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Mechanistic studies of NO_x reduction reactions involving copper complexes: encouragement of DFT calculations

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The reduction of nitrogen oxides (NO_x), which is mainly mediated by metalloenzymes and metal complexes, is a critical process in the nitrogen cycle and environmental remediation. This Frontier article highlights the importance of density functional theory (DFT) calculations to gain mechanistic insights into nitrite (NO₂[−]) and nitric oxide (NO) reduction reactions facilitated by copper complexes by focusing on two key processes: the reduction of NO₂[−] to NO by a monocopper complex, with special emphasis on the concerted proton–electron transfer, and the reduction of NO to N₂O by a dicopper complex, which involves N–N bond formation, N₂O₂ isomerization, and N–O bond cleavage. These findings underscore the utility of DFT calculations in unraveling complicated reaction mechanisms and offer a foundation for future research aimed at improving the reactivity of transition metal complexes in NO_x reduction reactions.

Introduction

In the denitrification process of the nitrogen cycle, nitrate (NO₃[−]) is sequentially reduced to nitrogen gas (N₂) according to the process NO₃[−] → NO₂[−] → NO → N₂O → N₂.^{1–3} Among these steps, the reduction of nitrite (NO₂[−]) and nitric oxide (NO) to produce NO and nitrous oxide (N₂O), respectively, is catalyzed by metalloenzymes.⁴ Inspired by these enzymes, diverse transition metal complexes that mediate NO₂[−] and NO reductions have been reported. The electronic structure of such reactions, particularly those involving nitrosyl complexes (M–NO), is complicated due to the non-innocent nature of the NO ligand.^{5,6} Experimentally, the oxidation state of the NO ligand is estimated from the M–N–O angle and N–O stretching vibration values. In contrast, density functional theory (DFT) calculations provide direct information about the electron densities in molecules and atoms.^{7–12} Additionally, DFT calculations allow elucidating complicated reaction mechanisms such as concerted proton–electron transfer (CPET).^{13–18} Therefore, DFT calculations are highly recommended for the analysis of the reaction mechanisms of NO₂[−]/NO reduction catalyzed by transition metal complexes. Here, we discuss two DFT studies on the copper-mediated reductions of NO₂[−] and NO. The former focuses on the CPET in a copper-mediated NO₂[−] reduction and quantitatively estimates the asynchronicity between proton transfer (PT) and electron transfer (ET) during CPET. The latter explores the complicated

electronic state changes in a NO reduction reaction mediated by a dicopper complex.

Reduction of NO₂[−] to NO mediated by a monocopper complex using a phenol derivative

The reduction of NO₂[−] to NO in the denitrification process is catalyzed by nitrite reductases (NiRs), including copper-containing and heme-containing NiRs.^{2–4,19–22} In addition, the NO₂[−] reduction complements a physiological synthesis of NO, which functions as a signal transducer in immune response, vasodilation, and neurotransmission.^{23,24} Enzymes such as Mo-containing NiRs, xanthine oxidase, aldehyde oxidase, sulfite oxidase, and globins also perform the NO₂[−] reduction as a NO reservoir.^{21,24–27}

Crystallographic studies have identified the mononuclear type 2 Cu catalytic active site in CuNiRs, where three histidines bind to the Cu center.^{19–22,28–30} This has inspired researchers to investigate the Cu-complex-mediated NO₂[−] reduction. Woollard-Shore *et al.*³¹ synthesized, characterized and evaluated the catalytic properties of four Cu(II) complexes as potential agents for NO₂ reduction. The crystallographic study revealed three distinct coordination modes: η¹-NO₂, η¹-ONO, and η²-ONO. Chandra Maji *et al.*³² conducted an electrochemical study and suggested that the coordination mode is isomerized from η²-ONO to η¹-NO₂ during NO₂[−] reduction by a Cu(II) complex. Kundu and coworkers have expanded the study of the phenol-mediated NO₂[−] reduction using Cu(II)

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complexes.^{33–35} Among them, a reminiscent model of the red copper site in nitrosocyanin exhibited the NO-oxidase activity as well as the NiR activity.³⁵ Cioncoloni *et al.*³⁶ showed, based on kinetic and DFT studies, the involvement of CPET in the electrocatalytic NO₂[−] reduction by two types of Cu complexes. However, differences of reactivity depending on the coordination mode of NO₂[−] and detailed mechanism of CPET in the Cu-mediated NO₂[−] reduction remain unclear.

This study aims to characterize the mechanism of the NO₂[−] reduction by Cu(II) complexes, as reported by Kundu *et al.*³³ They utilized a Cu(II) complex with a tripodal heteroditopic cryptand ligand (L) and phenol. The reaction of a phenol derivative (ArOH) with a Cu(II) nitrite complex ([Cu^{II}(L)(ONO)]⁺) forms a hydroxyl Cu(II) complex ([Cu^{II}(L)(OH)]⁺), NO, and a biphenol derivative, which was presumed to be formed *via* coupling with the corresponding phenoxyl radical (ArO[•]). Mechanistic studies indicated that CPET occurred during the NO₂[−] reduction. We recently reported a DFT study on the reaction mechanism of this NO₂[−] reduction.³⁷ DFT calculations were used to elucidate the mechanisms of the NO₂[−] reduction in O- and N-coordination modes, highlighting the differences in the CPET process between these modes.

The X-ray crystallography study performed by Kundu *et al.* shows that NO₂[−] is O-coordinated to Cu.³³ However, DFT calculations suggest the formation of an N-coordinated Cu(II)-nitro complex (IC_N) in addition to an O-coordinated Cu(II)-nitrite complex ([CuONO]⁺, IC_O). Therefore, two reaction pathways from IC_O (nitrite pathway) and IC_N (nitro pathway) were examined. Both reaction pathways finally lead to the same product. In the nitrite pathway (Fig. 1), the reaction of IC_O with ArOH forms the reactant complex RC_O. The simultaneous H atom migration and O–N bond cleavage through transition state TS_O produces a hydroxyl Cu(II) complex, ArO[•], and NO. The overall reaction in the nitrite pathway is endothermic by 17.8 kcal mol^{−1} with an activation energy of 25.2 kcal mol^{−1} for the H atom migration and O–N bond cleavage. Meanwhile, the nitro pathway

involves three steps: H atom migration, N–O bond cleavage, and NO desorption, as shown in Fig. 2. The reaction begins with the association of IC_N with ArOH to form the reactant complex RC_N. Then, H atom migration from ArOH to NO₂[−] occurs, leading to the HONO intermediate [Cu(HONO)]⁺ and ArO[•] (IM1_N). N1–O2 bond cleavage forms the nitrosyl hydroxyl complex [Cu(NO)(OH)]⁺ (IM2_N) having two spin states corresponding to Cu^INO⁺ and Cu^{II}NO⁰. The final step involves NO ligand dissociation. The rate-determining step, *i.e.*, H atom migration, has an activation energy of 22.8 kcal mol^{−1}. Thus, DFT calculations suggest that the nitrite pathway (ΔG_{TS_O}[‡] = 25.2) and the nitro pathway (ΔG_{TS1_N}[‡] = 22.8) are comparable.

The above mentioned approach allowed tracing changes on the potential energy surface by connecting discrete points of stable structures and transition states; however, further insights can be obtained by using intrinsic reaction coordinate (IRC) calculations to trace continuous changes in electronic energies, geometries, and electronic configurations, particularly regarding the CPET mechanism. To better understand the CPET-derived NO₂[−] reduction, we carefully examined the electronic state changes associated with the structural changes derived from the IRC analysis, revealing the PT and ET behavior along the reaction coordinate. Over the past decade, a mechanism involving the concerted but asynchronous movement of a proton and an electron (*i.e.*, asynchronous CPET) has been proposed.^{38–43} Given the existence of multiple CPET mechanisms, a detailed analysis requires information on the sequential changes in charge and structure along the reaction path. To this aim, IRC calculations, which can determine the reaction pathway itself, are highly valu-

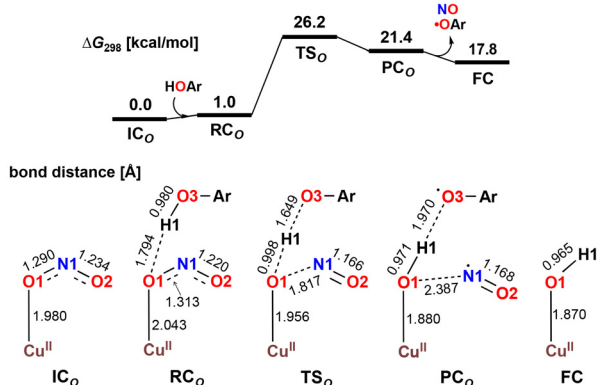


Fig. 1 Free energy profile and optimized structures for the reduction of NO₂[−] to NO in the nitrite pathway computed at the B3LYP-D3/6-31G** (Wachters-Hay for Cu) level of theory with the solvent effect of acetonitrile using the polarizable continuum model. Reproduced from ref. 37 with permission from the American Chemical Society, copyright 2023.



Fig. 2 Free energy profile (a) and optimized structures (b) for the reduction of NO₂[−] to NO at the B3LYP-D3/6-31G** (Wachters-Hay for Cu) level of theory with the solvent effect of acetonitrile using the polarizable continuum model. Free energy and bond distance are represented in kcal mol^{−1} and Å, respectively. Reproduced from ref. 37 with permission from the American Chemical Society, copyright 2023.



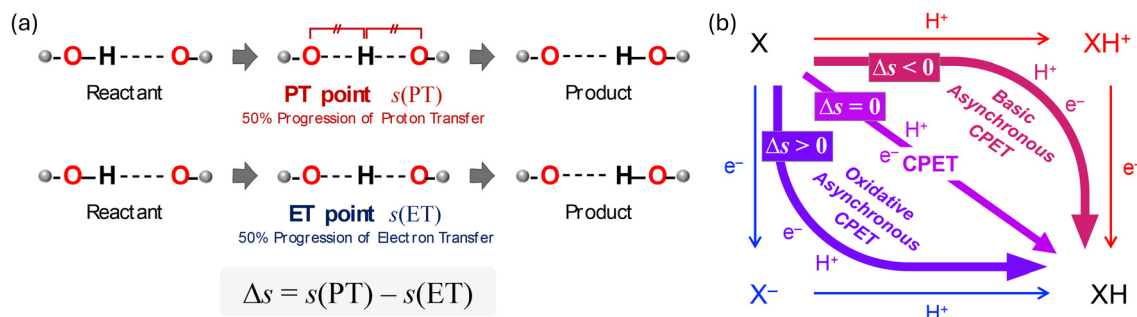


Fig. 3 (a) Conceptual representation of the definition of proton transfer (PT) point and electron transfer (ET) point. (b) Application of PT and ET points to the square scheme. When $\Delta s < 0$, PT occurs prior to ET (basic-asynchronous CPET). When $\Delta s = 0$, PT and ET proceed simultaneously (synchronous CPET). When $\Delta s > 0$, ET occurs before PT (oxidative-asynchronous CPET).

able. We recently introduced indices to spotlight the asynchronicity of PT and ET in a CPET reaction, which we called PT point and ET point, respectively. The PT and ET points designate the location of the middle progression of PT and ET in the reaction coordinate as depicted in Fig. 3. In particular, the PT point is the location where the two O-H bonds are of the same length and the ET point is the location where 50% of the total change in the spin density of the atoms involved in ET occurs. Calculating the

gap between PT and ET points (Δs) allows evaluating quantitatively their asynchronicity and classifying the type of CPET (Fig. 3(b)).

The IRC calculation results shown in Fig. 4 revealed changes in bond distance and spin density as a function of the reaction coordinate (s), where a reaction coordinate of $s = 0.0$ corresponds to a transition state. The values of two O-H distances, one N-O distance, and the spin densities for the Cu,



Fig. 4 Intrinsic coordinate plots of the changes in structures and spin densities from RC_0 to PC_0 via TS_0 in the nitrite pathway (a and c) and from RC_N to IM1_N via TS1_N in the nitro pathway (b and d) as a function of the reaction coordinate s [amu^{1/2} bohr]. The transition state is $s = 0.0$. The proton transfer (PT) point is defined as the intersection between the two O-H distances (O1-H1 and O3-H1 in (a) and O2-H1 and O3-H1 in (b)). The electron transfer (ET) point is defined as the point where the spin density for the N1 (Cu) atom reaches the midpoint of RC_0 and PC_0 in (c) (RC_N and IM1_N in (d)). Reproduced from ref. 37 with permission from the American Chemical Society, copyright 2023.

N1, O1, O2, and O3 atoms are shown. In the PT point, the two O–H bonds are of equal length, and the O–H distances change before and after PT. Meanwhile, in the ET point, 50% of the total change in spin density for the atoms involved in ET occurs. The results for the reaction from **RC_O** to **PC_O** via **TS_O** in the nitrite pathway are shown in Fig. 4(a) and (c). As depicted in Fig. 4(a), the increase in the O3–H1 distance and the decrease in the O1–H1 distance, which correspond to the migration of the H1 atom from the O3 atom to the O1 atom, is nearly complete before reaching the transition state ($s = 0.0$). However, the O1–N1 distance consistently increases throughout the reaction. Here, the PT point is defined as the intersection between the two plots of the O1–H1 and O3–H1 distances, which was determined to be $s(\text{PT}) = -1.72$. Fig. 4(c) shows that the α -spin emerges on the O3 atom, whereas the β -spin appears on the N1 and O2 atoms. The ET point is defined as the 50% completion point of the spin-density change for the N1 atom, which was identified as $s(\text{ET}) = -0.07$. Consequently, $\Delta s = s(\text{PT}) - s(\text{ET}) = -1.65$, indicating that PT occurs earlier than ET. These computational results indicate that **TS_O** represents a basic-asynchronous CPET, with “basic” referring to the dominance of the acid–base character.^{43–45}

Fig. 4(b) and (d) illustrate the results for the reaction from **RC_N** to **IM1_N** via **TS1_N** in nitro pathway. As shown in Fig. 4(b), the increase in the O3–H1 distance and the decrease in the O2–H1 distance, corresponding to the migration of the H1 atom from the O3 atom to the O2 atom, progress considerably near $s = 0$. The O1–N1 distance also increases substantially near $s = 0$; however, this change is minimal and can be attributed to the elongation of the N–O bond from a double bond to a single bond. In this case, the PT point, which is defined as the intersection between the two plots of the O2–H1 and O3–H1 distances, was determined to be $s(\text{PT}) = -0.18$. Fig. 4(d) shows an increase and a decrease in the spin densities on the Cu atom and the O3 atom, respectively. The ET point, defined as the 50% completion point of the spin-density change for the Cu atom, was identified as $s(\text{ET}) = -0.07$. Consequently, a gap of $\Delta s = -0.11$ exists between PT and ET, indicating that both processes progress almost synchronously. According to the IRC results, the reaction step via **TS1_N** was classified as a synchronous CPET. These characterizations of CPET are consistent with a previous work. Bím *et al.*⁴⁶ introduced the asynchronicity factor based on reduction potential and acidity constant, proposing an inverse correlation between asynchronicity and the imaginary frequency of the transition state of CPET. In our calculations, we also identified this inverse correlation; **TS_O** with a smaller imaginary frequency ($300i \text{ cm}^{-1}$) is characterized as basic-asynchronous CPET while **TS1_N** with a larger imaginary frequency ($1461i \text{ cm}^{-1}$) is classified as synchronous CPET.

The concept of PT and ET points allows evaluating simple reactions involving the transfer of one proton and one electron. However, it is expected to be applicable to the elucidation of more complex reaction mechanisms in biological systems, such as ion channels, which involve the transfer of multiple protons and electrons.

DFT study on the reduction of NO to N₂O catalyzed by a dicopper complex

The second topic of this Frontier article is the reduction of NO to N₂O, which is crucial in the processes of denitrification and catalytic flue gas purification. This reaction, which is catalyzed by NO reductases (NOR) and flavodiiron NO reductases, involves the formation of a hyponitrite ($\text{N}_2\text{O}_2^{2-}$) intermediate via N–N bond formation.^{4,47,48} However, details of the N–N bond formation, N–O bond cleavage, the possible intermediate species, and the precise timing of the metal redox shuttling remain to be elucidated. This knowledge will provide important insights into the design of NOR-related therapeutics and improved NO_x purification systems.

Theoretical studies on transition-metal-complex-mediated NO reduction reactions are limited. We previously reported the investigation of the mechanism of NO reduction reactions catalyzed by a dinuclear Ru complex⁴⁹ and Cu–ZSM-5⁵⁰ using DFT calculations. Recently, we focused on the experimental study by Tao *et al.*⁵¹ and theoretically elucidated the reaction mechanism of the NO reduction mediated by a dinuclear copper complex⁵² and the alkane hydroxylation performed using the oxo species produced after the reaction.⁵³ Our DFT analysis revealed an interesting ET during the NO reduction reaction.

The dinuclear copper complex bearing a 1,2-bis(di(pyridin-2-yl)methoxy)benzene ligand (**A**) reported by Tao *et al.* was found to activate NO.⁵¹ The reaction of **A** with 3 equiv. of NO yields a (μ -oxo)(μ -nitrosyl) dicopper complex and N₂O (Fig. 5). We investigated this reaction and proposed a mechanism for the reduction of NO to N₂O by a dicopper complex. In this type of reaction, a μ -oxo complex is generally considered as an intermediate or a product. Therefore, the reaction was presumed to proceed via the reaction of the dicopper complex **A** with 2 equiv. of NO to form μ -oxo complex **B** followed by the coordination of another NO molecule to **B** to give the complex



Fig. 5 Possible reaction pathway of the NO reduction using dicopper complex **A** reported by Tao *et al.* Reproduced from ref. 52 with permission from the Royal Society of Chemistry, copyright 2022.



product (Fig. 5). In that study, we focused on the NO reduction to form N_2O and B .

The reaction mechanism and ET proposed on the basis of the calculations are depicted in Fig. 6. The reaction comprises three steps: (1) N–N bond formation, (2) isomerization of N_2O_2 , and (3) N–O bond cleavage. In the initial complex, two NO molecules are coordinated to two Cu centers in an end-on manner. Initially, one electron from one Cu atom is transferred to NO, forming an intermediate with an N–N bond of 1.417 Å. The activation energy for this reaction is 21.8 kcal mol^{−1} and the formal charge of the N_2O_2 ligand is −1, which indicates that the coordination mode of the NO moiety changes from end-on to side-on. Subsequently, isomerization of the N_2O_2 ligand with rotation around the N–N bond occurs. In the transition state of this reaction, the N–N bond rotation barrier is lowered as the electrons of N_2O_2 temporarily return to the Cu atom, reducing the N–N bond order. This results in an activation energy of 14.0 kcal mol^{−1}. Regarding the electron configuration, the closed-shell singlet of the $\text{Cu}(\text{I})\text{Cu}(\text{I})$ moiety

is most stable in the transition state. After the reaction, two electrons are transferred from one of the Cu atoms to N_2O_2 , resulting in a $\text{Cu}(\text{I})\text{Cu}(\text{III})$ configuration. However, due to their instability, charge transfer between the Cu atoms leads to a $\text{Cu}(\text{II})\text{Cu}(\text{II})$ open-shell singlet or triplet as the ground state. The final step involves the cleavage of the bond between the bridged O atom and the adjacent N atom, yielding a N_2O molecule. The activation energy of this process is 7.4 kcal mol^{−1}, indicating that the N–O bond of the N_2O_2 ligand is easily broken. The overall reaction proceeds exothermically with an energy of 35.0 kcal mol^{−1}. The rate-determining step is the first step of the N–N bond formation, and its activation energy is 21.8 kcal mol^{−1}. During the NO reduction, the dicopper complex possesses various intermediates containing the *cis*- and *trans*- N_2O_2 isomers with different coordination modes to the Cu atoms. This flexibility of the dicopper complex allows adopting the most favored structures for each step in the NO reduction, *i.e.*, N–N bond formation, N_2O_2 isomerization, and N–O bond cleavage. In addition to this structural flexibility,

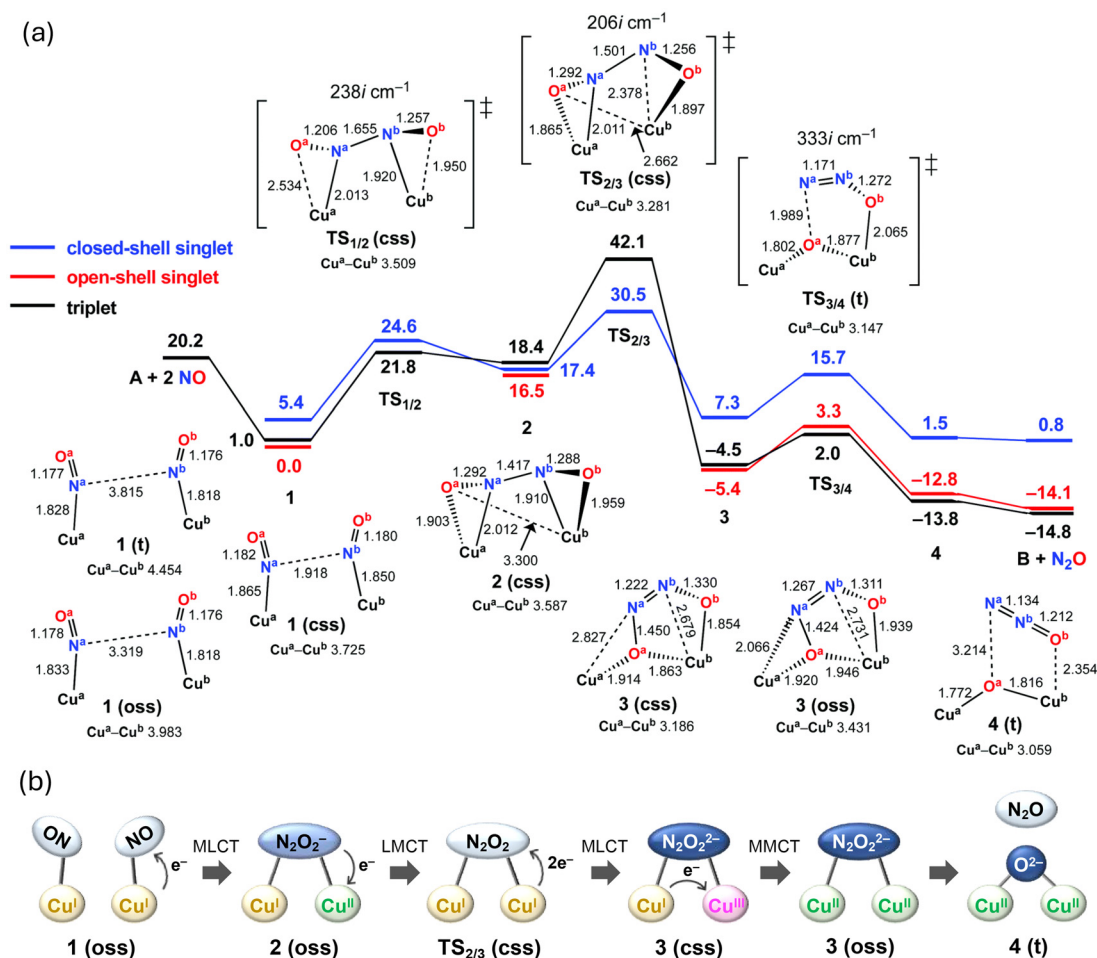


Fig. 6 (a) Free-energy profile for the reduction of NO to N_2O by dicopper complex A. Free energies are expressed in kcal mol^{−1}. Energies and optimized geometries were calculated at the B3LYP-D3/D95** (Wachters-Hay for Cu) level of theory with the solvent effect of tetrahydrofuran using the polarizable continuum model. (b) Schematic representation of the changes in electronic states during the NO reduction by the dicopper complex. Reproduced from ref. 52 with permission from the Royal Society of Chemistry, copyright 2022.

the frequent ET between the Cu atoms and the ligand, as illustrated in Fig. 6(b), facilitate the occurrence of the sequential reactions during the NO reduction. In a recent publication about the same dicopper complex, Tao *et al.* reported new kinetic data indicating the necessity of the third equivalent of NO in the reductive NO coupling.⁵⁴ The paper suggested that the third NO facilitates the N–N bond formation to form the μ -oxo μ -nitroxyl complex $[\text{Cu}_2(\text{NO})\text{O}]$ without through the μ -oxo complex $[\text{Cu}_2\text{O}]$. This mechanism is strongly linked to the disproportionation of NO ($3 \text{ NO} \rightarrow \text{NO}_2 + \text{N}_2\text{O}$). In light of these findings, it would be beneficial for future studies to consider the third NO in the mechanistic study of NO reduction reactions.

Conclusions

This Frontier article summarizes the mechanistic aspects of two important NO_x reduction processes: NO reduction and NO_2^- reduction. First, the mechanism of the reduction of NO_2^- to NO mediated by a Cu(II) complex is discussed. According to geometry optimization, we identified two types of $[\text{CuNO}_2]^+$ initial complexes, *i.e.*, an O-coordinated Cu(II) -nitrite complex (IC_O) and an N-coordinated Cu(II) -nitro complex (IC_N), which allowed proposing the nitrite and nitro reaction pathways, respectively. Given their energy profiles, the two pathways are comparable. In addition, to determine the exact nature of the CPET reactions in both pathways, we studied the changes in the geometric and electronic structures using an IRC analysis. Then, we introduced the PT and ET points as indices to evaluate the asynchronicity of PT and ET in a CPET reaction using the difference between the PT and ET points ($\Delta s = s(\text{PT}) - s(\text{ET})$, where the s value represents the location of PT and ET). In the nitrite pathway, $\Delta s = -1.65$, indicating that the PT point precedes the ET point, that is the reaction is a basic asynchronous CPET. In contrast, in the nitro pathway, the reaction is a synchronous CPET because the PT and ET points are close ($\Delta s = -0.11$). Although the two pathways lead to the same products, *i.e.*, $[\text{Cu(II)OH}]^+$, NO, and $\text{ArO}^\bullet$, their CPET mechanisms differ substantially in terms of the asynchronicity of PT and ET. Thus, by tracking the continuous change in electronic structure *via* IRC analysis, we successfully described the CPET behavior in the NO_2^- reduction. The concept of PT and ET points will facilitate the characterization and understanding of CPET reactions.

We also investigated the mechanism of the reduction of NO to N_2O by a dicopper complex using DFT calculations. The computed results indicated that the reaction consists of three fundamental steps: (1) N–N bond formation, (2) isomerization of the N_2O_2 moiety, and (3) N–O bond cleavage. The calculated reaction mechanism predicts the initial coupling of two NO molecules. These results are consistent with experimental observations, where N_2O is released in the presence of the dicopper complex. Additionally, the large binding energy of NO and Cu atoms in the reactant complex $[\text{Cu}_2(\text{NO})_2]^{2+}$ and the small binding energy of N_2O in the product complex

$[\text{Cu}_2(\text{N}_2\text{O})(\mu\text{-O})]^{2+}$ favor this catalytic cycle. Thus, DFT calculations allow following the frequent ET between Cu and the ligand in addition to structural changes in the reaction mechanism, providing a useful tool for the analysis of reaction pathways and revealing the details of the sequential N–N bond formation, isomerization of N_2O_2 , and N–O bond cleavage reactions.

Author contributions

YK – Conceptualization, writing – original draft. YS – Conceptualization, writing – review and editing.

Data availability

Data sharing is not applicable to this article as no new data were created or analyzed in this study.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 D. E. Canfield, A. N. Glazer and P. G. Falkowski, *Science*, 2010, **330**, 192–196.
- 2 D. J. Richardson and N. J. Watmough, *Curr. Opin. Chem. Biol.*, 1999, **3**, 207–219.
- 3 A. J. Timmons and M. D. Symes, *Chem. Soc. Rev.*, 2015, **44**, 6708–6722.
- 4 I. M. Wasser, S. de Vries, P. Moënne-Loccoz, I. Schröder and K. D. Karlin, *Chem. Rev.*, 2002, **102**, 1201–1234.
- 5 C. K. Jørgensen, *Coord. Chem. Rev.*, 1966, **1**, 164–178.
- 6 W. Kaim, *Inorg. Chem.*, 2011, **50**, 9752–9765.
- 7 G. Schenk, M. Y. M. Pau and E. I. Solomon, *J. Am. Chem. Soc.*, 2004, **126**, 505–515.
- 8 R. G. Serres, C. A. Grapperhaus, E. Bothe, E. Bill, T. Weyhermüller, F. Neese and K. Wieghardt, *J. Am. Chem. Soc.*, 2004, **126**, 5138–5153.
- 9 F. Roncaroli, M. Videla, L. D. Slep and J. A. Olabe, *Coord. Chem. Rev.*, 2007, **251**, 1903–1930.
- 10 V. K. K. Praneeth, F. Paulat, T. C. Berto, S. D. George, C. Näther, C. D. Sulok and N. Lehnert, *J. Am. Chem. Soc.*, 2008, **130**, 15288–15303.
- 11 M. Radon, E. Broclawik and K. Pierloot, *J. Phys. Chem. B*, 2010, **114**, 1518–1528.



- 12 M. A. Hough, J. Conradie, R. W. Strange, S. V. Antonyuk, R. R. Eady, A. Ghosh and S. S. Hasnain, *Chem. Sci.*, 2020, **11**, 12485–12492.
- 13 J. M. Mayer, D. A. Hrovat, J. L. Thomas and W. T. Borden, *J. Am. Chem. Soc.*, 2002, **124**, 11142–11147.
- 14 G. A. DiLabio and E. R. Johnson, *J. Am. Chem. Soc.*, 2007, **129**, 6199–6203.
- 15 D. Usharani, D. C. Lacy, A. S. Borovik and S. Shaik, *J. Am. Chem. Soc.*, 2013, **135**, 17090–17104.
- 16 T. Ishizuka, A. Watanabe, H. Kotani, D. C. Hong, K. Satonaka, T. Wada, Y. Shiota, K. Yoshizawa, K. Ohara, K. Yamaguchi, S. Kato, S. Fukuzumi and T. Kojima, *Inorg. Chem.*, 2016, **55**, 1154–1164.
- 17 K. Koshiba, K. Yamauchi and K. Sakai, *Angew. Chem., Int. Ed.*, 2017, **56**, 4247–4251.
- 18 M. J. Chalkley, T. J. Del Castillo, B. D. Matson and J. C. Peters, *J. Am. Chem. Soc.*, 2018, **140**, 6122–6129.
- 19 B. A. Averill, *Chem. Rev.*, 1996, **96**, 2951–2964.
- 20 S. Besson, M. G. Almeida and C. M. Silveira, *Coord. Chem. Rev.*, 2022, **464**, 214560.
- 21 L. B. Maia and J. J. G. Moura, *Chem. Rev.*, 2014, **114**, 5273–5357.
- 22 F. E. Dodd, J. Van Beeumen, R. R. Eady and S. S. Hasnain, *J. Mol. Biol.*, 1998, **282**, 369–382.
- 23 E. Culotta and D. E. Koshland, *Science*, 1992, **258**, 1862–1865.
- 24 J. O. Lundberg, M. T. Gladwin and E. Weitzberg, *Nat. Rev. Drug Discovery*, 2015, **14**, 623–641.
- 25 J. Wang, G. Keceli, R. Cao, J. T. Su and Z. Y. Mi, *Redox Rep.*, 2017, **22**, 17–25.
- 26 C. E. Sparacino-Watkins, J. Tejero, B. Sun, M. C. Gauthier, J. Thomas, V. Ragireddy, B. A. Merchant, J. Wang, I. Azarov, P. Basu and M. T. Gladwin, *J. Biol. Chem.*, 2014, **289**, 10345–10358.
- 27 M. T. Gladwin, R. Grubina and M. P. Doyle, *Acc. Chem. Res.*, 2009, **42**, 157–167.
- 28 J. W. Godden, S. Turley, D. C. Teller, E. T. Adman, M. Y. Liu, W. J. Payne and J. LeGall, *Science*, 1991, **253**, 438–442.
- 29 R. R. Eady and S. S. Hasnain, *Coord. Chem. Rev.*, 2022, **460**, 214463.
- 30 S. L. Rose, F. M. Ferroni, S. Horrell, C. D. Brondino, R. R. Eady, S. Jaho, M. A. Hough, R. L. Owen, S. V. Antonyuk and S. S. Hasnain, *J. Mol. Biol.*, 2024, **436**, 168706.
- 31 J. G. Woollard-Shore, J. P. Holland, M. W. Jones and J. R. Dilworth, *Dalton Trans.*, 2010, **39**, 1576–1585.
- 32 R. Chandra Maji, S. Mishra, A. Bhandari, R. Singh, M. M. Olmstead and A. K. Patra, *Inorg. Chem.*, 2018, **57**, 1550–1561.
- 33 A. Mondal, K. P. Reddy, J. A. Bertke and S. Kundu, *J. Am. Chem. Soc.*, 2020, **142**, 1726–1730.
- 34 S. Gupta, S. Arora, A. Mondal, S. C. E. Stieber, P. Gupta and S. Kundu, *Eur. J. Inorg. Chem.*, 2022, **15**, e202200105.
- 35 B. S. Anju, N. R. Nair and S. Kundu, *Angew. Chem., Int. Ed.*, 2023, **62**, e202311523.
- 36 G. Cioncoloni, I. Roger, P. S. Wheatley, C. Wilson, R. E. Morris, S. Sproules and M. D. Symes, *ACS Catal.*, 2018, **8**, 5070–5084.
- 37 Y. Kametani, K. Ikeda, K. Yoshizawa and Y. Shiota, *Inorg. Chem.*, 2023, **62**, 13765–13774.
- 38 J. W. Darcy, S. S. Kolmar and J. M. Mayer, *J. Am. Chem. Soc.*, 2019, **141**, 10777–10787.
- 39 S. C. Coste, A. C. Brezny, B. Koronkiewicz and J. M. Mayer, *Chem. Sci.*, 2021, **12**, 13127–13136.
- 40 R. Tyburski, T. F. Liu, S. D. Glover and L. Hammarström, *J. Am. Chem. Soc.*, 2021, **143**, 560–576.
- 41 H. Kotani, H. Shimomura, K. Ikeda, T. Ishizuka, Y. Shiota, K. Yoshizawa and T. Kojima, *J. Am. Chem. Soc.*, 2020, **142**, 16982–16989.
- 42 D. Bím, M. Maldonado-Domínguez, L. Rulíšek and M. Srnc, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, E10287–E10294.
- 43 J. S. Zhang, Y. M. Lee, M. S. Seo, Y. Kim, E. Lee, S. Fukuzumi and W. Nam, *Inorg. Chem. Front.*, 2022, **9**, 3233–3243.
- 44 M. K. Goetz and J. S. Anderson, *J. Am. Chem. Soc.*, 2019, **141**, 4051–4062.
- 45 X. H. Chen, Y.-F. Yang and Y. B. She, *Org. Chem. Front.*, 2024, **11**, 1039–1049.
- 46 D. Bím, M. Maldonado-Domínguez, L. Rulíšek and M. Srnc, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, E10287–E10294.
- 47 C. Varotsis, T. Ohta, T. Kitagawa, T. Soulimane and E. Pinakoulaki, *Angew. Chem., Int. Ed.*, 2007, **46**, 2210–2214.
- 48 T. Hayashi, J. D. Caranto, D. A. Wampler, D. M. Kurtz and P. Moënné-Loccoz, *Biochemistry*, 2010, **49**, 7040–7049.
- 49 T. Suzuki, H. Tanaka, Y. Shiota, P. K. Sajith, Y. Arikawa and K. Yoshizawa, *Inorg. Chem.*, 2015, **54**, 7181–7191.
- 50 P. K. Sajith, Y. Shiota and K. Yoshizawa, *ACS Catal.*, 2014, **4**, 2075–2085.
- 51 W. Tao, J. K. Bower, C. E. Moore and S. Zhang, *J. Am. Chem. Soc.*, 2019, **141**, 10159–10164.
- 52 Y. Kametani, T. Abe, K. Yoshizawa and Y. Shiota, *Dalton Trans.*, 2022, **51**, 5399–5403.
- 53 T. Abe, Y. Kametani, K. Yoshizawa and Y. Shiota, *Inorg. Chem.*, 2021, **60**, 4599–4609.
- 54 W. Tao, S. Carter, R. Trevino, W. Zhang, H. S. Shafaat and S. Zhang, *J. Am. Chem. Soc.*, 2022, **144**, 22633–22640.

