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An energy decomposition and extrapolation scheme for evaluating electron transfer rate constants: a case study on electron self-exchange reactions of transition metal complexes†

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A simple approach to the analysis of electron transfer (ET) reactions based on energy decomposition and extrapolation schemes is proposed. The present energy decomposition and extrapolation-based electron localization (EDEEL) method represents the diabatic energies for the initial and final states using the adiabatic energies of the donor and acceptor species and their complex. A scheme for the efficient estimation of ET rate constants is also proposed. EDEEL is semi-quantitative by directly evaluating the seam-of-crossing region of two diabatic potentials. In a numerical test, EDEEL successfully provided ET rate constants for electron self-exchange reactions of thirteen transition metal complexes with reasonable accuracy. In addition, its energy decomposition and extrapolation schemes provide all the energy values required for activation-strain model (ASM) analysis. The ASM analysis using EDEEL provided rational interpretations of the variation of the ET rate constants as a function of the transition metal complexes. These results suggest that EDEEL is useful for efficiently evaluating ET rate constants and obtaining a rational understanding of their magnitudes.

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Introduction

Electron transfer (ET) plays a fundamental role in various chemical systems, including artificial photosynthesis, photocatalysis, electrocatalysis, and electronic devices.^{1–6} Theoretical tools are needed to elucidate and design novel reactivities and functions triggered by ET.^{7–10} The ET rate constant is given by eqn (1).^{9,11,12}

$$k_{\text{ET}} = \left(\frac{RT}{p^0}\right)^q \kappa_{\text{el}} \nu_n \exp\left(-\frac{\Delta G^\ddagger}{RT}\right) \quad (1)$$

where R is the gas constant, T is the temperature, p^0 is the standard atmosphere, q is either 1 or 0 for intermolecular or intramolecular ET, κ_{el} is the electronic transmission coefficient,

ν_n is an effective nuclear frequency along the reaction coordinate, and ΔG^\ddagger is the Gibbs energy of activation on the adiabatic potential energy surface (PES). κ_{el} is given by eqn (2)–(4).

$$\kappa_{\text{el}} = \begin{cases} \frac{2P_{\text{LZ}}}{1 + P_{\text{LZ}}} \text{ (If } \Delta G^0 \geq -\lambda) \\ 2P_{\text{LZ}}(1 - P_{\text{LZ}}) \text{ (If } \Delta G^0 < -\lambda) \end{cases} \quad (2)$$

$$P_{\text{LZ}} = 1 - \exp(-2\pi\gamma) \quad (3)$$

$$2\pi\gamma = \frac{\pi^{3/2}|V_{12}|^2}{h\nu_n\sqrt{\lambda k_{\text{B}}T}} \quad (4)$$

where P_{LZ} is the Landau-Zener transition probability, k_{B} is the Boltzmann constant, h is the Planck constant, V_{12} is the coupling constant, and λ is the reorganization energy (Fig. 1). When V_{12} is relatively small and $2\pi\gamma \ll 1$, the Marcus theory equation¹³ eqn (5) is obtained from Taylor series of eqn (3).

$$k_{\text{ET}} = \left(\frac{RT}{p^0}\right)^q \frac{2|V_{12}|^2}{h} \left(\frac{\pi^3}{\lambda k_{\text{B}}T}\right)^{1/2} \exp\left(-\frac{\Delta G_{\text{na}}^\ddagger}{RT}\right) \quad (5)$$

The harmonic approximation is often used to describe the diabatic PESs of the initial and final states, which allows one to estimate the Gibbs energy of activation $\Delta G_{\text{na}}^\ddagger$ (subscript “na” means “non-adiabatic”) by the following eqn (6) without explicitly locating the intersection between the two diabatic

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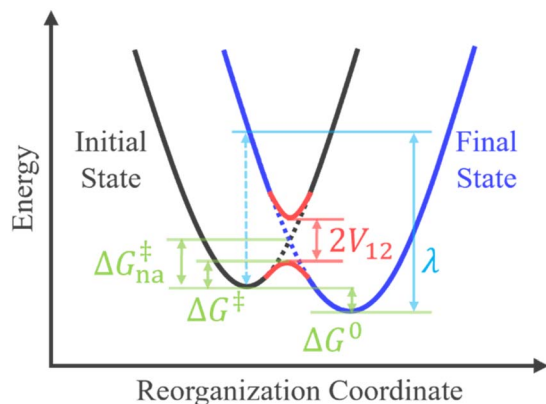


Fig. 1 Crossing of two diabatic potential energy surfaces and the relations among V_{12} , λ , ΔG^\ddagger , $\Delta G^\ddagger_{\text{na}}$, and ΔG^0 in the rate constant expressions.

Program	Step	Procedure
GRRM + G16	1	Monomer conformation search by SC-AFIR
GRRM + G16 + EDEEL	2	MESX optimization fixing D-A distance r_{DA}
	3	k_{ET} maximization along r_{DA} combined with single point calculations with larger basis set
	4	ASM analysis

Fig. 2 The entire workflow for obtaining an electron transfer rate constant by EDEEL and the calculation programs used in each procedure.

PESs on the multidimensional coordinate space. Note that the reaction barrier in eqn (1) is ΔG^\ddagger , not $\Delta G^\ddagger_{\text{na}}$ (Fig. 1).

$$\Delta G^\ddagger_{\text{na}} = \frac{(\lambda + \Delta G^0)^2}{4\lambda} \quad (6)$$

Since the diabaticization required to obtain the diabatic PESs and their coupling V_{12} is not unique, different approaches have been proposed depending on the purpose.^{14–23} For example, the generalized Mulliken–Hush (GMH) theory is for ET and generates diabatic states based on the assumption that the initial and final diabatic states are charge-localized states at different centers of the donor and acceptor.^{24,25} Constrained density functional theory (CDFT) is also useful for ET and forms diabatic states by imposing constraints on the partial charge (or

spin) of arbitrary molecules or fragments using the Lagrange multiplier.^{26,27}

To go beyond the framework based on the harmonic approximation, it is necessary to consider the anharmonic effect on the reaction coordinate. This requires explicit evaluations of PESs at around the non-adiabatic transition takes place. Molecular dynamics simulations can be performed to obtain accurate transition probabilities by calculating a large number of trajectories passing through such a region.^{9,28–31} A computationally less demanding alternative approach is to identify the seam-of-crossing (SX) or conical intersection (CI) geometry within the full-dimensional coordinate space.^{32–38} Simulations of non-adiabatic events based on SX or CI geometries have become increasingly common in recent years.^{39–45} In this approach, $\Delta G^\ddagger_{\text{na}}$ is estimated as the energy gap between the SX/CI geometry and the initial state equilibrium (EQ) geometry.

This study presents a semi-quantitative yet efficient algorithm for predicting the ET rate constant based eqn (1)–(4) rather than eqn (5) because our scheme tends to give relatively large coupling values as discussed in Results and discussion. The present method directly identifies the SX geometry. A simple diabaticization based on energy decomposition and extrapolation schemes^{46–48} allows one to optimize an SX geometry and evaluate ΔG^\ddagger , $\Delta G^\ddagger_{\text{na}}$, V_{12} , and λ . Rate constants for thirteen electron self-exchange reactions were calculated and compared with experimental^{49–59} and calculated^{60–66} values reported in the literature to investigate the performance of the present method. The calculated rate constants showed a good correlation with the experimental values. The energy decomposition scheme provided interpretations of the factors determining the magnitudes of the ET rate constants based on the activation strain model (ASM).^{67–69}

Method

In this study, a simple diabaticization scheme called energy decomposition and extrapolation-based electron localization (EDEEL) is proposed. In EDEEL, the diabatic energies for the initial and final states V_{11} and V_{22} are given by eqn (7) and (8), respectively.

$$V_{11}(\mathbf{R}) = E_{n+m}^{\text{C}}(\mathbf{R}) - E_n^{\text{D}}(\mathbf{R}^{\text{D}}) + E_{n+1}^{\text{D}}(\mathbf{R}^{\text{D}}) \quad (7)$$

$$V_{22}(\mathbf{R}) = E_{n+m}^{\text{C}}(\mathbf{R}) - E_m^{\text{A}}(\mathbf{R}^{\text{A}}) + E_{m+1}^{\text{A}}(\mathbf{R}^{\text{A}}) \quad (8)$$

where \mathbf{R} is the geometry of the donor–acceptor complex system, and \mathbf{R}^{D} and \mathbf{R}^{A} are its donor and acceptor components, respectively. The subscript for each E corresponds to the number of electrons in the corresponding system ($n = m$ in the electron self-exchange reactions). $E_{n+m}^{\text{C}}(\mathbf{R})$ is the adiabatic energy of the donor–acceptor complex without the moving electron. $E_n^{\text{D}}(\mathbf{R})$ and $E_{n+1}^{\text{D}}(\mathbf{R})$ are the adiabatic energies of the donor species without and with the moving electron, respectively. $E_m^{\text{A}}(\mathbf{R})$ and $E_{m+1}^{\text{A}}(\mathbf{R})$ are the adiabatic energies of the acceptor species without and with the moving electron, respectively. This scheme assumes that the moving electron is localized on either the donor or acceptor species and does not



Table 1 Spin multiplicity used in the calculations, experimental rate constants k_{expt} of the thirteen electron self-exchange reactions of transition metal complexes, and their data sources

Redox couple	Spin multiplicity (3+/2+)	Experimental values		
		$k_{\text{expt}}/\text{dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$	$\log_{10} k_{\text{expt}}$	References
$[\text{V}(\text{H}_2\text{O})_6]^{3+/2+}$	3/4	$3 \times 10^{-3}, 1 \times 10^{-2}$	-2.0	49 and 50
$[\text{Cr}(\text{H}_2\text{O})_6]^{3+/2+}$	4/5	$<2 \times 10^{-5}$	-4.7	51
$[\text{Fe}(\text{H}_2\text{O})_6]^{3+/2+}$	6/5	1.1	0.0	51
$[\text{Co}(\text{H}_2\text{O})_6]^{3+/2+}$	5/4	5	0.7	49 and 52
$[\text{Ru}(\text{H}_2\text{O})_6]^{3+/2+}$	2/1	$(6 \pm 4) \times 10^1$	1.8	53
$[\text{Co}(\text{NH}_3)_6]^{3+/2+}$	1/4	$>10^{-7}$	-7.0	51
$[\text{Ru}(\text{NH}_3)_6]^{3+/2+}$	2/1	4.3×10^3	3.6	49 and 54
$[\text{Ru}(\text{NH}_3)_5\text{py}]^{3+/2+}$	2/1	4.7×10^5	5.7	55
$[\text{Co}(\text{en})_3]^{3+/2+}$	1/4	7.7×10^{-5}	-4.1	56
$[\text{Ru}(\text{en})_3]^{3+/2+}$	2/1	2.8×10^4	4.4	51
$[\text{Fe}(\text{bpy})_3]^{3+/2+}$	2/1	3×10^8	8.5	57 and 58
$[\text{Co}(\text{bpy})_3]^{3+/2+}$	1/4 (1/2) ^a	18	1.3	51
$[\text{Ru}(\text{bpy})_3]^{3+/2+}$	2/1	1.2×10^9	9.1	59

^a Values for monomer and SX are shown outside and inside the parentheses, respectively.

Table 2 $\log_{10} k_{\text{calc}}$, Gibbs energies of activation ΔG^\ddagger , reorganization energy λ , diabatic coupling V_{12} , electronic transmission coefficient κ_{el} , and donor-acceptor distance r_{DA} by EDEEL at the U ω B97X-D/Def2-TZVP//Def2-SV(P) level

Redox couple	$\log_{10} k_{\text{calc}}$	$\Delta G^\ddagger/\text{eV}$	λ/eV	V_{12}/eV	κ_{el}	$r_{\text{DA}}/\text{\AA}$
$[\text{V}(\text{H}_2\text{O})_6]^{3+/2+}$	1.1	0.77	4.02	0.23	1.00	5.76
$[\text{Cr}(\text{H}_2\text{O})_6]^{3+/2+}$	-4.3	1.09	5.45	0.27	1.00	5.74
$[\text{Fe}(\text{H}_2\text{O})_6]^{3+/2+}$	0.2	0.83	4.29	0.24	1.00	5.72
$[\text{Co}(\text{H}_2\text{O})_6]^{3+/2+}$	1.0	0.78	4.10	0.24	1.00	5.53
$[\text{Ru}(\text{H}_2\text{O})_6]^{3+/2+}$	1.1	0.77	4.01	0.23	1.00	5.78
$[\text{Co}(\text{NH}_3)_6]^{3+/2+}$	-6.9	1.25	5.26	0.07	0.96	7.68
$[\text{Ru}(\text{NH}_3)_6]^{3+/2+}$	6.9	0.43	2.01	0.07	0.99	7.39
$[\text{Ru}(\text{NH}_3)_5\text{py}]^{3+/2+}$	8.6	0.33	1.65	0.08	1.00	9.51
$[\text{Co}(\text{en})_3]^{3+/2+}$	-5.5	1.16	4.86	0.05	0.91	7.96
$[\text{Ru}(\text{en})_3]^{3+/2+}$	7.1	0.42	1.90	0.06	0.98	8.02
$[\text{Fe}(\text{bpy})_3]^{3+/2+}$	11.9	0.08	0.87	0.13	1.00	9.08
$[\text{Co}(\text{bpy})_3]^{3+/2+}$	2.2	0.71	3.24	0.10	1.00	8.75
$[\text{Ru}(\text{bpy})_3]^{3+/2+}$	11.8	0.14	0.97	0.10	1.00	8.91

interact with the corresponding counterpart. In other words, this scheme sets these assumptions as the conditions for diabaticization. In the EDEEL scheme, an SX geometry between V_{11} and V_{22} is the critical point at which k_{ET} is evaluated.

$\Delta G_{\text{na}}^\ddagger = (G_{\text{SX}} - G_{\text{initial-EQ}})$ is the Gibbs energy gap between the SX and the initial state EQ geometries. Gibbs energy corrections at the EQ and SX are calculated by the normal mode analysis. The coupling V_{12} is computed as the energy difference $V_{12} = V_{11} - E_{n+m+1}^{\text{C}}$, where E_{n+m+1}^{C} is the lower adiabatic energy between those for the two states that contain the contributions of the two diabatic states the most. Thus, at the SX between V_{11} and V_{22} , *i.e.*, when $V_{11} = V_{22}$, the EDEEL scheme reproduces the diabatic-adiabatic relation of the two-state model in eqn (9).

$$E_{n+m+1}^{\text{C}} = \frac{1}{2}(V_{11} + V_{22}) - \frac{1}{2}\sqrt{(V_{11} - V_{22})^2 + 4V_{12}^2} \quad (9)$$

In the electron self-exchange reactions, E_{n+m+1}^{C} is the adiabatic energy of the donor-acceptor complex in its electronic ground state. The parameter λ is obtained as eqn (10) based on eqn (6).

$$\lambda = 2G_{\text{SX}} - G_{\text{initial-EQ}} - G_{\text{final-EQ}} + 2\sqrt{(G_{\text{SX}} - G_{\text{initial-EQ}})(G_{\text{SX}} - G_{\text{final-EQ}})} \quad (10)$$

where relations $\Delta G_{\text{na}}^\ddagger = G_{\text{SX}} - G_{\text{initial-EQ}}$ and $\Delta G^0 = G_{\text{final-EQ}} - G_{\text{initial-EQ}}$ are used, $G_{\text{final-EQ}}$ is Gibbs energy of the final state EQ, and eqn (6) gives $\lambda = 4\Delta G_{\text{na}}^\ddagger$ for electron self-exchange reactions.⁷⁰

The distance dependence of the ET reaction rate may have a distinct peak.⁶⁰ This study regards adiabatic ground state at the SX geometry between V_{11} and V_{22} as the approximate transition state and maximizes the ET rate constant k_{ET} within the SX region to obtain the final k_{ET} ($=k_{\text{calc}}$). This is done by taking the donor-acceptor distance r_{DA} (the metal-metal distance in the transition metal examples below) as the reaction coordinate, evaluating the k_{ET} at various r_{DA} , and maximizing the k_{ET} along r_{DA} . Details of how the initial $r_{\text{DA}}(\text{ini})$ is systematically determined and how the k_{ET} is maximized are described in the ESI.†

As shown in Fig. 2, the entire k_{calc} evaluation workflow consists of the following four steps.

(1) Systematic search for monomer conformations using the SC-AFIR^{71,72} method implemented in the Global Reaction Route Mapping (GRRM) program.⁷³

(2) Optimization of a minimum energy SX (MESX) geometry^{74,75} within the SX hypersurface between V_{11} and V_{22} while maintaining the donor-acceptor distance to $r_{\text{DA}}(\text{ini})$.

(3) k_{ET} maximization along r_{DA} by calculating k_{ET} for five r_{DA} values near $r_{\text{DA}}(\text{ini})$, where line search is done by simple quadratic curve fitting using three k_{ET} values in this study. In the calculation of k_{ET} with eqn (1)–(4), the Gibbs energies are extrapolated by the electronic energies obtained from single point calculations with larger basis set.



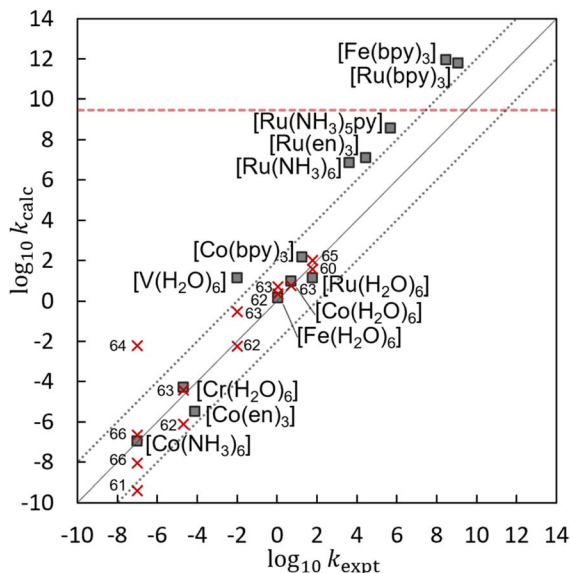


Fig. 3 A correlation between experimental $\log_{10} k_{\text{expt}}$ and calculated $\log_{10} k_{\text{calc}}$ by EDEEL at the $U\omega B97X-D/Def2-TZVP//Def2-SV(P)$ level (filled squares) or by the other method^{60–66} (red crosses). The red dotted line represents the diffusion-controlled rate constant.⁴⁹

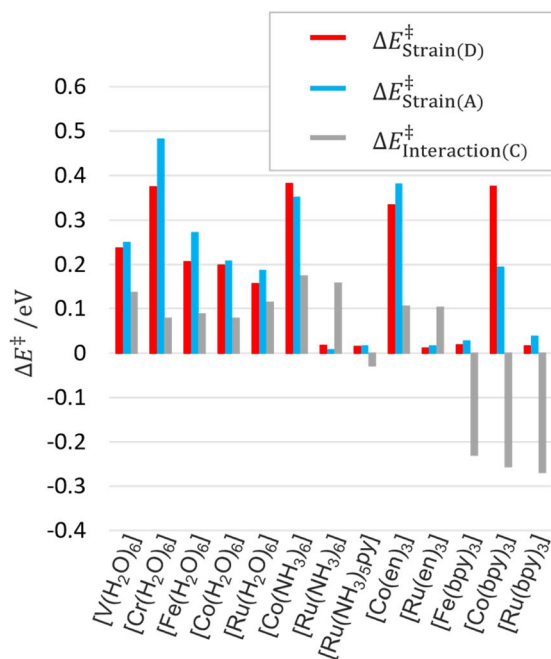


Fig. 4 Strain energies and interaction energies by ASM analysis at $U\omega B97X-D/Def2-TZVP$.

(4) ASM analysis at the geometry of the largest k_{ET} ($=k_{\text{calc}}$) obtained in step 3 (optional).

Results and discussion

The workflow in Fig. 2 was applied to the electron self-exchange reactions of thirteen transition metal complexes listed in Table

1. Electronic structure calculations were performed with the Gaussian 16 program⁷⁶ and geometry optimization was performed with the GRRM program at the $U\omega B97X-D/Def2-SV(P)$ ^{77,78} level taking into account the solvent effect of water by the conductor-like polarizable continuum model (C-PCM),^{79,80} where $U\omega B97X-D$ stands for a spin unrestricted DFT calculation using the $\omega B97X-D$ functional. k_{ET} was maximized using the electronic energy calculated at the $U\omega B97X-D/Def2-TZVP$ ⁸¹ level and the Gibbs energy correction at the $U\omega B97X-D/Def2-SV(P)$ level; the calculation level is represented as $U\omega B97X-D/Def2-TZVP//Def2-SV(P)$. In the calculation of k_{ET} , ν_n in eqn (1) was approximated as $k_{\text{B}}T/h$, the coefficient of transition state theory rate constant equation. The spin multiplicity was chosen to stabilize each state as much as possible and was set to the values listed in Table 1. The reaction set in Table 1 covers reactions of different timescales with rate constants in the wide range of 10^{-7} to $10^9 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$, and thus would be suitable as a test set. It was assumed that both +3 and +2 charged complexes were in the electronic ground state at SX, although there were possibilities for ET *via* metal-to-ligand-charge transfer (MLCT) or electronic excited states of each complex.⁸² Further computational details are presented in Computational section.

Table 2 lists the $\log_{10} k_{\text{calc}}$, ΔG^\ddagger , λ , V_{12} , κ_{el} , and r_{DA} values obtained by the present calculations at the $U\omega B97X-D/Def2-TZVP//Def2-SV(P)$ level. Fig. 3 shows the correlation between the calculated $\log_{10} k_{\text{calc}}$ and the experimental $\log_{10} k_{\text{expt}}$,^{49–59} and that the present calculation reproduces the experimental trend well. Fig. 3 also compares the $\log_{10} k_{\text{calc}}$ values with those obtained by the other theoretical methods.^{60–66} Despite the simplicity of the algorithm, which does not include a complex diabaticization scheme or any empirical factors, the present method provides an accuracy comparable to the other theoretical methods, making EDEEL promising for the semi-quantitative estimation of ET rate constants.

In Table 2, the coupling values are relatively large, making κ_{el} almost unity. This is because our diabaticization constraint assumes that there is no interaction between the electron to be transferred and the acceptor molecule. In other words, our constraint pushes the strong attraction between the negative charge on the moving electron and the +3 positive charge on the acceptor molecule into the coupling value, making the coupling values large. As for the non-adiabaticity, electron transfer reactions in $[M(\text{H}_2\text{O})_6]^{3+/2+}$, which showed the largest coupling values among the systems in our calculation, are known as non-adiabatic process with smaller actual coupling values ($<100 \text{ cm}^{-1}$).⁸³ The $\log_{10} k_{\text{calc}}$ does not include the diffusion-controlled rate constant, which is estimated to be $k_{\text{diff}} = 3 \times 10^9 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$ for these reactions.⁴⁹ In other words, k_{diff} would be a source of the large error seen for fast reactions such as $[\text{Fe}(\text{bpy})_3]^{3+/2+}$ and $[\text{Ru}(\text{bpy})_3]^{3+/2+}$. In our scheme, the C-PCM environments of the initial and final states are relaxed independently, which does not satisfy the Franck–Condon principle. This would have led to the underestimation of reaction barriers. For example, the outer-sphere reorganization energy of $[\text{Ru}(\text{H}_2\text{O})_6]^{3+/2+}$ was estimated to be 0.68 eV based on dielectric continuum theory.¹³ This implies additional reaction barrier of



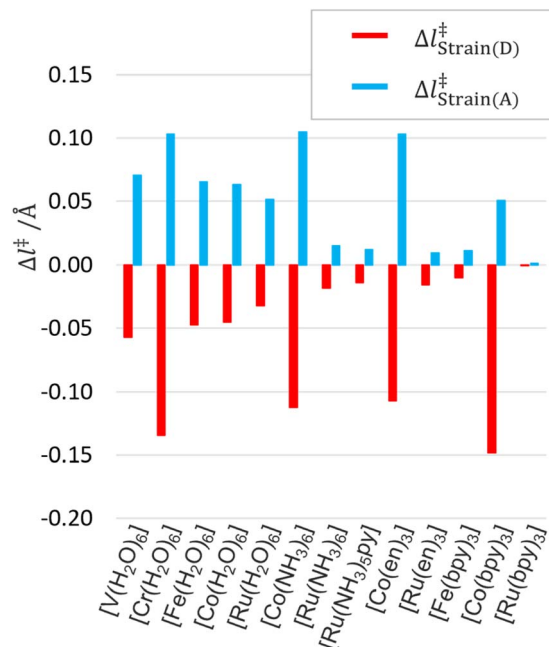


Fig. 5 Changes in mean coordination bond length.

0.17 eV, reducing its $\log_{10} k_{\text{ET}}$ from 1.1 to -1.7 , which is three-order of magnitude smaller than the experimental value. The simple addition of the outers-sphere reorganization energy correction is not sufficient in our scheme. The better treatment of outer-sphere reorganization in our scheme needs further study in the future. Many other factors such as explicit solvation, dynamic and quantum motion of the atoms, and higher order (or static) electron correlation would contribute to the error. However, further improvement of the accuracy considering these factors is beyond the scope of this study.

Thanks to the energy decomposition and extrapolation scheme of EDEEL, the ASM analysis^{67–69} can be performed using the energy components, *i.e.*, E_{n+m}^{C} , E_n^{D} , E_{n+1}^{D} , E_m^{A} and E_{m+1}^{A} , calculated during the EDEEL calculations, without any additional calculations. Note that the ASM analysis was performed using electronic energies rather than Gibbs energies for the SX geometry with the highest k_{ET} value. In the ASM/EDEEL analysis, the strain energies in donor $\Delta E_{\text{Strain(D)}}^{\ddagger}$ and acceptor $\Delta E_{\text{Strain(A)}}^{\ddagger}$ and their interaction energy $\Delta E_{\text{Interaction(C)}}^{\ddagger}$ are represented as follows.

$$\Delta E_{\text{Strain(D)}}^{\ddagger} = E_{n+1}^{\text{D(SX)}} - E_{n+1}^{\text{D(initial-EQ)}} \quad (11)$$

$$\Delta E_{\text{Strain(A)}}^{\ddagger} = E_m^{\text{A(SX)}} - E_m^{\text{A(initial-EQ)}} \quad (12)$$

$$\Delta E_{\text{Interaction(C)}}^{\ddagger} = E_{n+m}^{\text{C(SX)}} - E_n^{\text{D(SX)}} - E_m^{\text{A(SX)}} \quad (13)$$

where $E_{n+1}^{\text{D(SX)}}$, $E_n^{\text{D(SX)}}$, $E_m^{\text{A(SX)}}$ and $E_{n+m}^{\text{C(SX)}}$ are the electronic energies at the SX geometry of the donor with the moving electron, that of the donor without the moving electron, that of the acceptor without the moving electron, and that of the donor–acceptor complex without the moving electron, and $E_{n+1}^{\text{D(initial-EQ)}}$ and $E_m^{\text{A(initial-EQ)}}$ are the electronic energies at the initial state EQ

geometry of the donor with the moving electron and of the acceptor without the moving electron. The donor and acceptor in the initial EQ geometry are assumed to be infinitely far apart and not interacting. Fig. 4 and 5 show the results of the ASM/EDEEL analysis at the U ω B97X-D/Def2-TZVP level. The magnitudes of the strain energies of the aqua and cobalt complexes are large as shown in Fig. 4, reflecting the large oxidation/redox potentials of their +2/+3 species. In these systems, to compensate for the large energy gaps between the +2 and +3 species, large structural deformations ($\Delta l_{\text{Strain(D)}}^{\ddagger}$ and $\Delta l_{\text{Strain(A)}}^{\ddagger}$, changes in mean coordination bond length in donor and acceptor, respectively) are seen in the SXs around their central metal, as shown in Fig. 5. In pyridine and bipyridine complexes, attractive interactions are seen due to π -stacking between pyridine moieties and π -stacking between bipyridine molecules, respectively.^{13,49} On the other hand, aqua, ammonia, and ethylenediamine complexes show repulsive interactions, even though aqua complexes formed hydrogen bonding between water molecules. These interpretations by ASM/EDEEL would be beneficial in understanding ET rate constants quantitatively.

Conclusion

In this article, we have proposed a simple diabaticization scheme called EDEEL for the analysis of ET reactions. EDEEL represents the diabatic energies, V_{11} and V_{22} , by combining the adiabatic energies of the donor, acceptor, and their complex. Such a simple representation allows us to easily optimize the SX geometries between the two diabatic potentials. A scheme for estimating the ET rate constants at the SX geometries has also been introduced. The diabatic coupling V_{12} is also estimated using the adiabatic energy of the donor–acceptor complex. In other words, the present scheme allows one to obtain all the parameters necessary to estimate ET rate constants using only the adiabatic energies. Numerical tests with electron self-exchange reactions of thirteen transition metal complexes have shown that EDEEL reproduces the trend of the experimental rate constants well and is semi-quantitative.

Two advantages of using EDEEL can be suggested. One is that EDEEL can be combined with any *ab initio* method and program without touching their codes. This is because all diabatic potential elements, *i.e.*, V_{11} , V_{22} and V_{12} , are represented by the adiabatic energies. The other is that EDEEL provides all the energy components and SX geometries needed in the ASM analysis. The latter is helpful in interpreting the ET efficiency and further designing a system with higher ET efficiency. In the present applications, the EDEEL-based ASM analysis successfully provided rational explanations for the variation of the magnitudes of the ET rate constants depending on the transition metal complexes.

Computational section

The above procedures were implemented in an in-house Python script as an interface program between the GRRM program and the Gaussian 16 program.^{73,76} The script takes a geometry from GRRM, performs the necessary electronic structure calculations



to obtain the EDEEL PES at the geometry using Gaussian 16, and returns the energy and gradient (and Hessian if necessary) of the EDEEL PES at the geometry to GRRM. Geometry optimizations were performed by a developer version of the GRRM program at the U ω B97X-D/Def2-SV(P) level. The “Stable = Opt” option was also used to identify the electronic ground state configuration of a given spin multiplicity. The “Int(Grid = 99 590)” option was used in DFT calculations. Although some of the aqua-complexes are highly acidic and may not prevail as simple $[M(H_2O)_6]^{3+/2+}$ complexes in actual aqueous solution, the metal centers were assumed to be hexa-coordinated in this study as previous reports.^{60,62,63,65} Solvent water was modeled by the conductor-like polarizable continuum model (C-PCM).^{79,80} Gibbs energy corrections were obtained from the standard normal mode analysis of $3N-8$ dimensions (one direction orthogonal to SX and one direction along the vector between two metal atoms were removed from the full $3N-6$ dimensions) at $T = 298.15$ K and $p^0 = 1$ atm, with normal mode frequencies smaller than 100 cm^{-1} replaced by 100 cm^{-1} as suggested in the literature.⁸⁴ The bases of Gibbs activation energies were calculated as the sum of the extrapolated Gibbs energies of the most stable donor and acceptor monomers, where the most stable structures were identified by a systematic conformation search using SC-AFIR with possible spin multiplicities. An entropy correction $-4.3\text{ kcal mol}^{-1}$ was added to the Gibbs energy of the dimer complex in the activation energy calculation to account for the restriction of their mobility in the water solvent, following the previous studies.⁸⁵⁻⁸⁷ The rate constant maximization was done at the U ω B97X-D/Def2-TZVP//Def2-SV(P) level.

Conflicts of interest

There are no conflicts to declare.

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References

- D. Gust, T. A. Moore and A. L. Moore, *Acc. Chem. Res.*, 2009, **42**, 1890.
- M. H. Shaw, J. Twilton and D. W. C. MacMillan, *J. Org. Chem.*, 2016, **81**, 6898.
- N. A. Romero and D. A. Nicewicz, *Chem. Rev.*, 2016, **116**, 10075.
- N. T. Suen, S. F. Hung, Q. Quan, N. Zhang, Y. J. Xu and H. M. Chen, *Chem. Soc. Rev.*, 2017, **46**, 337.
- K. Tvrđy, P. A. Frantsuzov and P. V. Kamat, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **108**, 29.
- A. Hagfeldt, G. Boschloo, L. Sun, L. Kloo and H. Pettersson, *Chem. Rev.*, 2010, **110**, 6595.
- N. Sutin, *Prog. Inorg. Chem.*, 1983, **30**, 441.
- R. Borrelli and A. Peluso, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2013, **3**, 542.
- J. Blumberger, *Chem. Rev.*, 2015, **115**, 11191.
- S. Fletcher, *J. Solid State Electrochem.*, 2010, **14**, 705.
- M. D. Newton and N. Sutin, *Annu. Rev. Phys. Chem.*, 1984, **35**, 437.
- J. C. Tully, *J. Chem. Phys.*, 2012, **137**, 22A301.
- R. A. Marcus and N. Sutin, *Biochim. Biophys. Acta*, 1985, **811**, 265.
- C. A. Mead and D. G. Truhlar, *J. Chem. Phys.*, 1982, **77**, 6090.
- C. E. Hoyer, X. Xu, D. Ma, L. Gagliardi and D. G. Truhlar, *J. Chem. Phys.*, 2014, **141**, 114104.
- C. E. Hoyer, K. Parker, L. Gagliardi and D. G. Truhlar, *J. Chem. Phys.*, 2016, **144**, 194101.
- H. Nakamura and D. G. Truhlar, *J. Chem. Phys.*, 2001, **115**, 10353.
- H. Nakamura and D. G. Truhlar, *J. Chem. Phys.*, 2002, **117**, 5576.
- T. Lenzen and U. Manthe, *J. Chem. Phys.*, 2017, **147**, 084105.
- D. M. G. Williams, A. Viel and W. Eisfeld, *J. Chem. Phys.*, 2019, **151**, 164118.
- D. M. G. Williams and W. Eisfeld, *J. Chem. Phys.*, 2018, **149**, 204106.
- Y. Shu and D. G. Truhlar, *J. Chem. Theory Comput.*, 2020, **16**, 6456.
- Y. Shu, Z. Varga, A. G. S. de O. Filho and D. G. Truhlar, *J. Chem. Theory Comput.*, 2021, **17**, 1106.
- R. J. Cave and M. D. Newton, *Chem. Phys. Lett.*, 1996, **249**, 15.
- R. J. Cave and M. D. Newton, *J. Chem. Phys.*, 1997, **106**, 9213.
- B. Kaduk, T. Kowalczyk and T. V. Voorhis, *Chem. Rev.*, 2012, **112**, 321.
- F. Ding, H. Wang, Q. Wu, T. V. Voorhis, S. Chen and J. P. Konopelski, *J. Phys. Chem. A*, 2010, **114**, 6039.
- J. C. Tully, *J. Chem. Phys.*, 1990, **93**, 1061.
- A. V. Akimov, A. J. Neukirch and O. V. Prezhdo, *Chem. Rev.*, 2013, **113**, 4496.
- Z. Futera, K. Sodeyama, J. V. Burda, Y. Einaga and Y. Tateyama, *Phys. Chem. Chem. Phys.*, 2014, **16**, 19530.
- H. Nakano and H. Sato, *J. Chem. Phys.*, 2017, **146**, 154101.
- N. Koga and K. Morokuma, *Chem. Phys. Lett.*, 1985, **119**, 371.
- M. R. Manaa and D. R. Yarkony, *J. Chem. Phys.*, 1993, **99**, 5251.
- F. Bernardi, M. Olivucci and M. A. Robb, *Chem. Soc. Rev.*, 1996, **25**, 321.
- R. Poli and J. N. Harvey, *Chem. Soc. Rev.*, 2003, **32**, 1.
- B. G. Levine and T. J. Martínez, *Annu. Rev. Phys. Chem.*, 2007, **58**, 613.
- D. R. Yarkony, *Chem. Rev.*, 2012, **112**, 481.
- J. N. Harvey, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2014, **4**, 1.



- 39 S. Maeda, T. Taketsugu, K. Ohno and K. Morokuma, *J. Am. Chem. Soc.*, 2015, **137**, 3433.
- 40 Y. Harabuchi, T. Taketsugu and S. Maeda, *Phys. Chem. Chem. Phys.*, 2015, **17**, 22561.
- 41 M. Hatanaka, Y. Hirai, Y. Kitagawa, T. Nakanishi, Y. Hasegawa and K. Morokuma, *Chem. Sci.*, 2017, **8**, 423.
- 42 N. Aizawa, Y. Harabuchi, S. Maeda and Y. J. Pu, *Nat. Commun.*, 2020, **11**, 3909.
- 43 S. Suzuki, S. Sasaki, A. S. Sairi, R. Iwai, B. Z. Tang and G. Konishi, *Angew. Chem., Int. Ed.*, 2020, **59**, 9856.
- 44 E. Pieri, D. Lahana, A. M. Chang, C. R. Aldaz, K. C. Thompson and T. J. Martínez, *Chem. Sci.*, 2021, **12**, 7294.
- 45 Y. Harabuchi, H. Hayashi, H. Takano, T. Mita and S. Maeda, *Angew. Chem., Int. Ed.*, 2023, **62**, e202211936.
- 46 F. Maseras and K. Morokuma, *J. Comput. Chem.*, 1995, **16**, 1170.
- 47 L. W. Chung, H. Hirao, X. Li and K. Morokuma, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 327.
- 48 L. W. Chung, W. M. C. Sameera, R. Ramozzi, A. J. Page, M. Hatanaka, G. P. Petrova, T. V. Harris, X. Li, Z. Ke, F. Liu, H. B. Li, L. Ding and K. Morokuma, *Chem. Rev.*, 2015, **115**, 5678.
- 49 M. Chou, C. Creutz and N. Sutin, *J. Am. Chem. Soc.*, 1977, **99**, 5615.
- 50 K. V. Krishnamurty and A. C. Waiil, *J. Am. Chem. Soc.*, 1958, **80**, 5921.
- 51 B. S. Brunshwig, C. Creutz, D. H. Macartney, T. K. Sham and N. Sutin, *Faraday Discuss. Chem. Soc.*, 1982, **74**, 113.
- 52 N. A. Bonner and J. P. Hunt, *J. Am. Chem. Soc.*, 1960, **82**, 3826.
- 53 W. Bottcher, G. M. Brown and N. Sutin, *Inorg. Chem.*, 1979, **18**, 1447.
- 54 T. J. Meyer and H. Taube, *Inorg. Chem.*, 1968, **7**, 2369.
- 55 G. M. Brown, H. J. Krentzien, M. Abe and H. Taube, *Inorg. Chem.*, 1979, **18**, 3374.
- 56 F. P. Dwyer and A. M. Sargeson, *J. Phys. Chem.*, 1961, **65**, 1892.
- 57 I. Ruff and M. Zimonyi, *Electrochim. Acta*, 1973, **18**, 515.
- 58 D. H. Macartney and N. Sutin, *Inorg. Chem.*, 1983, **22**, 3530.
- 59 R. C. Young, F. R. Keene and T. J. Meyer, *J. Am. Chem. Soc.*, 1977, **99**, 2468.
- 60 H. Oberhofer and J. Blumberger, *Angew. Chem., Int. Ed.*, 2010, **49**, 3631.
- 61 M. D. Newton, *J. Phys. Chem.*, 1991, **95**, 30.
- 62 D. Zhang, B. Zhuang, D. Feng and C. Liu, *Int. J. Quantum Chem.*, 2000, **78**, 32.
- 63 K. M. Rosso and J. R. Rustad, *J. Phys. Chem. A*, 2000, **104**, 6718.
- 64 R. G. Endres, M. X. LaBute and D. L. Cox, *J. Chem. Phys.*, 2003, **118**, 8706.
- 65 X. Zeng, X. Hu and W. Yang, *J. Chem. Theory Comput.*, 2012, **8**, 4960.
- 66 R. L. Kenion and N. Ananth, *Phys. Chem. Chem. Phys.*, 2016, **18**, 26117.
- 67 F. M. Bickelhaupt, *J. Comput. Chem.*, 1999, **20**, 114.
- 68 L. P. Wolters and F. M. Bickelhaupt, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2015, **5**, 324.
- 69 F. M. Bickelhaupt and K. N. Houk, *Angew. Chem., Int. Ed.*, 2017, **56**, 10070.
- 70 F. P. Rotzinger, *Inorg. Chem.*, 2015, **54**, 10450.
- 71 S. Maeda, Y. Harabuchi, M. Takagi, K. Saita, K. Suzuki, T. Ichino, Y. Sumiya, K. Sugiyama and Y. Ono, *J. Comput. Chem.*, 2018, **39**, 233.
- 72 S. Maeda, Y. Harabuchi, M. Takagi, T. Taketsugu and K. Morokuma, *Chem. Rec.*, 2016, **16**, 2232.
- 73 S. Maeda, K. Ohno and K. Morokuma, *Phys. Chem. Chem. Phys.*, 2013, **15**, 3683.
- 74 M. J. Bearpark, M. A. Robb and H. B. Schlegel, *Chem. Phys. Lett.*, 1994, **223**, 269.
- 75 S. Maeda, K. Ohno and K. Morokuma, *J. Chem. Theory Comput.*, 2010, **6**, 1538.
- 76 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr, J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman, and D. J. Fox, *Gaussian 16, Revision C.02*, Gaussian, Inc., Wallingford CT, 2019.
- 77 J. D. Chai and M. H. Gordon, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6615.
- 78 F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297.
- 79 V. Barone and M. Cossi, *J. Phys. Chem. A*, 1998, **102**, 1995.
- 80 M. Cossi, N. Rega, G. Scalmani and V. Barone, *J. Comput. Chem.*, 2003, **24**, 669.
- 81 F. Weigend, *Phys. Chem. Chem. Phys.*, 2006, **8**, 1057.
- 82 C. Daniel, *Phys. Chem. Chem. Phys.*, 2021, **23**, 43.
- 83 Y. Bu, Y. Wang, F. Xu and C. Deng, *J. Mol. Struct.*, 1998, **453**, 43.
- 84 R. F. Ribeiro, A. V. Marenich, C. J. Cramer and D. G. Truhlar, *J. Phys. Chem. B*, 2011, **115**, 14556.
- 85 R. L. Martin, P. J. Hay and L. R. Pratt, *J. Phys. Chem. A*, 1998, **102**, 3565.
- 86 Y. Liu, Y. Tang, Y. Y. Jiang, X. Zhang, P. Li and S. Bi, *ACS Catal.*, 2017, **7**, 1886.
- 87 Z. Xu, Y. Yang, J. Jiang and Y. Fu, *Organometallics*, 2018, **37**, 1114.

