Long Range Proton-Coupled Electron Transfer Reactions of Bis(imidazole) Iron Tetraphenylporphyrins Linked to Benzoates

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Abstract
Concerted proton-electron transfer (CPET) reactions in iron carboxy-tetraphenylporphyrin complexes have been investigated using both experimental and theoretical methods. Synthetic heme models abstract H⁺ and e⁻ from the hydroxylamine TEMPOH or an ascorbate derivative, and the kinetics of the TEMPOH reaction indicate concerted transfer of H⁺ and e⁻. Phenylene linker domains vary the electron donor/acceptor separation by approximately 4 Å. The rate data and extensive molecular simulations show that the electronic coupling decay constant (β) depends on conformational flexibility and solvation associated with the linker domain. Our best estimate of β is 0.23 ± 0.07 Å⁻¹, a value that is near the low end of the range (0.2–0.5 Å⁻¹) established for electron transfer reactions involving related linkers. This is the first analysis of β for a CPET reaction.

Keywords
Proton-coupled electron transfer; concerted proton-electron transfer; electronic coupling; molecular dynamics; distance dependence

Electron transfer (ET) reactions accompanied by protonation changes are ubiquitous in chemistry and biology. Many of these proton-coupled electron transfer (PCET) reactions are ones in which both charged particles transfer in a single step (concerted PCET, denoted CPET). CPET is implicated in biological processes that often involve spatial separation of donor/acceptor sites,¹–³ as in the oxidation of tyrosine Z in photosystem II, which is thought to proceed by proton transfer (PT) to a nearby histidine in concert with ET to an oxidized chlorophyll P680⁺ approximately 10 Å away.¹ Separated CPET likely also is operative in ascorbate peroxidase,⁴ with ET between substrate and the heme iron (8 Å) and PT to or from the heme propionate.

A central question in both ET and CPET is how proton center separation and the composition of the intervening medium affect electron/ hole transport. This relation provides a basis for controlling the mechanisms and flow of charge in biological and synthetic systems. Studies of the distance dependence of pure ET revealed how linker domains and the associated

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ASSOCIATED CONTENT
Supporting Information. Experimental details and simulation protocols, thermodynamic and kinetics plots. This material is available free of charge via the Internet at http://pubs.asc.org.
bridge states impact the electronic coupling between redox sites. In the current study, we present the first corresponding analysis for a CPET reaction. We have employed rigid linkers in model porphyrin-based systems that allow for comparison of similar reactions at two different separations and therefore an estimate of the donor-acceptor electronic coupling constant ($\beta$). Subtle changes in both the preorganization and solvent reorganization energies are shown to play an important role in evaluation of $\beta$ values from the experimental rate constants.

Syntheses of 5-(4-carboxyphenyl)-10,15,20-triphenylporphyrin, 5-(1,1'-biphenyl-4-carboxylic acid)-10,15,20-triphenylporphyrin, and their iron complexes followed published procedures. UV-vis and $^1$H NMR spectra of the two bis(N-methyl imidazole) porphyrin complexes, denoted Fe$^{III}$PhCO$_2$H and Fe$^{III}$Ph$_2$CO$_2$H, exhibited similar features to those of the parent tetraphenylporphyrin (TPP) compounds.

Titrations of Fe$^{III}$PhCO$_2$H or Fe$^{III}$Ph$_2$CO$_2$H with the strong base 1,8-diazabicycloundecane (DBU, p$K_a$ = 24.3) in MeCN revealed small UV-vis spectroscopic changes, indicative of weak thermodynamic coupling between the iron and benzoate sites. Titrations of the ferric compounds with Et$_3$N gave the p$K_a$ values reported in Scheme 1, ca. 1 p$K_a$ unit lower than benzoic acid in MeCN (21.5). All titrations were reversible with addition of triflic acid (HOTf).

Cyclic voltammograms in MeCN (0.1 M $^4$Bu$_4$NPF$_6$) of Fe$^{III}$PhCO$_2$H and Fe$^{III}$Ph$_2$CO$_2$H were reversible with $E_{1/2}$ = −0.545 and −0.550 V vs. Cp$_2$Fe$^{+}/0$, respectively. Addition of 1 equiv. DBU (or $^4$Bu$_4$NOH) shifted the waves by −30 ± 10 and −15 ± 10 mV for Fe$^{III}$PhCO$_2$H and Fe$^{III}$Ph$_2$CO$_2$H. The shifts in potential were fully reversible upon addition of HOTf. The shifts in $E_{1/2}$ upon protonation indicate that the small coupling between the heme-iron and the benzoate decreases slightly with increasing linker length. Redox data were used to calculate the p$K_a$ of the Fe$^{II}$ porphyrins using Hess’ law (Scheme 1). The thermochemical properties of the two iron systems are very similar.

The deprotonated ferric compounds Fe$^{III}$PhCO$_2^−$ and Fe$^{III}$Ph$_2$CO$_2^−$ reacted with 2,2'-4,4'-tetramethylpiperidin-1-ol (TEMPOH, eq 1) or 5,6-isopropylidene ascorbate (iAscH$^-$, eq 2) to yield the corresponding protonated ferrous species Fe$^{II}$PhCO$_2^-$ and Fe$^{II}$Ph$_2$CO$_2^-$, as confirmed by UV-vis spectroscopy. The reactions with iAscH$^-$ and with ≥100 equivalents of TEMPO proceed to completion within minutes, before precipitation of the carboxylate species, which occurs over ~10 min for such dilute (µM) solutions under ambient conditions.

\[ \text{Fe}^{III}\text{PhCO}_{2}^{-} + \text{TEMPOH} \rightarrow \text{Fe}^{II}\text{PhCO}_{2}^{-} + \text{TEMPO} + \text{H}^{+} \]

\[ \text{Fe}^{III}\text{Ph}_{2}\text{CO}_{2}^{-} + \text{iAscH}^- \rightarrow \text{Fe}^{II}\text{Ph}_{2}\text{CO}_{2}^{-} + \text{iAsc}^{-} + \text{H}^{+} \]
Fe\textsuperscript{II}PhCO\textsubscript{2}H and Fe\textsuperscript{II}Ph\textsubscript{2}CO\textsubscript{2}H have similar O–H bond dissociation free energies (BDFEs, Scheme 1), indicating that their CPET reactions have similar driving forces. Reactions 1(n) (n = 1, 2; eq 1), under conditions of excess TEMPOH at 298 K, follow pseudo-first-order kinetics, with $k_1^{(1)} = 15.3 \pm 1.4$ M\textsuperscript{−1} s\textsuperscript{−1}; $k_1^{(2)} = 6.5 \pm 0.8$ M\textsuperscript{−1} s\textsuperscript{−1}. The corresponding reactions with TEMPO–D were markedly slower, such that kinetic measurements and quantitative deuterium kinetic isotope effects (KIEs) were complicated by the competing precipitation; however, the initial rates (d[Fe\textsuperscript{III}] / dt) were clearly slower for D-versus H-transfer in both cases.

Reactions 2(n) (n = 1, 2; eq 2) proceeded more quickly, requiring stopped-flow spectrophotometry to monitor the kinetics under pseudo-first-order conditions of excess iAscH\textsuperscript{−}. For the reaction of Fe\textsuperscript{III}PhCO\textsubscript{2}– with iAscH(D)\textsuperscript{−}, $k_2^{(1)} = (6.9 \pm 0.4) \times 10^{5}$ M\textsuperscript{−1} s\textsuperscript{−1}, and the corresponding rate constant for the deuterated system is (3.9 \pm 0.3) \times 10^{5} M\textsuperscript{−1} s\textsuperscript{−1}, such that $k_2^{(1)} / k_2^{(2)} = 1.8 \pm 0.2$. For the reaction of Fe\textsuperscript{III}Ph\textsubscript{2}CO\textsubscript{2}– with iAscH\textsuperscript{−}, $k_2^{(2)} = (4.7 \pm 0.5) \times 10^{5}$ M\textsuperscript{−1} s\textsuperscript{−1} [(4.2 \pm 0.4) \times 10^{5} M\textsuperscript{−1} s\textsuperscript{−1} for iAscD\textsuperscript{−}, $k_2^{(2)} / k_2^{(1)} = 1.12 \pm 0.15$]. These results are consistent with the small KIEs found for reactions of iAscH\textsuperscript{−} with other iron-porphyrin complexes.

Possible mechanisms for reactions 1 and 2 include initial ET followed by PT, initial PT followed by ET, or concerted transfer of H\textsuperscript{+} + e\textsuperscript{−}. \textsuperscript{1c} Fe\textsuperscript{III}PhCO\textsubscript{2}– and Fe\textsuperscript{III}Ph\textsubscript{2}CO\textsubscript{2}– have little bias towards CPET, since there is virtually no thermodynamic coupling between the redox and acid-base sites (Scheme 1), while TEMPOH strongly favors concerted H\textsuperscript{+} / e\textsuperscript{−} transfer to avoid high-energy intermediates generated from individual ET or PT. \textsuperscript{1c} $\Delta G_{\text{CPET}}^{\circ}(1)$ for concerted transfer from TEMPOH to Fe\textsuperscript{III}PhCO\textsubscript{2}– ($−3.5 \pm 1.1$ kcal/mol) is more favorable than $\Delta G_{\text{PT}}^{\circ}(1)$ or $\Delta G_{\text{ET}}^{\circ}(1)$ (+28.5, +29.6 kcal/mol, respectively). The experimentally measured barrier $\Delta G^{\ddagger}(1) = 15.8$ kcal/mol is below the $\Delta G_{\text{PT}}^{\circ}(1)$ or $\Delta G_{\text{ET}}^{\circ}(1)$, ruling against these pathways in favor of a CPET mechanism. CPET is similarly likely for reaction 1(2). \textsuperscript{18}

Thermochemical analysis of reaction 2 did not provide evidence for one specific mechanism, \textsuperscript{9} although the primary KIEs are consistent with CPET. Also, reactions 1 and 2 exhibited qualitatively similar behavior (i.e., $k_2^{(1)} > k_2^{(2)}$), consistent with CPET mediated by a (Ph)\textsubscript{n} bridge.

The rate constants for the reactions of the complex with the n = 2 linker are approximately two times slower than for the analogous $k$ values for the complex with the n = 1 linker (Table 1). This is crudely consistent with previous studies of intramolecular ET. \textsuperscript{5} To better analyze the measured rate constants for the CPET reactions involving TEMPOH we turned to molecular simulation techniques and the theory of CPET reaction rates.

Investigations of ET distance dependence employed photo-initiated unimolecular systems, which constrain the redox centers to a fixed distance, avoiding the formation of multiple, distinct precursor complexes and their associated free energy of formation. \textsuperscript{5} For the bimolecular CPET reactions studied here, the formation of a hydrogen bond between the donor and acceptor simplifies the analysis of these reactions by providing a well-defined
precursor complex. In the following analysis, the subscripts that distinguish between reactions 1 and 2 have been suppressed; we focus exclusively on reaction 1 (TEMPOH), which clearly occurs via CPET.

The thermal reaction rate for electronically nonadiabatic CPET in the normal regime assumes the golden-rule form

\[ k = \frac{\pi}{\hbar} \left[ \frac{H_{AB}^2(R)}{\Delta(R)} \right] e^{-\frac{1}{\hbar^2} \left( \frac{\lambda(R)}{2} + \frac{\Delta G^0(R)}{2} \right)} e^{-\frac{w_p(R)}{kT}}, \]  

(3)

where \( R \) is the electron donor-acceptor distance, \( \lambda(R) \) is the CPET reorganization energy given by the sum of the inner- and outer-sphere reorganization energies, \( w_p(R) \) is the work required for precursor complex formation, \( \Delta G^0(R) \) is the driving force for CPET, \( H_{AB}(R) \) is the electronic coupling matrix element, and \( \gamma \) includes the \( (n- \) and \( R \)-independent) proton coupling matrix elements. Derivation of eq 3 uses the assumption that the proton and electron donor-acceptor distances are statistically uncorrelated for configurations that dominantly contribute to the CPET rate; numerical validation of this assumption is provided in Fig. S18. Numerical support for the \( n \)- and \( R \)-independence of \( \gamma \) also is provided in Fig. S18 and Section 8.6.

At a given separation distance, \( \Delta G^0(R) \) can be related to the corresponding value at infinite separation, \( \Delta G_{\text{CPET}}^0 \), and the difference between the works of bringing together the reactant and product species.

\[ \Delta G^0(R) = \Delta G_{\text{CPET}}^0 + w_p(R) - w_p(R) \]  

(4)

In the electronically nonadiabatic regime, \( H_{AB}(R) \) varies exponentially with the electron donor-acceptor distance,

\[ H_{AB}(R) = H_{AB}^0 \exp \left( -\frac{1}{2} \beta(R - R^0) \right) \]  

(5)

Here, \( H_{AB}^0 \) is the coupling matrix element at a reference donor-acceptor distance \( R^0 \), and \( \beta \) is the associated decay constant. The ratio of the CPET rates for the complexes with \( n = 1 \) and \( n = 2 \) phenylene linkers is given by eqs 6–8.

\[ \frac{k^{(1)}}{k^{(2)}} = e^{-\frac{1}{2\beta} \Delta \Delta G^0} \int dR f^{(1)}(R) \int dR f^{(2)}(R) \]  

(6)

\[ f^{(0)}(R) = \sqrt{\frac{1}{\lambda^{(0)}(R)}} e^{-\frac{1}{2\beta} \left( \lambda^{(0)}(R) + 2w_p^{(0)}(R) + 2w_p^{(0)}(R) \right)} \]  

(7)

\[ \Delta \Delta G^0 = \Delta G_{\text{CPET}}^{(1)} - \Delta G_{\text{CPET}}^{(2)} \]  

(8)

The decay constant \( \beta \) can be obtained via numerical solution of eq 6 using the rate constants in Table 1. However, this requires terms \( w_p(R) \), \( w_p(R) \), \( \Delta \Delta G^0 \), and \( \lambda(R) \). Using a
combination of MD and electronic structure techniques, we obtained these terms for reactions 1\(^{(n)}\) and investigated their relative impact on the interpretation of \(\beta\).

We first considered the work of preorganization for the reactant and product complexes. Calculation of the reactant and product free energy profiles, \(w_r(n)(R)\) and \(w_p(n)(R)\), was performed using all-atom MD simulations with a novel setup\(^9\) that allows for the direct comparison of free energies obtained with different linker numbers, \(n\). Atomic charges for the MD simulations were obtained at the B3LYP/6-31G(d,p) level of theory. Efficient sampling was achieved by combining two-dimensional umbrella sampling with a three-dimensional implementation of the weighted histogram analysis method;\(^9\) nonetheless, a total of 400 ns of simulation time was needed to obtain sufficiently tight convergence for the analysis presented here. The reactant and product free energy profiles are presented in Figure 1.

The reactant profiles confirm that the longer (\(n = 2\)) linker domain increases the electron donor-acceptor distance by approximately 4 Å. The longer linker also leads to a wider basin of stability and slightly stronger binding of TEMPOH due to improved solvation of the hydrogen-bound complex. The product free energy profiles are qualitatively similar, but contain additional structure arising from the torsional potential associated with rotation of the carboxylic acid OH moiety (see inset structures).

Next, we computed the CPET reorganization energy \(\lambda(n)(R)\) for reactions 1\(^{(n)}\). As in the calculation of the preorganization work, the outer-sphere contribution to the reorganization energy, \(\lambda_o(n)(R)\), was obtained using all-atom MD simulations.\(^9\) We found that \(\lambda_o(n)(R)\) is only weakly sensitive to the number of phenylene linkers, \(n\), and to the electron donor-acceptor distance, \(R\) (Fig. S11); thermally averaging over \(R\) yields \(\lambda_o^{(1)} = 17.6 \pm 0.1\) kcal/mol and \(\lambda_o^{(2)} = 18.9 \pm 0.1\) kcal/mol.\(^9\) For comparison, we also evaluated the outer-sphere reorganization energy using the frequency-resolved cavity mode (FRCM),\(^{26,27}\) which provides a continuum dielectric representation for the MeCN solvent, yielding \(\lambda_o^{(1)} = 11.0 \pm 0.1\) kcal/mol and \(\lambda_o^{(2)} = 15.6 \pm 0.1\) kcal/mol.\(^9\) While qualitatively similar, FRCM predicts a stronger \(n\)-dependence for \(\lambda_o(n)(R)\) than the simulations with explicit solvent. The CPET inner-sphere reorganization energy, \(\lambda_i(n)(R)\), was computed at the B3LYP/6-31G(d,p)||B3LYP/TZVP level of theory\(^9\) using a version of Nelsen’s four-point method applied to CPET,\(^9\) yielding values of \(\lambda_i^{(1)} = 24.95\) kcal/mol and \(\lambda_i^{(2)} = 25.11\) kcal/mol.

Finally, we computed \(\Delta \Delta G^0\), the difference in driving forces for the TEMPOH reactions at infinite separation (eq 8). Driving forces for individual reactions, \(\Delta G_{\text{CPET}}^{(1)} = -3.43\) kcal/mol and \(\Delta G_{\text{CPET}}^{(2)} = -3.69\) kcal/mol, were obtained at the B3LYP/6-31G(d,p)||B3LYP/TZVP level of theory with solvation effects included via the polarizable continuum model.\(^9,28,29\) The computed values for the driving forces are in striking agreement with the corresponding experimental measurements (\(\Delta G_{\text{CPET}}^{(1)} = -3.5 \pm 1.1\) kcal/mol and \(\Delta G_{\text{CPET}}^{(2)} = -3.7 \pm 1.3\) kcal/mol) and yielded a difference of \(\Delta \Delta G^0 = 0.26\) kcal/mol, which we employ throughout the following analysis.

Having computed \(w_r(n)(R)\), \(w_p(n)(R)\), \(\lambda(n)(R)\), and \(\Delta \Delta G^0\), we examined the impact of these terms in the calculation of the decay constant \(\beta\) using eq 6. We considered a series of three cases (I–III), which provided increasingly complete descriptions of these terms.

In the simplest treatment of eq 6 (Case I), the CPET reaction was assumed to involve only a single electron donor-acceptor distance, \(R^{(n)}\), and the terms \(w_p\), \(w_r\) and \(\lambda\) were each assumed
to be independent of the number of phenylene linkers. Equation 6 then simplifies to a form that resembles what has been employed in the theoretical analysis of ET reactions, \(^{23,23}\)

\[
\frac{k^{(1)}}{k^{(2)}} = e^{-\frac{1}{kT} \Delta G^{\circ} \exp \left[ -\beta \left( \hat{R}^{(1)} - \hat{R}^{(2)} \right) \right]}
\]  

(9)

In this simplest case, the electron donor-acceptor distances for reactions \(1^{(1)}\) and \(1^{(2)}\) were estimated from the distances between the metal center and the carboxyl oxygen in the crystal structures of iron(III) tetra-4-carboxyphenylporphyrin chloride\(^{30}\) and silver(II) 5,10,15,20-tetrakis(4-carboxy-2,6-dimethylbiphenyl)porphyrin,\(^{31}\) respectively. Inserting the relative difference \(\hat{R}^{(1)} - \hat{R}^{(2)} = -4.2 \, \text{Å}\) and the experimental values for \(k^{(1)}_{1}\) and \(k^{(2)}_{1}\) into eq 9 gave \(\beta = 0.26 \pm 0.04 \, \text{Å}^{-1}\). The error bar in this estimate for \(\beta\) reflects the experimental uncertainty in the rate constants; error bars in the subsequent calculations for \(\beta\) additionally incorporate the statistical uncertainty of the numerical simulations.

For Case II, a more detailed treatment of eq 6 included the distance dependence of \(w^{(a)}_{r}(R)\) and \(w^{(a)}_{p}(R)\), while the dependence of the reorganization energy on the electron donor-acceptor distance and on the linker number was still neglected. Using the experimental values for \(k^{(1)}_{1}\) and \(k^{(2)}_{1}\), solution of eq 6 via numerical quadrature yielded \(\beta = 0.35 \pm 0.06 \, \text{Å}^{-1}\).

In the most complete treatment of eq 6 (Case III), we included the distance dependence of \(w^{(a)}_{r}(R)\) and \(w^{(a)}_{p}(R)\), and the distance- and \(n\)-dependence of \(\lambda\). Using the values for \(\lambda^{(a)}_{o}(R)\) obtained from the explicit-solvent MD simulations, eq 6 yielded \(\beta = 0.23 \pm 0.07 \, \text{Å}^{-1}\). However, using the (physically reasonable) values for \(\lambda^{(a)}_{o}(R)\) from the continuum-solvent FRCM yielded the unphysical result of \(\beta = -0.10 \pm 0.06 \, \text{Å}^{-1}\).

Table 2 presents a summary of these three analyses. Each case yielded a value for \(\beta\) that falls within the established range for ET across phenylene bridges (0.2–0.5 \, \text{Å}^{-1}),\(^{13}\) but inclusion of the various preorganization and solvent reorganization contributions was found to significantly shift \(\beta\) within that range. Specifically, comparison of Cases I and II indicates that inclusion of the difference in preorganization work for the bimolecular reactions leads to an increase of 0.09 \pm 0.04 \, \text{Å}^{-1} in the estimated value of \(\beta\), emphasizing that even weak \(n\)-dependence in the energy of preorganization (~0.3 kcal/mol) leads to a substantial difference in the calculated \(\beta\). Moreover, comparison of Cases II and III indicates that the weak \(n\)-dependence in the solvent reorganization energy leads to a decrease of 0.12 \pm 0.04 \, \text{Å}^{-1} in the calculated \(\beta\). Although the combined effects of \(w^{(a)}_{r}(R)\), \(w^{(a)}_{p}(R)\), and \(\lambda^{(a)}_{o}(R)\) in Table 2 nearly cancel for this current study, this must be regarded as fortuitous; for systems that do not exhibit this cancellation of error, a careful treatment of conformational flexibility and solvent reorganization energy would be necessary to avoid miscalculation of \(\beta\). The point is further underscored by our finding that approximations in the description of the solvent reorganization (i.e., using implicit vs. explicit solvation) can lead to an unphysical estimate for \(\beta\).

In conclusion, we have examined the bimolecular PCET reactions of Fe\(^{III}\)PhCO\(_2^{-}\) or Fe\(^{III}\)Ph\(_2\)CO\(_2^{-}\) with TEMPOH and AscH\(^-\). Thermochemical arguments show that the reactions between Fe\(^{III}\)Ph\(_n\)CO\(_2^{-}\) and TEMPOH (\(n = 1, 2\); eq 1) follow a concerted mechanism, which may also be exhibited by the ascorbate reactions (eq 2). Adding a phenyl linker and lengthening the electron donor-acceptor distance by ~4 Å leads to a decrease in CPET rate constant of only a factor of 2. Combination of the experimental results with extensive molecular simulations for reactions \(1^{(a)}\) gives the first test of the dependence of CPET reaction rates on the electron donor-acceptor distance. Notably, analysis of the value
of β for CPET in reaction 1 is sensitive to the treatment of the preorganization and solvent reorganization energies. The most detailed treatment provided here yields an estimate of $\beta \sim 0.23 \text{ Å}^{-1}$, which is on the low end of the range of values for ET across phenylene bridges ($\beta = 0.2-0.5 \text{ Å}^{-1}$). While this is only the first step in characterizing the relationship between β in ET and CPET reactions, the current results suggest that values of β obtained from ET reactions have relevance for electronically nonadiabatic CPET reactions that are central to biological catalysis and energy production.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

**Acknowledgments**

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**References**


9. Full details are given in the supporting information


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18. For FeIIIPh2CO2− + TEMPOH, ΔG°CPET, ΔG°PT, ΔG°ET, and ΔG1,2‡ = −3.7, 28.2, 29.4 and 16.3 kcal/mol, respectively


Figure 1.
(A) Free energy profiles for reactant preorganization, $w_{r}^{(n)}(R)$, associated with the TEMPOH-Fe$^{III}$Ph$_n$CO$_2^-$ complexes with $n = 1$ (red) and $n = 2$ (blue) linkers. The profiles are plotted as a function of $R$, defined as the separation between the TEMPOH oxygen and the 5-carbon of the porphyrin ring. Thermally accessible (+1 $kT$) regions of the free energy profiles are shaded. (B) The corresponding product free energy profiles, $w_{p}^{(n)}(R)$. Molecular structures indicate representative configurations for the complex associated with two basins of stability in the product profiles (with TEMPO$^*$ oriented toward the linker or into solution).
Scheme 1.
Thermochemical cycle for PCET reactions.
Table 1

PCET rate constants for reactions 1 and 2.\textsuperscript{a}

<table>
<thead>
<tr>
<th></th>
<th>Fe\textsuperscript{III}Ph\textsuperscript{−}CO\textsuperscript{2−}</th>
<th>Fe\textsuperscript{III}Ph\textsuperscript{2−}CO\textsuperscript{2−}</th>
<th>k\textsuperscript{(1)}(k\textsuperscript{(2)})</th>
</tr>
</thead>
<tbody>
<tr>
<td>iAscH</td>
<td>6.9(4) × 10\textsuperscript{5}</td>
<td>4.7(5) × 10\textsuperscript{5}</td>
<td>1.5</td>
</tr>
<tr>
<td>TEMPOH</td>
<td>15(1)</td>
<td>6.5(8)</td>
<td>2.4</td>
</tr>
</tbody>
</table>

\textsuperscript{a}Rate constants in M\textsuperscript{−1} s\textsuperscript{−1} for reactions at 298 K. Uncertainty in the last digit indicated in parentheses.
Electronic decay constant $\beta$ for three cases.

<table>
<thead>
<tr>
<th>Case</th>
<th>Explicitly in eq 6</th>
<th>calculated terms</th>
<th>$\beta / \text{Å}^{-1}$</th>
<th>$\Delta \beta / \text{Å}^{-1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>$\Delta \Delta G^0$</td>
<td></td>
<td>0.26(4)</td>
<td>---</td>
</tr>
<tr>
<td>II</td>
<td>$w_w^{\text{on}}(R), w_p^{\text{on}}(R) \Delta \Delta G^0$</td>
<td>0.35(6)</td>
<td>+ 0.09(4)$^b$</td>
<td></td>
</tr>
<tr>
<td>III$^a$</td>
<td>$w_w^{\text{on}}(R), w_p^{\text{on}}(R), \lambda / \Delta \Delta G^0$</td>
<td>0.23(7)</td>
<td>−0.12(4)$^c$</td>
<td></td>
</tr>
</tbody>
</table>

$^a$Using the explicit solvent results for $\lambda(R)$.

$^b$Relative to Case I.

$^c$Relative to Case II.