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Understanding the factors governing the ammonia oxidation reaction by a mononuclear ruthenium complex[†]

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Precise regulation of the active site of molecular catalysts is appealing because it could provide insights into the catalytic mechanism and possibly provide a new strategy for catalyst design. A ruthenium complex, $[\text{Ru}(\text{dppMe}, \text{COMe})(\text{bipy})(\text{Cl})]$ (CSU-3), containing $-\text{Me}$ and $-\text{COMe}$ substituted dipyridylpyrrole as a pincer ligand, was designed and synthesized. The CSU-3 complex featured a Cl^- ligand at the axial position as the active site for ammonia oxidation (AO), and is structurally analogous to AO catalyst $[\text{Ru}(\text{trpy})(\text{dmabpy})(\text{NH}_3)][\text{PF}_6]_2$ (1) bearing a terpyridine ligand, but different from AO catalyst $[\text{Ru}(\text{dpp})(\text{bipy})(\text{NH}_3)]$ (CSU-2) containing unsubstituted dipyridylpyrrole as a hemilabile ligand with the active site at an equatorial position. To gain insight into the role of active-site and ligand regulation in the AO reaction, the structure and electrochemical properties of CSU-3 and its catalytic performance and mechanism for the AO reaction were comparably studied. Complex CSU-3 has good selective catalytic performance for the oxidation of ammonia to hydrazine with a turnover frequency (TOF) of 258.8 h^{-1} and N_2H_4 formation selectivity of 84.7% at E_{app} of 1.0 V. The DFT calculations reveal that N_2H_4 as a dominant product is generated *via* an ammonia nucleophilic attack of ruthenium(IV)-imide to form N_2H_4 followed by N_2H_4 -by- NH_3 substitution.

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Introduction

Ammonia (NH_3) as a carbon-free alternative fuel is considered as one of the most important hydrogen carriers;^{1–3} however, the classical heterogeneous catalytic cracking reaction for NH_3 -to- H_2 conversion requires a precious metal catalyst and high temperature, and is therefore relatively high cost. Molecular catalytic ammonia splitting is one of the appealing alternative methodologies to produce H_2 .^{4,5} Ammonia oxidation (AO) is a critical half-reaction in NH_3 -to- H_2 conversion, and due to the involvement of multiple electron and proton transfers it is kinetically demanding. Molecular catalysts based on transition metals for the oxidation of ammonia might offer many attractive attributes, and a myriad of spectroscopic, kinetic,

thermodynamic, and electrochemical techniques have been used to gain detailed insights into the bond-breaking and bond-forming processes.^{6,7}

Numerous transition-metal complexes based on Ru,^{8–15} Fe,^{16–18} Cu,¹⁹ Ni,²⁰ and Mn,²¹ with various ligands for catalytic AO have been developed,^{22–24} since the pioneering work using a $[\text{Ru}(\text{trpy})(\text{dmabpy})(\text{NH}_3)][\text{PF}_6]_2$ (1, Scheme 1, trpy = 2,2':6',2''-terpyridine, dmabpy = 4,4,-bis(dimethylamino)-2,2'-bipyridine) catalyst⁸ for AO was reported by Smith III, Hamann and co-workers. The electron rich $-\text{NMe}_2$ group in 1 makes the catalyst decrease the onset potential of AO and triggers electrocatalytic oxidation of NH_3 to generate N_2 *via* an ammonia nucleophilic attack (ANA) mechanism, with a TOF_{N_2} of 0.7 h^{-1} . The N_2H_4 -ligated intermediates were determined through NMR spectra. As shown in Scheme 1, we recently reported a distinct example of a Ru-based catalyst,^{25a} $[\text{Ru}(\text{K}^3-\text{N},\text{N}',\text{N}'',\text{dpp})(\text{bipy})(\text{dmso})][\text{PF}_6]$ (CSU-1, bipy = 2,2'-bipyridine, Hdpp = 2,5-di(pyridin-2-yl)-1H-pyrrole) and $[\text{Ru}(\text{K}^2-\text{N},\text{N}'-\text{dpp})(\text{bipy})(\text{dmso})(\text{NH}_3)][\text{PF}_6]$ (CSU-2), which exhibits excellent electrocatalysis activity for the AO reaction to generate N_2H_4 with high selectivity (>99%) and high efficiency ($\text{TOF}_{\text{N}_2\text{H}_4} > 100 \text{ h}^{-1}$). The mechanism studies show that it benefits from the lower barrier in N_2H_4 formation involving a bimolecular coupling of Ru^{II} -aminyl or Ru^{III} -iminy intermediate, but is unfavorable for N_2 formation *via* the ANA mechanism like in complex 1.

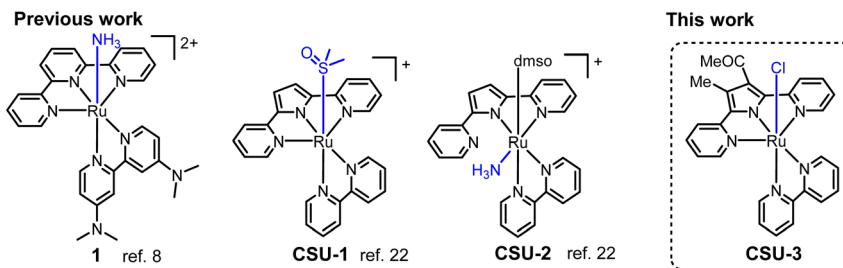
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[†] Electronic supplementary information (ESI) available: Complex synthesis and structure characterization, electrochemical property measurement methods, ammonia, hydrazine, hydrogen and nitrogen detection methods, and supplementary figures and tables. CCDC 2330568 and 2329730. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4sc02360a>

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Scheme 1 Molecular AO catalysts relevant to this work.

The anionic dipyridylpyrrole dpp⁻ ligand in **CSU-1** and **CSU-2** is structurally analogous with the neutral trpy ligand in **1**; however, their ligated ruthenium complexes exhibit significant differences in their spatial configurations, selectivity, catalytic efficiency and even mechanism for AO. The factors governing ammonia oxidation seem very complicated and deserve further in-depth research. To gain an insight into the role of the active-site in catalysis for the oxidation of ammonia, herein we design a mononuclear ruthenium(II) complex [Ru(dpp_{Me, COMe})(bipy)(Cl)] (**CSU-3**, Hdpp_{Me, COMe} = 1-(4-methyl-2,5-di(pyridin-2-yl)-1H-pyrrol-3-yl)ethan-1-one, Scheme 1) and comparably study its electrocatalysis of the AO reaction in CH₃CN media.

To avoid the hemilability of the dpp⁻ ligand to form bidentate K²-N,N'-coordination modes like in **CSU-2**, we chose the dpp_{Me, COMe}⁻ ligand in **CSU-3**. The steric effect of the substituted -Me and -COMe groups causes a smaller interior bond angle (av. 116.3°) between the pyrrole and pyridine in the free dpp_{Me, COMe}⁻ ligand (Fig. S1–S4,† crystallographic data in Tables S1 and S2†) compared to the 121.4° in the free dpp⁻ ligand.²³ This leads to an increase in the binding strength of the Ru–N bond between the Ru and N of dpp_{Me, COMe}⁻ in **CSU-3**, and causes dpp_{Me, COMe}⁻ binding to the Ru center with a N⁺N⁺N coordination mode, like trpy in **1**. Thus, the active site in **CSU-3** was regulated to the axial position for a comparative study.

Results and discussion

Synthesis and characterization

Treatment of [Ru(dmso)₄(Cl)₂], Hdpp_{Me, COMe} and bipy in the presence of Et₃N under refluxing toluene gives a red solid, which is redissolved in MeOH and refluxed for 5 d to afford **CSU-3** in 23% yield. Complex **CSU-3** was fully characterized using NMR, elemental analysis, and infrared spectroscopy (Fig. S5–S8†). The ESI-MS results display a parent peak at *m/z* 569.0282 assigned to [M]⁺. Compared to the ¹H NMR of **CSU-1**, the resonance signal of ligated dmso is absent in **CSU-3**. This result is consistent with what was observed in single crystal X-ray diffraction analysis. As shown in Fig. 1, **CSU-3** displays a slightly distorted octahedral geometry around the ruthenium center. The Cl⁻ ligand in **CSU-3** coordinates with the ruthenium center at the axial position, unlike in **CSU-1** with a dmso ligand at the axial position. Due to the steric effect of the substituted -Me and -COMe groups on the pyrrole unit, as mentioned in the introduction section, the dpp_{Me, COMe}⁻ ligand in **CSU-3** as an N⁺N⁺N pincer ligand strongly coordinates to the ruthenium

center at the equatorial position. This is mirrored by the shorter bond distance of the ruthenium center and the terminal pyridine of the dpp_{Me, COMe}⁻ ligand in **CSU-3** with Ru1–N1 of 2.107(5) Å and Ru1–N3 of 2.101(4) Å compared to that in **CSU-1** (2.136(2) Å and 2.137(2) Å). Unlike in **CSU-1**, in complex **CSU-3**, the outer donor group is not readily decoordination to supply a vacant site for coordination of the incoming NH₃. The NMR and UV-vis monitoring experiments also confirm that **CSU-3** is very stable in the presence of a coordinating solvent and even NH₃ (Fig. S9†) due to the negative charge of the Cl⁻ ligand, which makes it a poor leaving group. Thus, the corresponding NH₃-ligated complex could not be obtained by direct Cl⁻-by-NH₃ substitution of **CSU-3**.

Complex **CSU-3** is treated with AgOTf (OTf⁻ = trifluoromethylsulfonate) in CH₃CN to remove the Cl⁻ ligand, and then ammonia gas is bubbled into the filtrate solution to give NH₃-ligated complex [Ru(dpp_{Me, COMe})(bipy)(NH₃)]OTf ([CSU-3-NH₃]OTf) (Fig. S10–S13†). Its ¹H NMR spectrum shows a newly added single broad peak at 2.47 ppm due to the incoming NH₃ (Fig. S10†), which is consistent with what was observed in its ESI-MS spectrum with a parent peak at *m/z* 551.1135 for [CSU-3-NH₃]⁺ (Fig. S12†) and elemental analysis.

Electrochemical and electrocatalytic performances

The electrochemical behavior of **CSU-3** was studied using 0.1 M Bu₄NPF₆ in CH₃CN as the electrolyte, glassy carbon as the

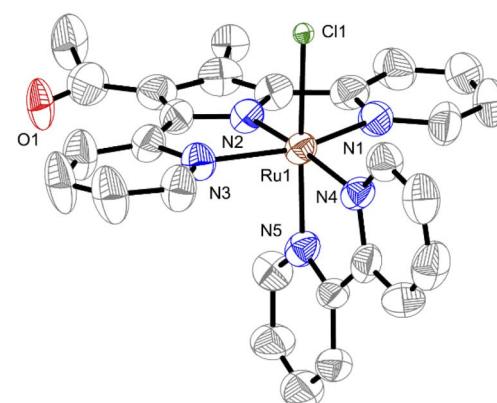


Fig. 1 Solid-state structure of **CSU-3**. The hydrogen atoms are omitted for clarity. Bond distances (Å): Ru1–N1, 2.107(5); Ru1–N2, 1.907(4); Ru1–N3, 2.101(4); Ru1–N4, 2.033(5); Ru1–N5, 2.065(4); Ru1–Cl1, 2.4297(14).



working electrode, Pt wire as the counter electrode and Ag/AgCl in saturated KCl aqueous solution as the reference electrode. Unless otherwise specified, all potentials are converted into $E_{1/2}$ versus $\text{Cp}_2\text{Fe}^{+0}$ in CH_3CN by adding -0.43 V to the measured potential.

As shown in Fig. 2a, the cyclic voltammogram (CV) of **CSU-3** displays a reversible wave at -0.14 V followed by two irreversible waves at 0.98 and 1.21 V, which are assigned to $\text{Ru}^{\text{III}/\text{II}}$, $\text{Ru}^{\text{IV}/\text{III}}$ and the ligand oxidation, respectively. The redox potential of $\text{Ru}^{\text{III}/\text{II}}$ is significantly lower than that of **1** ($\text{Ru}^{\text{III}/\text{II}}$ 0.055 V vs. $\text{Cp}_2\text{Fe}^{+0}$ in THF), **CSU-1** (0.47 V vs. $\text{Cp}_2\text{Fe}^{+0}$ in CH_3CN) and **CSU-2** (0.43 V vs. $\text{Cp}_2\text{Fe}^{+0}$ in CH_3CN). Obviously, the redox behavior of the metal center in **CSU-3** is sensitive to the electron donor nature of the dpp_{Me} , COMe^- ligand with a methyl substituent. Compared to **CSU-1** and **CSU-2** with a π -accepting dmsO ligand at the axial position, the π -electron donating Cl^- ligand at the axial position is also a possible reason for the significant negative-shift of redox potential in **CSU-3**.

For complex $[\text{CSU-3-NH}_3]\text{OTf}$, the first reversible wave (0.09 V) and the second irreversible wave (0.97 V) were attributed to continued oxidation of the ruthenium center ($\text{Ru}^{\text{II}} \square \text{Ru}^{\text{III}} \square \text{Ru}^{\text{IV}}$). The third irreversible wave at 1.34 V is due to ligand oxidation. Compared to **CSU-3**, the redox potential of the first reversible $\text{Ru}^{\text{III}/\text{II}}$ wave in $[\text{CSU-3-NH}_3]\text{OTf}$ is positively shifted. This suggests that the electron donating ability of NH_3 is weaker than that of the negatively charged Cl^- ligand. The latter acts as a π donor increasing the electron density of the ruthenium center.

The CV plot of **CSU-3** in the presence of NH_3 with various concentrations (0.01 – 0.05 M), as shown in Fig. 2b, illustrates

that the $\text{Ru}^{\text{III}/\text{II}}$ couple remains unchanged, and a new oxidation wave (~ 1.06 V) appears for the Ru^{IV} species. Subsequently, a strong catalytic current (i_{cat}) is observed (Fig. 2c), suggesting that the Ru^{IV} species triggers the oxidation of ammonia. Obviously, when the ruthenium center of **CSU-3** is oxidized to the Ru^{IV} oxidation state, an EC process occurs to generate $\text{Ru}^{\text{IV}}-\text{NH}_3$ via Cl -by- NH_3 substitution of the Ru^{IV} species from 2e^- oxidation of **CSU-3**, which is also supported by the DFT calculations. According to the CV plot of $[\text{CSU-3-NH}_3]\text{OTf}$ (Fig. 2d), in addition to the Ru^{IV} species, the Ru^{III} species from $[\text{CSU-3-NH}_3]\text{OTf}$ also triggers the oxidation of ammonia, which is very similar to what is observed in the **CSU-1**, **CSU-2** and $[\text{Ru}(\text{K}^3\text{-N},\text{N}',\text{N}'\text{-dpp})(\text{bpy})(\text{L})]\cdot\text{PF}_6$ (L = pyridine; 4-methylpyridine; pyrimidine; isoquinoline) catalyst systems and $[\text{Ru}(\text{K}^3\text{-N}',\text{N}'\text{-dpp})(\text{trpy})(\text{NH}_3)]\cdot\text{PF}_6$.^{25b,c}

As shown in Fig. S14 and S15, \dagger complexes **CSU-3** and $[\text{CSU-3-NH}_3]\text{OTf}$ exhibit satisfactory stability, corroborated by 100 consecutive cyclic voltammetry cycles, in which no new redox wave appears and the attenuation of the catalytic current is not significant. A post-catalysis and thoroughly rinsed carbon cloth working electrode displayed no catalytic activity (Fig. S16 \dagger). This indicates that the catalytic process of **CSU-3** and $[\text{CSU-3-NH}_3]\text{OTf}$ is homogenous.

Controlled potential coulometry (CPC) experiments were conducted in a sealed Schlenk electrolytic cell with a 0.01 mM ruthenium catalyst solution containing 0.2 M (or 2.0 M) NH_3 and 0.1 M $[\text{Bu}_4\text{N}]\text{PF}_6$ supporting electrolyte in anhydrous MeCN . The detection method for the possible products (H_2 , N_2 , N_2H_4 , NO_2^- and NO_3^-) and blank experiments is described in

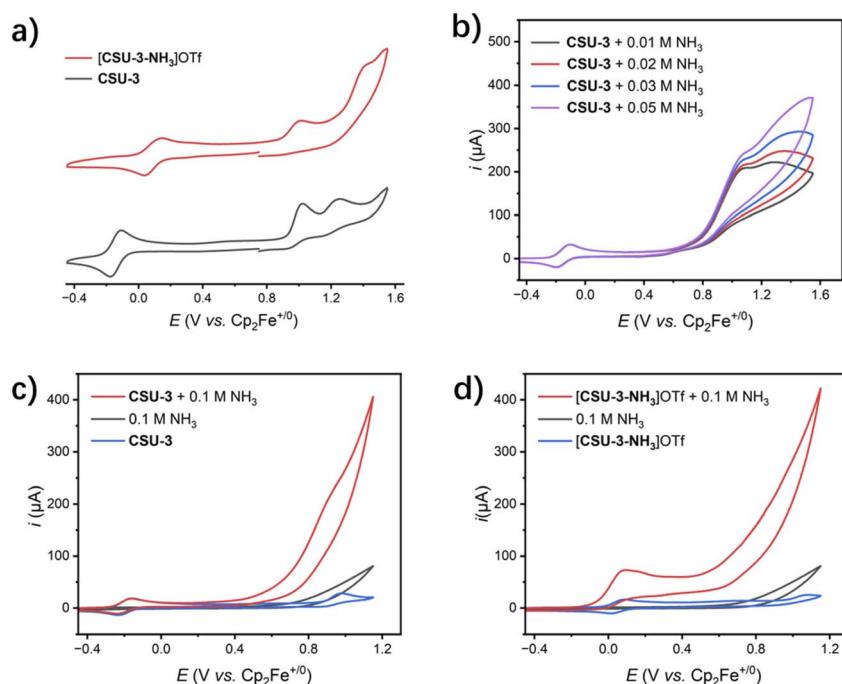


Fig. 2 (a) CV of **CSU-3** and $[\text{CSU-3-NH}_3]\text{OTf}$ in MeCN solution; (b) CV of **CSU-3** in MeCN in the presence of NH_3 with various concentrations (0.01 – 0.05 mM); (c) CV of **CSU-3** in MeCN in the presence of 0.10 M NH_3 ; (d) CV of $[\text{CSU-3-NH}_3]\text{OTf}$ in MeCN in the presence of 0.10 M NH_3 . Conditions: $[\text{Ru}] = 1$ mM, rate 0.1 V s^{-1} , 0.1 M $[\text{Bu}_4\text{N}]\text{PF}_6$ as the supporting electrolyte, platinum wire as the working electrode, potential referenced to the $\text{Cp}_2\text{Fe}^{+0}$ redox couple.



the ESI (Fig. S17–S21 and Table S5).[†] The data of catalytic performance are listed in Table 1 and Fig. S22.[†]

For complex **CSU-3**, the applied potential (E_{app}) is fixed at 0.2 V (entry 1, 2) to only generate Ru^{III} species, and as expected, in CV studies, no oxidation products of ammonia are detected. When holding the E_{app} at 1.0 V for a low concentration ammonia solution (0.2 M) for 1 h, the gas products of H₂ (113.1 μmol, 141.4 equiv. based on Ru) and N₂ (0.8 μmol, 1 equiv. based on Ru) in the headspace and N₂H₄ (112.2 μmol, 140.3 equiv. based on Ru) in the electrolyte solution are determined (entry 3). When the concentration of ammonia is increased to 2.0 M, the catalytic efficiency of **CSU-3** is approximately doubled. In addition, the selectivity of N₂H₄ formation and Faraday efficiency (FE) is almost unchanged with changes in ammonia concentration, maintaining a level of 85.6 and 84.7%, respectively. This indicates that NH₃ is possibly involved in the N₂H₄ formation step *via* an ammonia nucleophilic attack mechanism and/or N₂H₄ release *via* N₂H₄-by-NH₃ substitution.

In the **[CSU-3-NH₃]⁺OTf** catalyst system, only N₂H₄ as an anodic product is generated at low electrolytic potential (0.2 V) to only generate Ru^{III} species as the intermediate (entry 5, 6), which is very similar to the results for **CSU-1**, **CSU-2** and **[Ru(K³-N,N',N'',dpp)(bpy)(L)]⁺PF₆**.²⁵ This suggests that a bimolecular coupling mechanism of ruthenium amide is possibly involved. Holding the E_{app} at 1.0 V (Ru^{IV} species generated at this potential), the catalytic efficiency is greatly enhanced (entry 7, 8). TOF_{N₂H₄}, FE_{N₂H₄}, and S_{N₂H₄} reach 350.5 s⁻¹, 87.9% and 98.9%, respectively.

Mechanism

The full mechanism of the AO reaction catalysed by **CSU-3** is proposed by theoretical calculations. As shown in Fig. 3a, **CSU-3** is firstly oxidized to **[Ru^{III}-Cl]⁺** ($\Delta G = -8.3$ kcal mol⁻¹). The corresponding calculated Ru^{III/II} redox potential is -0.36 V. Direct Cl-by-NH₃ substitution of **[Ru^{III}-Cl]⁺** and **CSU-3** to generate **[Ru^{III}-NH₃]⁺**

and **[Ru^{II}-NH₃]⁺**, respectively, is unfavourable due to the high energy barrier (Fig. S25[†]). For example, two possible substitution pathways, namely the concerted associative pathway (*I_a*) and dissociative pathway (*D*), are considered. The energy barriers of the *I_a* pathway ($\Delta G^\ddagger = 26.2$ kcal mol⁻¹) and *D* pathway ($\Delta G^\ddagger = 24.7$ kcal mol⁻¹) are high enough to hinder direct Cl-by-NH₃ substitution of **[Ru^{III}-Cl]⁺**, which is in agreement with what is observed in CV studies and synthetic experiments. A subsequent oxidation of **[Ru^{III}-Cl]⁺** generates **[Ru^{IV}-Cl]²⁺** ($\Delta G = 22.5$ kcal mol⁻¹, $E_{cal.} = 0.98$ V). The subsequent Cl-by-NH₃ substitution of **[Ru^{IV}-Cl]²⁺** to produce **[Ru^{IV}-NH₃]³⁺** is an endergonic step with ΔG of = 12.3 kcal mol⁻¹). Subsequently, deprotonation of **[Ru^{IV}-NH₃]³⁺** affords ruthenium(IV)-imido complex **[Ru^{IV}-NH₂]²⁺**, which is a key intermediate for N₂H₄ formation.

Complex **[Ru^{IV}-NH₂]²⁺** is nucleophilically attacked by NH₃ to produce terminal hydrazinium-ligated **[Ru^{II}-NH₂NH₃]²⁺**, only overcoming an energy barrier of 1.7 kcal mol⁻¹, followed by an energetically favourable deprotonation process to generate terminal hydrazine-ligated Ru^{II}-intermediate **[Ru^{II}-NH₂NH₂]⁺** ($\Delta G = -17.4$ kcal mol⁻¹). Obviously, an N–N bond is readily formed *via* ammonia nucleophilic attack of **[Ru^{IV}-NH₂]²⁺** of **CSU-3** ($\Delta G^\ddagger = 1.7$ kcal mol⁻¹), unlike **1** and **CSU-2** *via* ammonia nucleophilic attack of Ru^{IV}-imide with higher barriers ($\Delta G^\ddagger = 24.1$ and 7.7 kcal mol⁻¹, respectively). The single-site molecular catalytic pathway of **CSU-3** is confirmed by the linear relationship between the catalytic current and concentration of ammonia and catalyst (Fig. S23 and S24[†]). Furthermore, the pathway of generating hydrazine-bridged bimetallic **[Ru^{III}-μ-N₂H₄-Ru^{III}]⁴⁺** *via* bimolecular N–N coupling **[Ru^{IV}-NH₂]²⁺** (grey line in Fig. 3b) is excluded due to the high energetic barrier ($\Delta G^\ddagger = 10.8$ kcal mol⁻¹).

N₂H₄/N₂ selectivity is usually based on the hydrazine-ligated Ru^{II}-intermediate, which could oxidize the ruthenium centre leading to hydrazine oxidation to generate N₂,^{8,17,19a,20a} but also could cause N₂H₄-by-NH₃ substitution to produce N₂H₄. As shown in Fig. 3c, N₂H₄ release through N₂H₄-by-NH₃ substitution to generate **[Ru^{II}-NH₃]⁺** *via* the *I_a* and *D* pathways was

Table 1 The electrocatalytic performances of **CSU-3** and **[CSU-3-NH₃]⁺OTf^a**

Entry	Cat.	c_{NH_3} (mol L ⁻¹)	E_{app} (V)	TOF _{H₂} (h ⁻¹) n_{H_2} (μmol)	TOF _{N₂H₄} (h ⁻¹) $n_{N_2H_4}$ (μmol)	TOF _{N₂} (h ⁻¹) n_{N_2} (μmol)	Q ^b (C)	FE _{N₂H₄} ^c (%)	S _{N₂H₄} ^d (%)
1	CSU-3	0.2	0.2	Trace	Trace	Trace	—	—	—
2	CSU-3	2.0	0.2	Trace	Trace	Trace	—	—	—
3	CSU-3	0.2	1.0	141.4 113.1	140.3 112.2	1.0 0.8	25.3	85.6	99.3
4	CSU-3	2.0	1.0	276.0 220.8	258.5 206.8	2.9 2.3	47.1	84.7	98.9
5	[CSU-3-NH₃]⁺OTf	0.2	0.2	5.0 4.0	4.8 3.8	Trace	0.8	87.9	100
6	[CSU-3-NH₃]⁺OTf	2.0	0.2	19.5 15.6	19.1 15.3	Trace	3.3	89.5	100
7	[CSU-3-NH₃]⁺OTf	0.2	1.0	169.3 135.4	165.3 132.2	1.8 1.4	28.5	89.5	98.9
8	[CSU-3-NH₃]⁺OTf	2.0	1.0	366.9 293.5	350.5 280.4	3.8 3.0	60.7	89.1	98.9

^a [Cat.] = 0.01 mM; electrolysis time, 1 h; E_{app} vs. Cp₂Fe^{+/-}; carbon cloth (1 cm²) as the working electrode; molar ratio of N₂, N₂H₄, and H₂ determined by taking the average of two tests for the electrolyte in CPC experiments, and the generation of these compounds in the control CPC experiment (Table S5) is subtracted. The maximum relative errors of 1.5%, 2.2% and 3.5% for production of H₂, N₂H₄ and N₂. ^b Charge passed in CPC experiments in 1 h. ^c FE_{N₂H₄} = $n_{N_2H_4}/Q \times 100\%$. ^d S_{N₂H₄} = $n_{N_2H_4}/(n_{N_2H_4} + n_{N_2}) \times 100\%$.



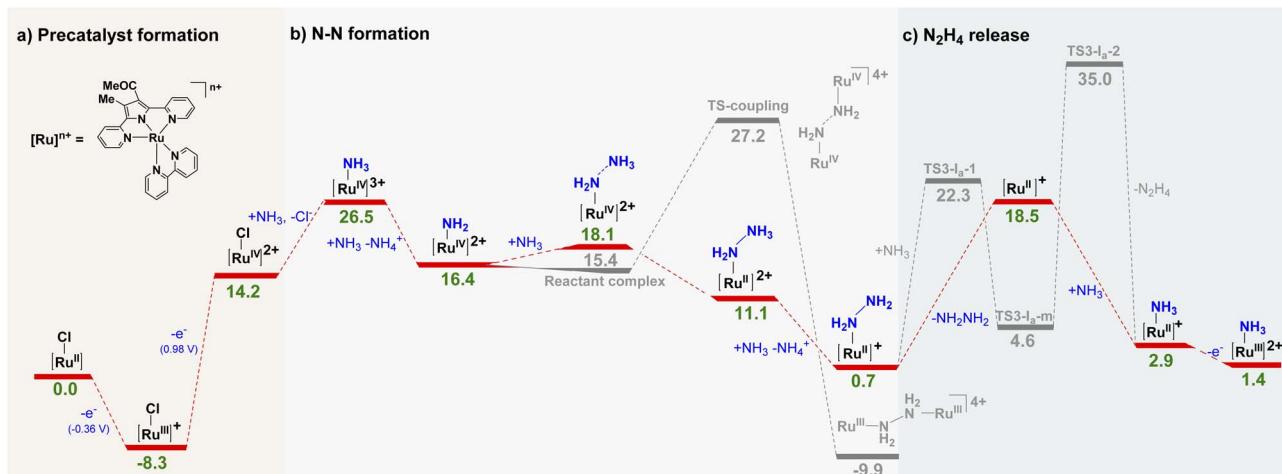


Fig. 3 The possible mechanism of AO catalyzed by CSU-3. (a) Precatalyst formation, (b) N-N formation, and (c) N₂H₄ release. The free energy changes (ΔG) are presented in the individual reaction steps in kcal mol⁻¹, with the calculated potentials in parentheses versus $\text{Cp}_2\text{Fe}^{+/0}$ in CH_3CN .

considered. Compared to the I_a mechanism with two transition states with large energetic barriers ($\Delta G^\ddagger = 21.6$ and 30.4 kcal mol⁻¹), the release of N₂H₄ via the D pathway is more favourable due to the lower energetic barrier of 17.8 kcal mol⁻¹, which is also lower than that in the CSU-2 catalytic AO system ($\Delta G^\ddagger = 23.4$ kcal mol⁻¹).

Orbital interaction and electrostatic force between Ru and hydrazine in $[\text{Ru}^{\text{II}}\text{-NH}_2\text{NH}_2]^+$ play a key role in the stabilization of the binding of the dative ligand. The energy of the lowest unoccupied molecular orbital of $[\text{Ru}^{\text{II}}]^+$ from CSU-3 shows a higher value of -0.078 au compared to $[\text{Ru}(\text{trpy})(\text{dmabpy})]^+$ from 1 (-0.093 au), indicating the relatively weaker Ru-N₂H₄ bond in $[\text{Ru}^{\text{II}}\text{-NH}_2\text{NH}_2]^+$, which is more labile (Fig. S26†). Meanwhile, natural population analysis shows that the partial charge at the ruthenium centre of $[\text{Ru}^{\text{II}}\text{-NH}_2\text{NH}_2]^+$ from CSU-3 is more positive than that from 1 (Table S6†), indicating that the influence of the electrostatic interaction is not as large as that of orbital interaction because N₂H₄ binds less strongly to the complex, where the partial charge at the ruthenium is larger. After N₂H₄-by-NH₃ substitution, the formed $[\text{Ru}^{\text{II}}\text{-NH}_3]^+$ is continuously oxidized to $[\text{Ru}^{\text{III}}\text{-NH}_3]^{2+}$, and the catalytic cycle restarts. Obviously, except for the Cl-by-NH₃ substitution in the precatalyst formation step, N₂H₄-by-NH₃ substitution (or N₂H₄ release) is the rate-determining step for the catalytic oxidation of ammonia to hydrazine. According to the literature,⁷ one-electron metal-based oxidation of $[\text{Ru}^{\text{II}}\text{-NH}_2\text{NH}_2]^+$ to $[\text{Ru}^{\text{III}}\text{-NH}_2\text{NH}_2]^{2+}$ in the complex 1 catalytic system is calculated to be the most endergonic step (31.7 kcal mol⁻¹) in the AO reaction. This electron transfer step seems to be a key ingredient in NH₃ conversion into N₂. Hence, we believe that this very thermodynamically demanding step is the possible reason that N₂ generation is unfavourable in the CSU-3 catalytic system.

Conclusions

In summary, a mononuclear ruthenium complex CSU-3 and its selective catalysis for ammonia oxidation is reported. The dpp_{Me},

COME⁻ as a pincer ligand coordinates to the ruthenium center, and the Cl⁻ ligand occupies the axial position. The redox potential of Ru^{III/II} in CSU-3 is negatively shifted to about 0.14 V and 0.61 V due to the electron donor nature of the dipyridylpyrrole ligand and the Cl⁻ axially coordinated ligand, compared to structurally analogous complex 1 and CSU-1, respectively. Complex CSU-3 could selectively catalyze the oxidation of ammonia to generate N₂H₄ as a dominant product via ammonia nucleophilic attack of the ruthenium(IV) imide forming a N-N bond, followed by an N₂H₄-by-NH₃ substitution, which is significantly distinguished from the structurally analogous 1 producing N₂ via an ANA mechanism and is also different from CSU-1 and CSU-2 bearing similar hemilabile dipyridylpyrrolide ligands, which more efficiently give the N₂H₄ product via a bimolecular coupling mechanism of the ruthenium(III)-imanyl radical. DFT calculation indicates that N₂H₄ release is the rate-determining step for NH₃-to-N₂H₄ conversion catalysed by CSU-3. The weak orbital interaction between the HOMO of N₂H₄ and the LUMO of D (the LUMO is the orbital that the dissociated N₂H₄ binds to) in the CSU-3 catalyst system may be the main reason why N₂H₄ is more easily released than in the 1 catalyst system.

Data availability

All data included in this paper are available upon request by contact with the corresponding author.

Author contributions

X.-Y. Yi designed research; G. Chen, X.-L. Ding, P. He, and T. Cheng performed research; G. Chen, X.-L. Ding, Y. Chen, J. Lin, X. Zhang, S. Zhao and N. Qiao analyzed data; G. Chen and X.-Y. Yi wrote the paper.

Conflicts of interest

There are no conflicts to declare.

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