})\text{S}^-$
Ferulic acid decarboxylase (AnFdc1)	$\text{C}_6\text{H}_5\text{CH}=\text{CHCOOH} + \text{CO}_2 \xrightleftharpoons{\text{K}^+, \text{Mn}^{2+}, \text{prFMN}} \text{C}_6\text{H}_5\text{CH}=\text{CHCOO}^- + \text{CO}_2$
3,4-Dihydroxybenzoic acid decarboxylases (AroY)	$\text{C}_6\text{H}_2(\text{OH})_2\text{COOH} + \text{CO}_2 \xrightleftharpoons{\text{K}^+, \text{Mn}^{2+}, \text{prFMN}} \text{C}_6\text{H}_2(\text{OH})_2\text{COO}^- + \text{CO}_2$
Pyrrole-2-carboxylic acid decarboxylase (HudA)	$\text{C}_4\text{H}_3\text{N} + \text{CO}_2 \xrightleftharpoons{\text{K}^+, \text{Mn}^{2+}, \text{prFMN}} \text{C}_4\text{H}_3\text{NCOOH} + \text{CO}_2$
Decarboxylases from amidohydrolase superfamily (AHS)	$\text{C}_6\text{H}_4(\text{OH})\text{R} + \text{CO}_2 \xrightleftharpoons{\text{Mn}^{2+}/\text{Mg}^{2+}} \text{C}_6\text{H}_4(\text{OH})\text{COOH} + \text{CO}_2$

^a Reactions of decarboxylases with carboxylation activity are depicted in the carboxylation direction.

barrier for the reduction of CO₂ is 19.0 kcal mol^{−1}, calculated from the resting state after the addition of the (H⁺, e[−]) couple to the hydride-transfer transition state (Fig. 1c). The added electron turns out to stabilize the binding of CO₂. The overall reduction reaction energy is +0.2 kcal mol^{−1}, which aligns well with the experimental result, indicating that this process is barrierless at a −0.4 V redox potential.⁴³ For the oxidation direction, the rate-limiting barrier is 16.9 kcal mol^{−1}, calculated from the resting point after CO₂ has left, followed by the release of (H⁺, e[−]), up to the hydride-transfer transition state. This study considered the entire oxidation cycle, rather than focusing solely on the chemical steps as in the previous study, thereby offering a more comprehensive view of CO₂ reduction catalyzed by Mo-FDH.

3.1.2 Ni, Fe-containing carbon monoxide dehydrogenase (CODH). The Ni, Fe-containing carbon monoxide dehydrogenase (CODH) catalyzes the reversible reduction of carbon dioxide (CO₂) to carbon monoxide (CO) and water *via* a tetranuclear

[NiFe₃S₄] cluster.^{27,48} A mechanistic study on this enzyme concluded that one sulfide bridge in the NiFe cluster should be protonated.⁴⁹ The NiFe cluster was assigned as Ni^{II}3Fe^{II}Fe^{III} based on the X-ray structure, CO₂ binding mode, and the doublet state of this cluster as indicated by EPR. The calculated energies of the model, refined by the addition of a proton and a water molecule, align well with experimental results. The reduction of CO₂ involves the cleavage of the C–O bond in CO₂ and two reductions, each involving a simultaneous addition of a proton and an electron (Scheme 1). During the reduction, nickel was proposed to occur as nickel(II) or nickel(I), but not nickel(0). The calculated energy of CO₂ replacing CO to start the new cycle suggests that this exchange occurs almost without energy cost. Additionally, the absence of CO in the resolved structure was suggested to be caused by the over-reduction of the cluster by X-rays during crystallization (PDB ID: 3B51).⁵⁰

The binding and release of CO₂ by CODH have also been investigated using quantum mechanics calculations.⁵¹ All possible

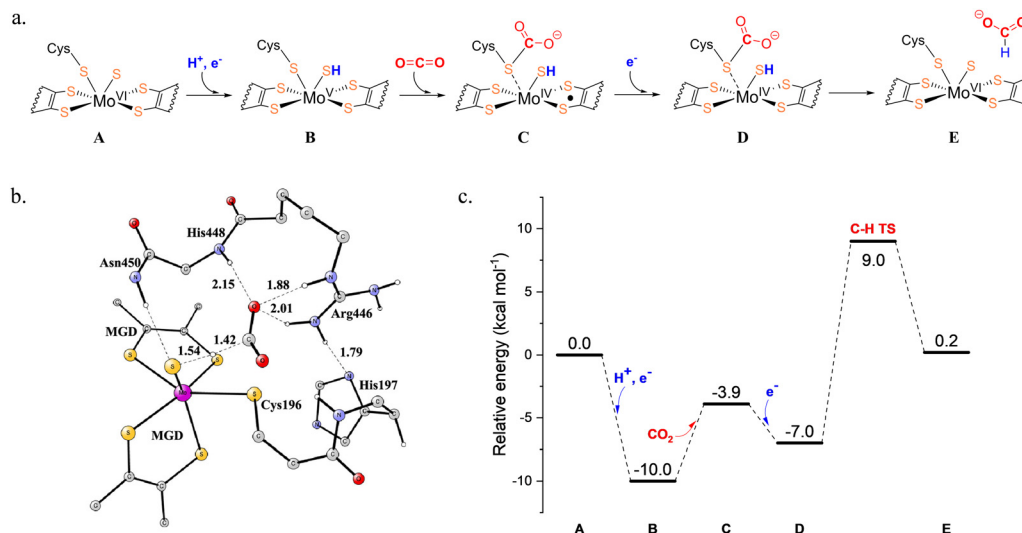
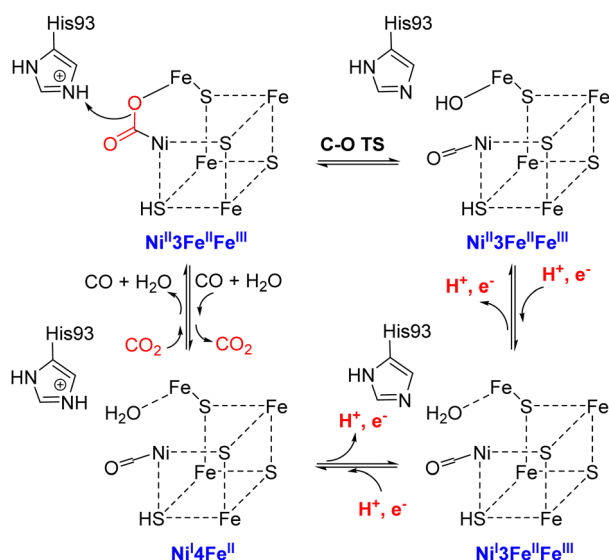


Fig. 1 (a) Proposed mechanism of the molybdenum-containing formate dehydrogenase (Mo-FDH) for CO₂ reduction. (b) The optimized transition state for the hydride transfer to CO₂ and (c) the calculated energy profile of CO₂ reduction. Reprinted with permission from ref. 47, Copyright 2022 American Chemical Society and CC-BY 4.0.



Scheme 1 Reaction mechanism of Ni, Fe-containing carbon monoxide dehydrogenase (CODH). Adapted with permission from ref. 49, Copyright 2019 American Chemical Society.

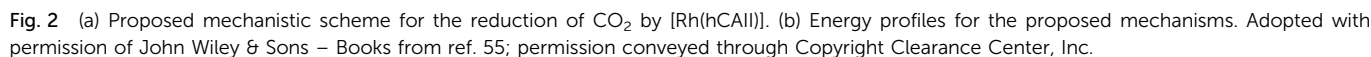
spin-coupling schemes for three reduced C-cluster states were investigated using a small model, and the states with low energies were finally used for a large model. The calculations of the CO₂ binding and release processes revealed that these two steps highly depend on the protonation state of His93, and the oxidation state of the asymmetrical [Ni-Fe-S] cluster. In detail, CO₂ binds to the dielectronic reduction state of C-cluster when His93 is doubly protonated, and is released when His93 is singly protonated and the C-cluster is in an oxidized state.

3.1.3 Artificial human carbonic anhydrase (hCAII). Human carbonic anhydrase II (hCAII) is a zinc-dependent enzyme that catalyzes the reversible conversion between CO₂ and

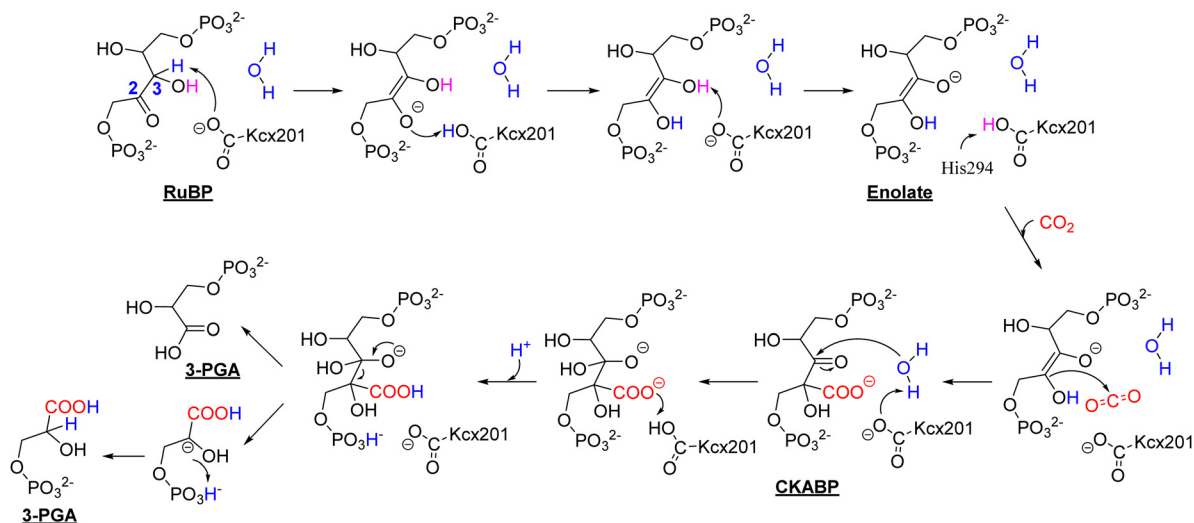
bicarbonate. Experimental studies have shown that, when the Zn(II) ion in the catalytic center of hCAII is replaced by Rh(I), the resulting artificial enzyme exhibits the capability to enable the stereoselective hydrogenation of olefins in the presence of H₂.^{52,53} Inspired by this reduction ability, the feasibility of this artificial enzyme in catalyzing the conversion of CO₂ to HCOOH through a direct hydrogenation reaction was evaluated using quantum chemical calculations.^{54,55}

Possible reaction mechanisms of hCAII-catalyzed conversion of CO₂ to HCOOH were proposed and evaluated.⁵⁴ The simulations using a cluster model of 50 atoms demonstrated that the initial step of the catalytic cycle, namely the activation of H₂ by Rh(I), occurs without an energy barrier. Subsequently, CO₂, acting as a Lewis base, coordinates to Rh(III) through one oxygen atom, then inserts into the Rh(III)-H bond, finally giving rise to the **1d** species. For the subsequent formation of formic acid, both reductive elimination and σ -bond metathesis mechanisms were considered, with the latter being preferred according to the calculated energies (Fig. 2a). The release of formic acid and the restoration of the catalytic cycle were also examined. It was found that the energy barrier for the release of the product is 12.8 kcal mol⁻¹ (**6b-6d**, Fig. 2b), which is the rate-determining step of the overall reaction.

In a later study, the role of explicit water molecules in the catalysis was examined using the ONIOM QM/QM' (B3LYP/PM6) method, defining a high-level layer QM region and a low-level layer QM' region.⁵⁵ The energy profile obtained from the QM/QM' simulations follows the same trend as the one from the QM cluster calculations (Fig. 2b). Consistent with the previous cluster calculations, the catalytic process was shown to follow the σ -bond metathesis mechanism, and the rate-limiting step is the release of formic acid. An interesting finding was that for the product release the energy barrier from the QM/QM' investigation, obtained without the presence of water



The conserved lysine (Lys175) was believed to provide the necessary proton in stereospecific protonation of C2.^{56,57} To investigate the feasibility of this assumption, Cummins *et al.* used DFT(B3LYP) and *ab initio* molecular dynamics simulations with umbrella sampling to determine the mechanisms of the C2–C3 bond scission and the stereospecific protonation.⁶² It was concluded that the Lys175 is not the direct proton donor to C2 during the product formation according to the free energy change. Instead, a proton from the P1 phosphate of RuBP is plausibly transported *via* a water molecule to C2, following the Grothuss mechanism (Scheme 2).



Scheme 2 Reaction mechanism of RuBisCO: the enolization of RuBP, the carboxylation and hydration steps, the scission of C2–C3, and the stereospecific protonation of C2.^{58–60}

According to these calculations, the RuBisCO carboxylation process starts with the abstraction of a proton from the carbon C3 of ribulose-1,5-bisphosphate (RuBP) by the carbamylated Lys201, forming an enolate intermediate. Then, this intermediate undergoes a proton rearrangement, and subsequently performs a nucleophilic attack on CO₂, adding CO₂ to its C2 atom to form 2-carboxy-3-keto-D-arabinitol-1,5-bisphosphate (CKABP). The reaction further proceeds with the hydration of CKABP and the C2–C3 bond cleavage concurrent with C2 stereospecific protonation, ultimately producing two molecules of 3-phospho-D-glyceric acid (3-PGA) (Scheme 2). These mechanistic insights can be helpful in investigating the catalytic limitation on the reactivity of RuBisCO, whose rate constants are much lower than those of classical central metabolic enzymes.

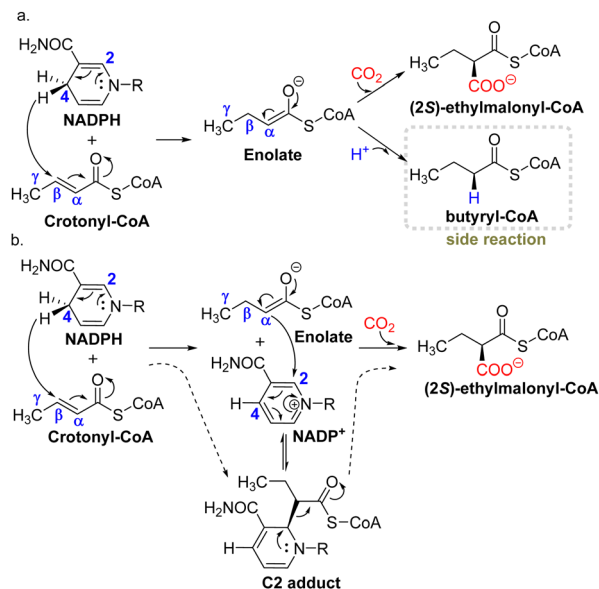
3.2.2 Crotonyl-CoA carboxylases/reductase (CCR). Without any side reactivity with O₂, the NADPH-dependent crotonyl-CoA carboxylase/reductase (CCR), which is a member of the enoyl-CoA carboxylases/reductases (ECRs) family, can exclusively fix CO₂ and exceed the catalytic efficiency of RuBisCO by more than tenfold.^{63–65}

The catalytic reaction of CCR follows the general mechanism of ECR enzymes. This process involves a hydride transfer from NADPH to the β-carbon of crotonyl-CoA, forming an enolate intermediate. The enolate then undergoes a nucleophilic attack on CO₂ to form (2*S*)-ethylmalonyl-CoA. Stoffel *et al.* revealed the function of four amino acids that contribute significantly to the high efficiency of the carboxylation activity of CCR through a combination of QM/MM approach and experimental assays.⁶³ It was demonstrated that Asn81 is crucial for positioning the CO₂ molecule correctly through its carboxamide NH₂ interacting with CO₂. His365 and Glu171 jointly coordinate an ordered water molecule, forming a water network that helps control the positioning of CO₂. Meanwhile, Phe170 shields the active site from water, preventing the enolate from being protonated into the reduction side product butyryl-CoA.

Interestingly, a C2-ene adduct was experimentally detected in CCR reaction in the absence of CO₂.^{66,67} Similar adduct has also been experimentally characterized in mammalian fatty acid synthase, which was then computationally confirmed.⁶⁸ To reveal the role of this adduct intermediate in the catalysis, Recabarren *et al.* have evaluated two possible reaction pathways for CCR, including a direct mechanism and a C2 mechanism (Scheme 3).⁶⁹ Instead of directly transferring a hydride from N4 of NADPH to the Cβ of the substrate, forming an enolate that reacts with CO₂ in the direct mechanism, the C2 mechanism involves a Michael addition between the C2 of NADPH and the Cα of the substrate to form a C2-ene adduct. This adduct is followed by subsequent carboxylation (Scheme 3). According to the calculated free energy landscapes, this adduct can react with CO₂ with a similar activation barrier to the direct reaction pathway. It was thus proposed that the formation of the C2-ene adduct is a strategy employed by this enzyme to store enolate intermediate when CO₂ is absent from the active site. It becomes reactive when CO₂ is available, thereby improving CO₂-fixing efficiency. This adduct also acts as a stable intermediate to avoid the reduction side reaction.

In conclusion, factors that determine the high efficiency of CO₂ fixation by CCR may include: the unidirectional formation of the enolate; key residues that control the position of CO₂ being attacked; suppression of competing (re)protonation reactions by shielding the active pocket from water; and the formation of a C2-ene adduct, which restores the enolate intermediate and prevents side reduction reactions.

3.2.3 Pyruvate carboxylase (PC). Pyruvate carboxylase (PC, EC 6.4.1.1), a biotin-dependent carboxylase, catalyzes the carboxylation of pyruvate to oxaloacetate, a process regulated by acetyl-CoA. This enzyme functions through the collaborative actions of three key domains: the biotin carboxyl carrier protein (BCCP) domain, the biotin carboxylase (BC) domain, and the carboxyl transferase (CT) domain.^{70,71} In this catalytic process, biotin is covalently attached to a lysine residue within the BCCP



Scheme 3 Reaction mechanism of crotonyl-CoA carboxylases/reductases: direct mechanism (a) and C2 mechanism (b). Adapted with permission from ref. 69, Copyright 2023 American Chemical Society.

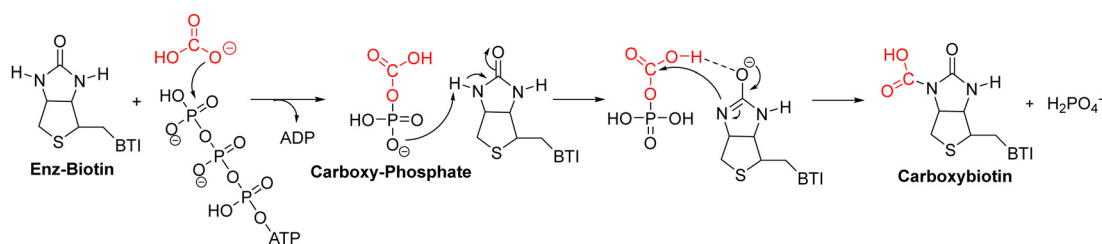
domain. As this domain moves, it translocates the carboxylated biotin (carboxybiotin) between the BC domain and the CT domain, enabling the transfer of carboxyl groups. Two reactions sequentially take place within the BC and CT domains separately. For the first reaction, biotin accepts the carboxyl group from bicarbonate in the BC domain, requiring ATP in complex with Mg (Mg-ATP) as the cofactor. Subsequently, the carboxyl group of carboxybiotin is transferred to pyruvate to form oxaloacetate in CT domain. The catalytic mechanism and the energetic characteristics of pyruvate carboxylase have been sketched through quantum chemical studies.

Yoshizawa *et al.* investigated the plausible reaction mechanisms accomplished in the BC domain using quantum calculations.⁷² According to the detailed analysis of potential energy surfaces, a two-step reaction mechanism was proposed (Scheme 4). Initially, bicarbonate is phosphorylated by attacking on the γ -phosphate group of ATP to form a short-lived carboxy-phosphate (CP) intermediate. During this process, the Mg^{2+} ion promotes the formation of CP by alleviating the electrostatic repulsion between bicarbonate and ATP. In the second step, CP serves as a general acid-base catalyst, facilitating the enolization of biotin and subsequently

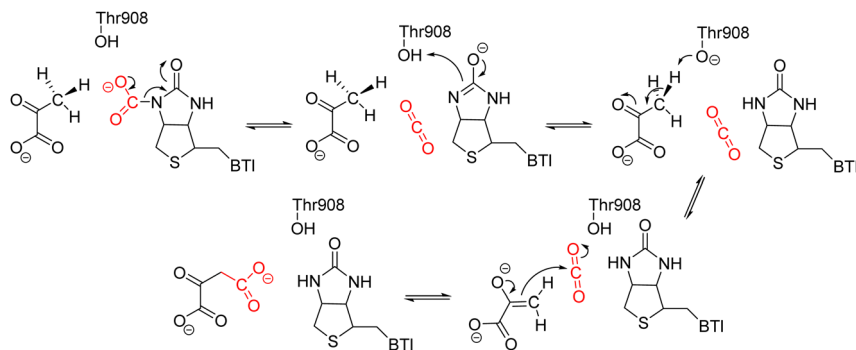
generating a bent CO_2 to react with enolic biotin. This step can be interpreted as the nucleophilic attack by the ureido nitrogen of biotin on the carboxyl group of CP to form carboxybiotin. The rate-limiting step is the formation of CP in the first step.

Sheng *et al.* conducted a successive study on the catalytic mechanism of the CT domain in pyruvate carboxylase from *Staphylococcus aureus*,^{73,74} exploring how posttranslational modifications (PTMs) affect enzyme activity. At first, the elementary steps in the catalytic cycle occurring within the CT domain were investigated.⁷³ Four steps were proposed, including the decarboxylation of carboxybiotin to generate an enol-biotin intermediate and a free CO_2 molecule, the proton transfer from T908 to enol-biotin, the proton transfer from pyruvate to T908, and the nucleophilic attack by enol-pyruvate at the generated CO_2 (Scheme 5). Subsequently, Sheng *et al.* investigated the roles of the carboxylated lysine and a water molecule coordinated to Zn^{2+} in the catalysis.⁷⁴ While following the same reaction mechanism, the new model, which includes the entire modified lysine and a water molecule in the QM region of QM/MM calculations, showed significantly lower energy barriers of all the elementary steps compared to the previous calculations, which was based on a model without the inclusion of the carboxylate group on the modified lysine and the water molecule. In particular, the energy barrier of the overall reaction using the new model is approximately 14 kcal mol^{-1} , which is much lower than that in the previous study. The Zn-coordinated water molecule was explained to enhance the stabilization of the transition state during the carboxylation of enol-pyruvate. Meanwhile, the modified lysine was proposed to facilitate the reaction by positioning and orienting the substrate and cofactor in a proper position during the catalysis, and also by regulating the charge of zinc cation.

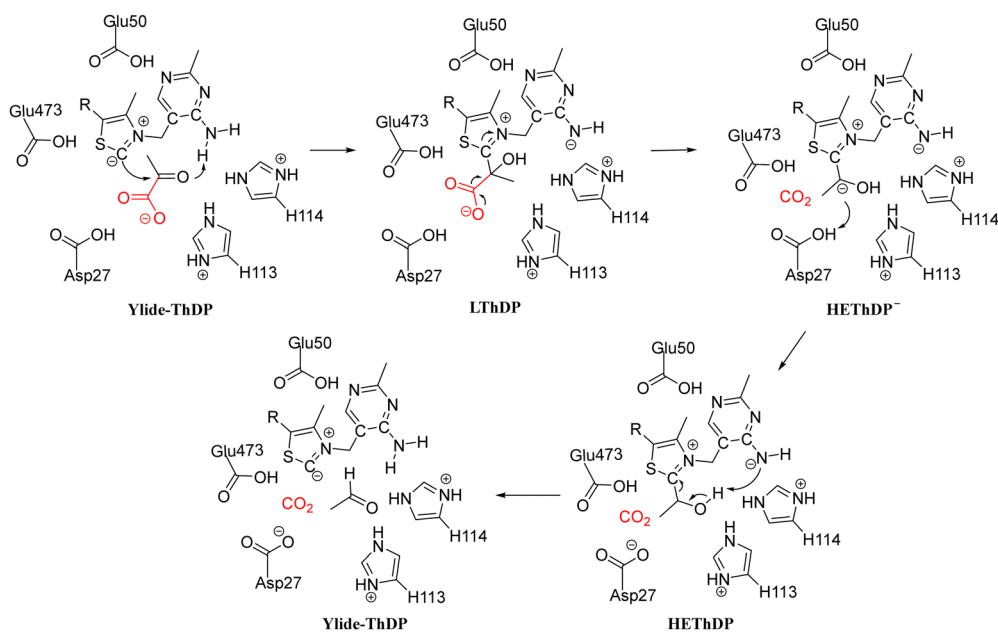
3.2.4 Pyruvate decarboxylase (PDC). Pyruvate decarboxylase (PDC) is a thiamine diphosphate (ThDP)-dependent keto acid decarboxylase that catalyzes the nonoxidative decarboxylation of pyruvate to acetaldehyde. Interestingly, the PDC from *Saccharomyces cerevisiae* (ScPDC) has been demonstrated to catalyze the carboxylation of acetaldehyde into pyruvic acid under conditions of high sodium bicarbonate buffer (optimum at 500 mM NaHCO_3 - Na_2CO_3 buffer) and strongly alkaline conditions (optimum at pH 11).⁷⁵ For the decarboxylation direction, the entire catalytic cycle of PDC includes the formation of a ylide by the deprotonation of C2 of thiazolium, the covalent addition of pyruvate to the ylide to form the lactylthiamin diphosphate (LThDP) intermediate, the decarboxylation of LThDP,



Scheme 4 Reaction mechanism of the biotin carboxylation in the BC domain of pyruvate carboxylase.⁷²



Scheme 5 Reaction mechanism of the pyruvate carboxylation in the CT domain of pyruvate carboxylase.⁷³



Scheme 6 Decarboxylation mechanism of pyruvate decarboxylase from *Zymomonas mobilis* (ZmPDC).⁷⁸

the protonation of the α -carbanion to form 2-(1-hydroxyethyl)-thiamin diphosphate (HEThDP), and finally the release of acetaldehyde (Scheme 6).^{76,77}

Using density functional theory calculations, Wang *et al.* investigated the reaction mechanism of *ScPDC*.⁷⁹ According to the calculations, a proton relay among Glu51 (equivalent to Glu50 in Scheme 6), N1' and the 4'-amino group of ThDP facilitates the deprotonation of C2, forming the ylide intermediate. This process also affects the basicity of the 4'-amino group, which deprotonates the α -hydroxyl of HEThDP, thereby accelerating the release of acetaldehyde. Glu477 and Asp28 facilitate the decarboxylation of LThDP by forming strong hydrogen bonds with its carboxylate group, making the reaction endothermic. Subsequently, a proton is transferred from His115 *via* Asp28 to the α -carbanion in a concerted manner to form HEThDP. However, the cluster model used in this study was small, and the roles of other key active site residues were not discussed.

A QM/MM theoretical study of PDC from *Zymomonas mobilis* (ZmPDC) was carried out to determine the protonation states

and the roles of active site catalytic residues.⁷⁸ Three models with varying protonation states of the key active site residues were considered for different mechanisms. In the most plausible mechanism, the ionizable residues in the active site, including Asp27, Glu50, His113, His114 and Glu473, were proposed to be in their protonated states. Asp27 was suggested to provide a proton to the HEThDP carbanion/enamine intermediate, differing from the proposed mechanism of *ScPDC* discussed above.⁷⁹ His113 and His114 interact with the substrate and intermediates during the reaction process. According to the energy profile, the LThDP and HEThDP intermediates have lower energies than all the other intermediates, including the enzyme-substrate and enzyme-product complexes. This is consistent with the experimentally observed accumulation of these two compounds.⁸⁰ The decarboxylation of this enzyme was calculated to be the rate-determining step.

These calculations have confirmed the protonation states of the ionizable residues in the active site of PDC, and also the roles of these residues in the catalysis.

3.2.5 Pyruvate dehydrogenase (PDHc). As one subunit of the pyruvate dehydrogenase multienzyme complex (PDHc), the ThDP-dependent E1 subunit catalyzes the reversible decarboxylation of pyruvate and the transfer of the resulting acetyl group to the lipoyl group of the E2 subunit. The catalytic mechanism for the PDHc E1 subunit has been previously proposed.^{76,77} It starts with the activation of ThDP, generating the ThDP ylide. Then, the C2 carbanion of ThDP ylide attacks the C2 α of pyruvate, forming the LThDP intermediate. Subsequently, the cleavage of the C2 α –C2 β bond takes place, giving the HETHP intermediate and releasing CO₂. The C2 α of HETHP then attacks the S6 of the lipoyl group, concurrent with the protonation of S8. Finally, the C2–C2 α bond breaks, resulting in the formation of the acetyl–lipoyl adduct and the regeneration of ThDP ylide.

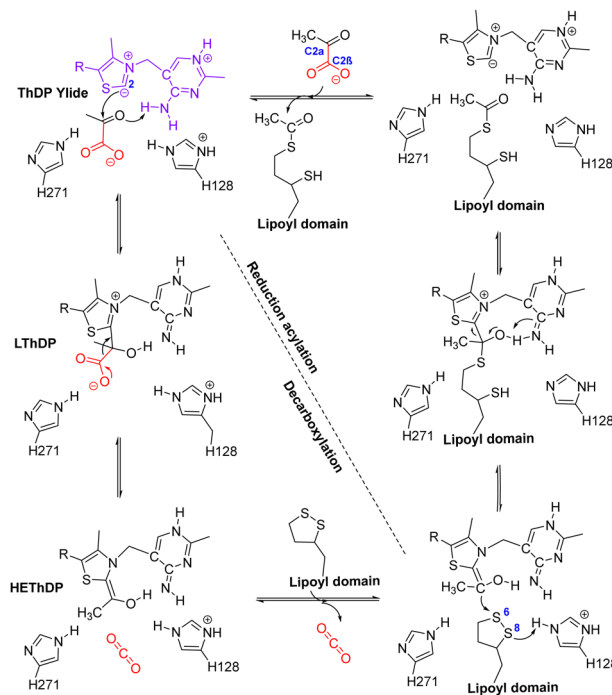
The quantum chemical studies on the PDHc E1 subunit provided support to the reaction mechanism discussed above. In the mechanistic study on the PDHc E1 from *Escherichia coli* using the cluster approach,⁸¹ the roles of several active site residues were confirmed. In particular, Glu571 and Glu522 play essential roles in ThDP ylide formation, pyruvate addition, acetaldehyde release, and the positioning of the carboxyl of pyruvate, while His640 and a water molecule are involved in a concerted proton transfer processes to protonate the enamine.

In a subsequent study, also using the cluster approach, Sheng and Liu investigated four possible pathways for the PDHc E1 from *Bacillus stearothermophilus*. In these pathways, the 4'-amino of the ylide, His271 (a residue not considered in the study on the PDHc E1 from *E. coli* discussed above), and His128 (equivalent to His640 in the PDHc E1 from *E. coli*) were proposed to act as the general acid/base in various combinations in the processes of decarboxylation and reductive acylation.⁸² It was found that the most favorable pathway involved the 4'-amino group of the ylide and protonated His128 acting as acid/base groups, with His271 deprotonated (Scheme 7). The C–C bond cleavage that releases CO₂ was calculated to be the rate-limiting step.

The computational studies on PDC and PDHc E1 showed that the decarboxylation reactions follow a similar mechanism. However, the energetics, protonation states, and roles of active site residues can vary between different enzymes. This highlights the importance of calculating the entire mechanisms in detail and the corresponding energy profiles to gain accurate insights into the reaction.

3.3 Fixation of carbon dioxide to form aromatic acids

3.3.1 Enzymes belonging to UbiD enzyme family. The enzymes from UbiD-family are reversible decarboxylases believed to be involved in the degradation of aromatic compounds. Their catalytic activities require the prFMN co-factor, which is formed by the prenylation of FMN catalyzed by UbiX or other functionally identical enzymes. Various aromatic compounds, such as styrene derivatives, phenol derivatives, and heteroaromatic substrates, can be carboxylated by the UbiD enzymes, though only sluggish to moderate conversions have been observed.^{3,7,24} Distinct mechanisms have been proposed for different members of this family.

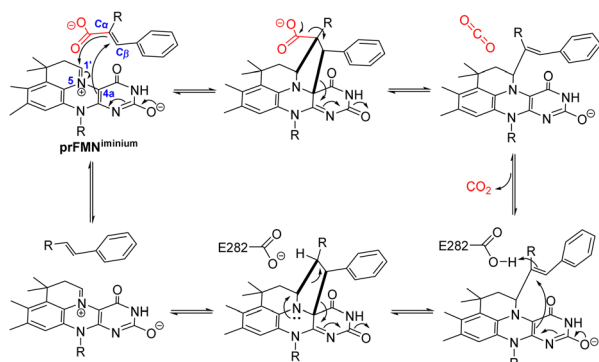


Scheme 7 The reaction mechanism of PDHc E1 subunit. Adapted with permission from ref. 82, Copyright 2013 American Chemical Society.

A 1,3-dipolar cycloaddition mechanism was first proposed for the UbiD enzymes on the basis of the study of the ferulic acid decarboxylase (Fdc1) from *Aspergillus niger*.⁸³ Given the azomethine ylide character of prFMN and the alkene dipolarophiles of α,β -unsaturated carbonyl group, the 1,3-cycloaddition reaction was suggested to occur between the co-factor and the substrate. This leads to the formation of the [3+2] covalent substrate–prFMN pyrrolidine adduct, which involves the substrate C α binding to the C1' of prFMN and the substrate C β binding to the C4 α of prFMN concurrently. The reaction proceeds with decarboxylation coupled with the breaking of the C β –C4 α bond, the protonation of phenylpyruvate adduct by E282, the formation of the second cycloadduct species, and then the cycloelimination to form final styrene product (Scheme 8).

The decarboxylation mechanism of Fdc1 was investigated by using both the cluster approach⁸⁴ and the QM/MM method.⁸⁵ According to the calculations, the Fdc1 reaction indeed follows the 1,3-cycloaddition mechanism and Glu282 was confirmed to play important role in catalysis being as a general acid group. In the former study, using α -methylcinnamic acid as the representative substrate, the decomposition of the second pyrrolidine intermediate was calculated to be the rate-limiting step.⁸⁴ While in the latter one, the calculations showed that for the decarboxylation of cinnamic acid the protonation step is rate-limiting.⁸⁵

Quantum chemical cluster calculations revealed a distinct reaction mechanism from 1,3-cycloaddition for 3,4-dihydroxybenzoic acid decarboxylases (AroY), which also belongs to the UbiD enzyme family.⁸⁶ The reversible reaction of aromatic



Scheme 8 Proposed 1,3-cycloaddition mechanism for ferulic acid decarboxylase (Fdc1).⁸³

catechol substrates was suggested to initiate by a nucleophilic attack of the substrate C α to the C1' of prFMN. Instead of forming 1,3-dipolar cycloaddition intermediate, a quinoid binding to the cofactor adduct *via* a single covalent bond is formed (Scheme 9). The overall barrier of the reaction is 19.9 kcal mol⁻¹, according to the calculated energy profile. Moreover, the calculations confirmed that a cycloadduct intermediate is unlikely to form in the case of AroY, as the energy of such an intermediate is as high as 37 kcal mol⁻¹. Interestingly, by using also the cluster approach, a similar intermediate was later found in the cinnamic acid cycloaddition by Fdc1.²⁹ However, such intermediates were not located in the reaction of phenylpropionic or crotonic acid by Fdc1.

Theoretical analysis of UbiD enzyme pyrrole-2-carboxylic acid decarboxylase (HudA),⁸⁷ a virulence attenuation factor from *Pseudomonas aeruginosa*, suggests a nucleophilic attack of C1' at the pyrrole C2 position of heteroaromatic substrates. It follows the electrophilic aromatic substitution mechanism, with a Wheland-type intermediate formed. A recent mechanistic study of HudA and two other prFMN-dependent decarboxylases (HmfF and InD) using heteroaromatic substrates suggested that the decarboxylation adopts a nucleophilic attack mechanism while the carboxylation follows a 1,3-dipolar

cycloaddition mechanism.⁸⁸ For the carboxylation of furoic acid, pyrrole, and indole, it was proposed that they favor the formation of a 1,3-dipolar cycloadduct in the initial step because of the lack of a carboxyl group on one side. The “absent” carboxyl group was proposed to hinder electron donation to the cofactor, making it unfavourable to form the Wheland-type intermediate.

3.3.2 Enzymes belonging to amidohydrolase superfamily (AHS). The decarboxylases belonging to the amidohydrolase superfamily (AHS) are non-redox metal-dependent enzymes that catalyze the reversible decarboxylation of aromatic acids. The quantum chemical cluster chemical calculations were performed on a number of AHS decarboxylases, including 5-carboxyvanillate decarboxylase (LigW),⁸⁹ γ -resorcyate decarboxylase (γ -RSD),⁹⁰ 2,3-dihydroxybenzoic acid decarboxylase (2,3-DHBD),⁹¹ salicylic acid decarboxylase (SDC),⁹² and isorotinate decarboxylase (IDCase).⁹³ Detailed information of the mechanisms can facilitate the improvement of their further utilization in converting CO₂ into value-added aromatic chemicals.

Decarboxylation mechanisms of five members of this family have been theoretically investigated, which mainly consist of two key steps: the protonation of the carbon of the substrate and the cleavage of the C–C bond.⁸⁹ In combination with crystallographic study, the substrates were confirmed to bind to the metals in a bidentate mode. Upon the binding, the hydroxyl groups lose their protons. The initial reaction products of the enzymatic decarboxylation were confirmed to be CO₂ rather than the bicarbonate proposed in previous studies,⁹⁴ because a high-energy hydrated carboxylate intermediate is involved in the reaction pathway to form bicarbonate.

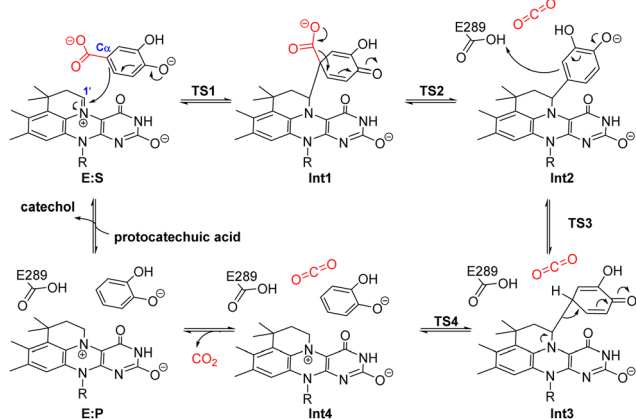
Among these decarboxylases being investigated, the rate-limiting steps are not the same for each one (Table 2). The proton transfer is the rate-limiting step for LigW,⁸⁹ γ -RSD,⁹⁰ 2,3-DHBD⁹¹ and SDC,⁹² while the C–C bond cleavage is rate-limiting for IDCase.⁹³ Initially, the divalent metal ion dependency of the AHS superfamily was believed to be Zn²⁺.^{94–96} However, according to the five research studies carried out using quantum chemical calculations in combination with experimental validation (Table 2), 2,3-DHBD is active with both Mn²⁺ and Mg²⁺,⁹¹ LigW,⁸⁹ IDCase⁹³ and γ -RSD⁹⁰ prefer to depend on Mn²⁺, while SDC is Mg²⁺-dependent.⁹²

These calculations on AHS decarboxylases show that the quantum chemical modeling is a powerful tool in investigating reaction mechanism, and can also be used in clarifying the metal identity and other properties of the enzymes.

4. Perspective

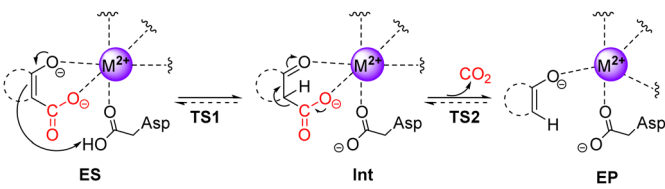
Our perspective closes with a viewpoint on the potential of combining machine learning (ML) and quantum descriptions in investigating enzymatic CO₂ conversion.

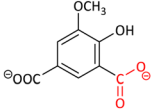
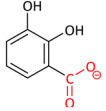
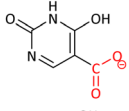
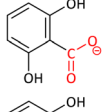
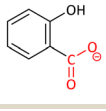
CO₂-converting enzymes generally catalyze at rather low k_{cat} values, ranging from 10 s⁻¹ (average for RubisCO)⁹⁷ to 100 s⁻¹ (ECRs).^{9,98} Therefore, there is an urgent need to enhance the



Scheme 9 Proposed nucleophilic attack mechanism for 3,4-dihydroxybenzoic acid decarboxylases (AroY).⁸⁶

Table 2 A general mechanism for AHS decarboxylases, and calculated energies (kcal mol⁻¹) of the intermediates and transition states of each enzyme



Enzyme	Substrate	ES	TS1	Int	TS2	EP	M ²⁺
LigW ⁸⁹		0.0	16.8	9.9	14.4	-4.0	Mn ²⁺
2,3-DHBD ⁹¹		0.0	15.8	8.8	11.7	-3.1	Mn ²⁺ /Mg ²⁺
IDCase ⁹³		0.0	14.1	9.2	20.7	2.8	Mn ²⁺
γ-RSD ⁹⁰		0.0	14.8	8.1	11.4	3.3	Mn ²⁺
SDC ⁹²		0.0	16.6	9.4	15.8	0.8	Mg ²⁺

performance of existing carbon-fixation enzymes or to design new-to-nature CO₂-converting enzymes. Recently, ML has shown significant promise in accelerating enzyme engineering. This is due to substantial improvements in protein representations and deep learning models, which can incorporate numerous parameters as needed.

Using data obtained from quantum chemistry calculations to train ML models is an efficient way to predict the electronic properties of reactants and the thermodynamic properties of reactions, thereby to predict the reactivity and selectivity of CO₂ conversion reactions. ML has successfully been applied to study the chemisorption, binding, activation, and hydrogenation of CO₂, in combination with quantum chemical computations in chemical systems.^{99–102} For example, Zheng *et al.* reported quantum-informed machine-learning force fields (QMLFFs) that can predict the binding free energy landscape of CO₂ chemisorption in Mg-MOF-74.⁹⁹ Chen *et al.* developed neural network (NN) ML models based on DFT calculations to discover catalysts capable of circumventing the scaling relation for CO₂ reduction reactions.¹⁰³ Zhu *et al.* proposed ML models based on DFT calculations to predict the free energy changes along the reaction pathway and the product probability for methanol, methane and formaldehyde in CO₂ reduction reaction catalyzed by metal-zeolites.¹⁰⁴

With the ability to analyze and learn from large size of data, ML methods can dramatically expand the applicability range of

conventional QM-based simulations to biological systems. We believe that combining quantum descriptions with machine learning can be valuable in addressing the reaction mechanisms of specific CO₂-converting enzymes and can contribute to the discovery of novel enzyme activities, the rational design of enzymes, or the improvement of efficiency in biocatalytic CO₂ conversion.

Author contributions

Baoyan Liu: investigation, writing – original draft, visualization; Beibei Lin: visualization; Hao Su: visualization, writing – original draft; Xiang Sheng: conceptualization, supervision, visualization, writing – review & editing.

Data availability

No primary research results, software or code have been included and no new data were generated or analyzed as part of this perspective.

Conflicts of interest

There are no conflicts to declare.

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