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Metal vs. ligand protonation and the alleged proton-shuttling role of the azadithiolate ligand in catalytic H₂ formation with FeFe hydrogenase model complexes†

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Electron and proton transfer reactions of diiron complexes [Fe₂adt(CO)₆] (1) and [Fe₂adt(CO)₄(PMe₃)₂] (4), with the biomimetic azadithiolate (adt) bridging ligand, have been investigated by real-time IR- and UV-vis-spectroscopic observation to elucidate the role of the adt-N as a potential proton shuttle in catalytic H₂ formation. Protonation of the one-electron reduced complex, 1^{•-}, occurs on the adt-N yielding 1H and the same species is obtained by one-electron reduction of 1H⁺. The preference for ligand vs. metal protonation in the Fe₂(I,0) state is presumably kinetic but no evidence for tautomerization of 1H to the hydride 1Hy was observed. This shows that the adt ligand does not work as a proton relay in the formation of hydride intermediates in the reduced catalyst. A hydride intermediate 1HHy⁺ is formed only by protonation of 1H with stronger acid. Adt protonation results in reduction of the catalyst at much less negative potential, but subsequent protonation of the metal centers is not slowed down, as would be expected according to the decrease in basicity. Thus, the adtH⁺ complex retains a high turnover frequency at the lowered overpotential. Instead of proton shuttling, we propose that this gain in catalytic performance compared to the propyldithiolate analogue might be rationalized in terms of lower reorganization energy for hydride formation with bulk acid upon adt protonation.

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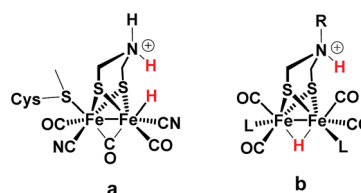
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Introduction

Proton binding sites in the second coordination sphere of catalytic metal centres are widely considered as an important design principle for facilitating proton coupled electron transfer in molecular catalysts for *e.g.* water splitting,^{1–5} H₂ formation and activation^{6–12} or CO₂ reduction.^{13–21} For the directed design of synthetic catalysts it is, however, important to note that improvements to catalytic performance (TOF, overpotential) brought about by basic sites in the second coordination sphere are not necessarily arising from a proton relay activity. Instead, the modulation of redox and acid/base properties of the metal centre upon changes in protonation state in the second coordination sphere may be the actual origin of improved catalytic performance.^{14–17}

Regarding catalysts for H₂ formation, the N₂P₂ ligands introduced by DuBois and co-workers,^{6,7,12} and the azadithiolate (adt) bridging ligand in diiron complexes of the general formula Fe₂adt(CO)₄X₂ (X = *e.g.* CO, R₃P) modelled after the FeFe-Hase active site,^{9,22–24} are prominent examples of ligand motifs with basic sites in the second coordination sphere. The amine function of the adt ligand is believed to assist enzymatic H₂ formation and activation by shuttling of protons to or from the distal iron centre of the active site where the crucial terminal hydride intermediate is formed (Scheme 1a).^{25,26} A similar role of the adt ligand in catalysis by the synthetic models considered here seems less likely given the preferred bridging coordination of the hydride (Scheme 1b).²³ The superior catalytic performance of Fe₂(adt)(CO)₆, 1, over its propyldithiolate (pdt)



Scheme 1 FeFe Hase active site (a) and synthetic model (b) bearing a hydride and a proton.

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which avoids protonation of the neutral parent complex, shifts all three ν_{C-O} bands by about 20 cm^{-1} to higher wavenumbers (2025 , 1965 and 1935 cm^{-1}), as shown in Fig. 2. The magnitude of the shift is similar to what is observed upon protonation of the neutral parent complex with stronger acids (*cf.* Fig. 3a) and characteristic for protonation of the adt-N.²⁹



Metal protonation of $\mathbf{1}^-$ is on the other hand expected to shift the carbonyl bands by about 80 cm^{-1} to higher wavenumbers. Shifts of this magnitude have been established for the bridging hydrides formed by protonation of $\text{Fe}_2(\text{I},\text{I})$ complexes with electron donor ligands like $4'$ (ref. 35) as well as transiently generated $\text{Fe}_2(\text{I},\text{0})$ hexacarbonyl complexes 2^- and 3^- .^{32,33} Mutually cancelling shifts of the carbonyl bands upon one-electron reduction and metal protonation were further evidenced with help of complex 4 (*vide infra*). The transient IR spectrum of the protonation product hence proves that the adt ligand remains the preferred site of protonation, at least kinetically, also upon one-electron reduction of $\mathbf{1}$. This conclusion is corroborated by the UV-vis spectrum of $\mathbf{1H}$ (ESI[†]), which is similar to its precursor $\mathbf{1}^-$, while metal protonation of 2^- was previously shown to cause complete bleach of its visible absorption bands that arise from transitions involving predominantly metal-based orbitals. The observed pseudo-first order rate constants for formation of $\mathbf{1H}$ follow a linear dependence on acid concentration before becoming limited by

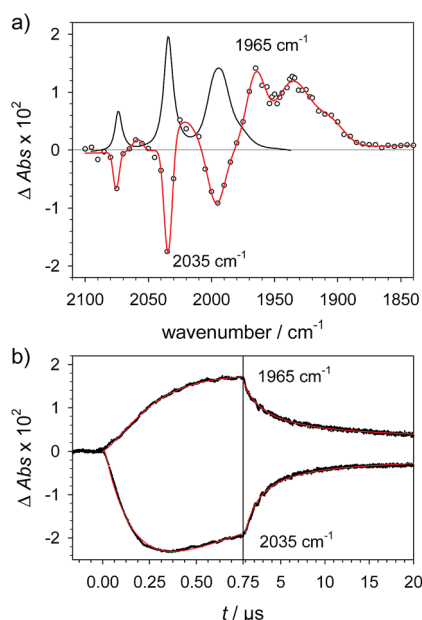


Fig. 2 (a) Transient IR spectrum $1\ \mu\text{s}$ after excitation (O, $\color{red}{\text{---}}$) and normalized IR spectrum of the starting state (---) for formation of $\mathbf{1H}$ by protonation of $\mathbf{1}^-$ with Cl_3CCOOH (13 mM) following reduction of $\mathbf{1}$ (1.3 mM) by flash-quench generated $[\text{Ru}(\text{dmb})_3]^+$ in acetonitrile. (b) Kinetic traces (\cdots) and fits ($\color{red}{\text{---}}$) monitoring pseudo-first order formation of $\mathbf{1H}$ (1965 cm^{-1}) and bleach of $\mathbf{1}$ (2035 cm^{-1}) followed by second order charge recombination with TTF^+ .



Fig. 3 (a) FTIR spectra of 4 (1.5 mM) (---) in acetonitrile, 4H^+ ($\color{blue}{\text{---}}$) obtained by protonation with toluenesulfonic acid (3 mM), and 4Hy^+ ($\color{red}{\text{---}}$) obtained by protonation with toluenesulfonic acid (3 mM) in presence of tetrabutyl-ammonium chloride (1.5 mM) after correction of the raw spectrum ($\color{red}{\text{---}}$) for minor contribution from 4H^+ . (b) Transient IR spectrum upon reduction of $[\text{Ru}(\text{dmb})_3]^+$ (O, $\color{red}{\text{---}}$, $1\ \mu\text{s}$ after excitation) and normalized IR spectra of 4Hy^+ ($\color{red}{\text{---}}$) and 4 (---).

the preceding electron transfer step (ESI[†]). The straightforward concentration dependence excludes any complications of the protonation kinetics due to dimerization of the acetic acids that was previously found to result in quadratic concentration dependence corresponding to an increase in effective acid strength.³⁶ Interestingly, protonation of $\mathbf{1}^-$, which occurs unambiguously at the adt-N, proceeds remarkably slowly with bimolecular rate constants k_{PT1} of $4 \times 10^8\text{ M}^{-1}\text{ s}^{-1}$ and $2 \times 10^7\text{ M}^{-1}\text{ s}^{-1}$ for Cl_3CCOOH and ClH_2CCOOH , respectively.

Notably, the protonation rate constant is almost two orders of magnitude below the diffusion controlled limit even for the strongly exergonic reaction with Cl_3CCOOH ($\Delta\text{p}K_{\text{a}} = 3$). This is quite untypical for protonation of an amine base and indicative of significant reorganization associated with ligand protonation.

Spectral comparison against the one-electron reduced hydride 4Hy

The above assignment of the protonation reaction to ligand protonation builds to a large extent on the general notion of nearly perfectly cancelling shifts in carbonyl frequencies caused by one-electron reduction and formation of a bridging hydride. This expectation could previously only be supported by the vanishing transient absorption observed upon protonation of complexes 2^- and 3^- .^{32,33} Here, these assignments could be



eventually verified by the positive observation of a one-electron reduced hydride **4Hy** by laser flash induced reduction of hydride complex **4Hy**⁺. The latter can be obtained by the Cl⁻-catalysed protonation of **4** that yields the thermodynamically stable hydride instead of the kinetically preferred ligand protonated **4H**⁺ in analogy to the previously reported behaviour of **4**.³⁷ Upon one electron reduction of **4Hy**⁺ the carbonyl bands of the product **4Hy** coincide almost perfectly with those of the parent complex **4** (Fig. 3) verifying the close spectral resemblance of the diiron carbonyl complexes and their bridging hydride derivatives in the Fe₂(I,0) oxidation state.

Probing possible tautomerization of **1H**

According to the above arguments, also formation of a possibly more stable hydride **1Hy** by tautomerization (5) should lead to complete decay of all transient absorption given the predictable spectral similarity between **1** and **1Hy**.



While the expected spectral changes are the same as for charge recombination between **1H** and TTF⁺ (6), tautomerization should nevertheless accelerate the decay of **1H** if it proceeds with a rate that is at least comparable to charge recombination (6).



In flash photolysis experiments, **1H** decays with second order kinetics on a time scale of about 10 μs in transient IR and about 100 μs with the lower transient concentrations in UV-vis experiments. These time scales are typical for diffusion controlled recombination ($k_{\text{rec},2} = 1.5 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$) and put an upper limit on the order of 10^4 s^{-1} on the rate constant of a possibly competing tautomerization reaction.

As a turnover frequency (TOF) of 10^3 to 10^4 s^{-1} has been proposed from electrocatalytic experiments ($\log(k_{\text{cat}}/\text{s}^{-1}) = 3.9$ and 2.6 for 6 mM Cl₃CCOOH and ClH₂CCOOH, respectively),²⁷ we wanted to probe the reactivity of **1H** on a longer time scale than 100 μs. Thus, **1H**⁺ was chemically reduced by cobaltocene in IR stopped flow experiments (rapid-scan FTIR). **1H**⁺ was obtained by addition of two equivalents of 2,5-dichlorobenzenesulfonic acid (Cl₂BSA, $\text{p}K_{\text{a}} = 6.7$)³⁴ to warrant quantitative protonation of **1**. Reduction to **1H** occurred on a shorter time scale than the time resolution of the experiment. Importantly, **1H** was observed for several seconds (Fig. 4). Its decay with a lifetime of 1.3 s yields a product spectrum that can be assigned to the parent complex **1** or the tautomerization product **1Hy** given the expected similarity of their spectra. The former could be generated by catalytic turnover that leads to depletion of acid and therefore regenerates unreduced, unprotonated catalyst. Whether the slow decay of **1H** is predominantly due to turnover or tautomerization cannot be discriminated but the observed lifetime demonstrates in either case that the rate constant for tautomerization, if this occurs at all, has to be smaller than 1 s^{-1} . We note that tautomerization

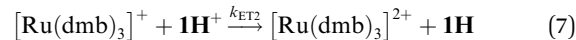


Fig. 4 Rapid scan FTIR spectra in acetonitrile between 50 ms (—) and 3 s (—) after mixing of **1H**⁺ (1.3 mM **1**, 2.6 mM Cl₂BSA) with cobaltocene (20 mM) (a) and kinetic traces (○) with exponential fits (—) monitoring decay of **1H** at 1960 cm⁻¹ (b) and formation of **1** at 2034 cm⁻¹ (c).

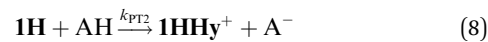
would be an intramolecular process that is independent on external conditions, such as identity and concentration of acid and the method of reduction (chemical or electrochemical).

Reduction and subsequent protonation of **1H**⁺

Alternative formation of **1H** via the PT-ET route was studied by laser-flash induced reduction of **1H**⁺ (7).



1H⁺ was obtained by protonation of **1** with Cl₂BSA that shifts the IR peaks by about 20 cm⁻¹ to higher wavenumbers (2090, 2051 and 2015 cm⁻¹).²⁹ The transient IR absorption spectrum after reduction of **1H**⁺ (Fig. 5) shows the bleaching of all three ν_{C-O} peaks of **1H**⁺ together with the same product absorption peaks previously attributed to **1H** (2025, 1965, 1935 cm⁻¹). While the ET-PT and PT-ET sequences lead expectedly to the same product **1H**, its decay is much faster when generated in presence of the stronger acid required for the initial ligand protonation. The accelerated decay of **1H** (Fig. 5) without appearance of new transient signals can be assigned to its metal protonation yielding **1HHy**⁺ (8).



This assignment is based on the expected IR absorption of **1HHy**⁺ that is likely to cancel the bleach of absorption from the starting material **1H**⁺ as previously found for hydrides **2Hy** and **3Hy**.^{32,33} Protonation is also manifested in a slower charge recombination of **1HHy**⁺ with TTF⁺ (ESI†). Kinetic analysis of both **1H** and TTF⁺ signals results in a protonation rate constant of $k_{\text{PT}2} = 6 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ (see ESI†). The analysis eliminates the possible effect of accelerated recombination with accumulated TTF⁺ in the presence of acid. With a rate constant of $6 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ formation of the hydride **1HHy**⁺ by metal protonation of **1H** with Cl₂BSA ($\text{p}K_{\text{a}} 6.7$)³⁴ occurs about as fast as metal protonation of the pdt analogue **2**⁻ ($7 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$)³² by TSOH



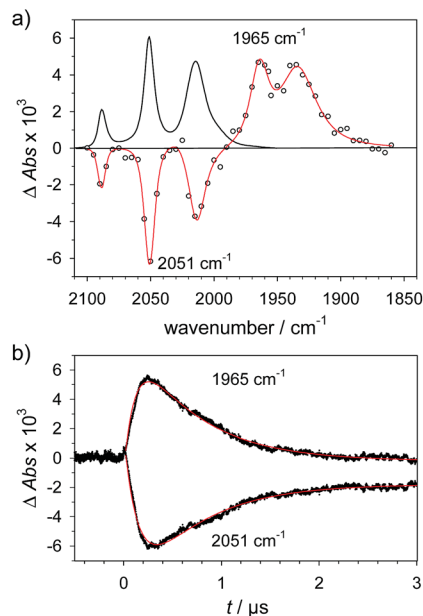


Fig. 5 (a) Transient IR spectrum 0.3 μs after excitation (O, ---) and normalized IR spectrum of the starting state (---) for formation of 1H by reduction of 1H^+ (obtained by protonation of 1 with 2.6 mM Cl_2BSA) with flash-quench generated $[\text{Ru}(\text{dmb})_3]^+$ in acetonitrile. (b) Kinetic traces (\cdots) and fits (---) monitoring pseudo-first order formation of 1H (1965 cm^{-1}) and bleach of 1H^+ (2051 cm^{-1}) followed by pseudo-first order decay of 1H by protonation with Cl_2BSA and charge recombination with TTF^+ .

($\text{p}K_{\text{a}}$ 8–8.7).³⁴ Both complexes are in the same $\text{Fe}_2(\text{l},0)$ oxidation state but the $\text{p}K_{\text{a}}$ of the hydride should drop by about 5 units according to a shift of +330 mV in reduction potential for the $1\text{H}^+/1\text{H}$ couple compared to $1/1^-$ or $2/2^-$. Protonation of 1H , even by the stronger Cl_2BSA , is therefore thermodynamically less favourable than protonation of 2^- by TsOH , with $\Delta\text{p}K_{\text{a}}$ lowered by about 3 for the former reaction. Applying the free energy relationship (Brønsted coefficient of 0.4) established for protonation of 2^- ,³² the protonation of 1H to 1HHy^+ is hence one to two orders of magnitude faster than expected for protonation of its pdt analogue 2^- with the same driving force. Intrinsically faster protonation of the adt-protonated 1H would prevent this step from becoming rate limiting when reducing weaker acids. Below, we discuss the possible origin of this favourable effect.

Implications for (electro)catalytic H_2 formation

From previous electrochemical experiments comparing 1 and 2 it has been concluded that catalytic H_2 formation from the same acids proceeds with similar second order catalytic rate constants despite the lower overpotential arising from the easier reduction of adt-protonated catalyst.²⁷ Undiminished reactivity towards protonation despite the lowered driving force would imply that 1 breaks the usual scaling relation between overpotential and rate,³⁸ an effect that has been attributed to the proton shuttling role of the adt²⁷ ligand. By putting an upper limit of 1 s^{-1} on the first order rate constant for intramolecular

tautomerization, our results exclude however that a mechanism involving hydride formation by proton shuttling at the $\text{Fe}_2(0,\text{l})$ redox level could under any conditions result in the reported pseudo-first order catalytic rate constant on the order of 10^3 to 10^4 s^{-1} , as previously proposed.²⁷ It is important to note that the reported kinetics refer to the first catalytic wave (-1.4 V) that has been attributed to mechanisms where the hydride is formed prior to the second electron transfer, *i.e.* a CECE sequence with stronger acids or an ECCE sequence with weak acids. Therefore, possible formation of the hydride *via* tautomerization at the $\text{Fe}_2(0,0)$ redox level ($1\text{H}^- \rightarrow 1\text{Hy}^-$) cannot explain the superior catalytic performance of the adt complex.

An alternative advantage of the adt ligand might instead arise from the lowered barrier for hydride formation due to the structural changes at the iron core induced by the preceding ligand protonation. This notion would be in line with the relative sluggishness of the latter reaction (see above) and is corroborated by structural data that has been previously obtained from EXAFS spectra³⁹ of ligand- and metal-protonated states of the related $\text{Fe}_2(\text{l},\text{l})$ complex $4'$.³⁷ Specifically, protonation of the adt ligand in $4'$ results in an elongated Fe–Fe bond distance in $4'\text{H}^+$ that is close to the distance found in the bridging hydride $4'\text{HHy}^{2+}$. Formation of the hydride from the ligand protonated complex requires hence much less reorganization of this coordinate than formation of hydride $4'\text{Hy}^+$ from $4'$.

Generally, a proton shuttling ligand could accelerate not only the formation of the hydride intermediate but also its subsequent coupling with a proton. In case of adt complex 1 , with its bridging hydride intermediate, neither of the two steps is, however, likely to involve the adt-N, for steric reasons. In contrast to the terminal hydride of the enzyme active site, the hydride is bridging in the model complex (*cf.* Scheme 1), as evidenced by the conserved symmetry shown by the IR spectra, and the absence of a bridging CO IR band. The superior catalytic performance of 1 over its pdt analogue 2 is instead attributed to the observed acceleration of hydride formation with bulk acid as described above.

Conclusions

In summary, we could demonstrate by real-time spectroscopic observation that the initial protonation of hydrogenase model complex 1 , also in its one-electron reduced $\text{Fe}_2(\text{l},0)$ state 1^- , occurs on the adt ligand rather than the Fe_2 core. The same ligand protonated intermediate 1H was alternatively generated by one-electron reduction of 1H^+ and no evidence for tautomerization of 1H to a potentially more stable hydride 1Hy was observed in either case. 1H was stable on the time scale of 1 s , which shows that intramolecular proton shuttling cannot be responsible for the reported catalytic TOFs of 10^3 to 10^4 s^{-1} .²⁷ For catalytic H_2 generation from weak acids this implies that no hydride intermediate is formed on the $\text{Fe}_2(\text{l},0)$ level; instead, hydride formation may occur after further reduction, by direct protonation in the $\text{Fe}_2(0,0)$ state. Only with strong acids a doubly protonated intermediate 1HHy^+ is formed in the $\text{Fe}_2(\text{l},0)$ state, by direct metal protonation of 1H , not by ligand



protonation of a hydride precursor. Generally, it can therefore be concluded that, in contrast to the enzymatic reaction, the adt ligand of model complex **1** has no proton relay function that enables rapid formation of the bridging hydride intermediates on the Fe₂(μ ,0) level. The protonation kinetics reveal however an exceptionally high barrier for protonation of the adt ligand that we suggest in turn may lower the barrier for hydride formation, presumably due to rearrangements of the metal core upon ligand protonation. In this way the basic site in the second coordination sphere might allow the catalyst to overcome the usual trade-off between overpotential and rate without invoking any proton-shuttling role.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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Notes and references

† Protonation kinetics of the phosphine complexes **4** (ref. 29) and **4'** (ref. 37) have so far been studied only in the Fe₂(μ , μ) state where hydride formation is extremely slow.

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