



Cite this: *RSC Adv.*, 2020, 10, 32069

Heterodinuclear nickel(II)–iron(II) azadithiolates as structural and functional models for the active site of [NiFe]-hydrogenases†

Li-Cheng Song, ^{ab} Bei-Bei Liu,^a Wen-Bo Liu^a and Zheng-Lei Tan^a

To develop the biomimetic chemistry of [NiFe]-H₂ases, the first azadithiolato-bridged NiFe model complexes [CpNi((μ-SCH₂)₂NR)Fe(CO)(diphos)]BF₄ (5, R = Ph, diphos = dppv; 6, 4-ClC₆H₄, dppv; 7, 4-MeC₆H₄, dppv; 8, CO₂CH₂Ph, dppe; 9, H, dppe) have been synthesized *via* well-designed synthetic routes. Thus, treatment of RN(CH₂S(O)CMe)₂ with *t*-BuONa followed by reaction of the resulting intermediates RN(CH₂SNa)₂ with (dppv)Fe(CO)₂Cl₂ or (dppe)Fe(CO)₂Cl₂ gave the N-substituted azadithiolato-chelated Fe complexes [RN(CH₂S)₂]Fe(CO)₂(diphos) (1, R = Ph, diphos = dppv; 2, 4-ClC₆H₄, dppv; 3, 4-MeC₆H₄, dppv; 4, CO₂CH₂Ph, dppe). Further treatment of 1–4 with nickelocene in the presence of HBF₄·Et₂O afforded the corresponding N-substituted azadithiolato-bridged NiFe model complexes 5–8, while treatment of 8 with HBF₄·Et₂O resulted in formation of the parent azadithiolato-bridged model complex 9. While all the new complexes 1–9 were characterized by elemental analysis and spectroscopy, the molecular structures of model complexes 6–8 were confirmed by X-ray crystallographic study. In addition, model complexes 7 and 9 were found to be catalysts for H₂ production with moderate *i*_{cat}/*i*_p and overpotential values from TFA under CV conditions.

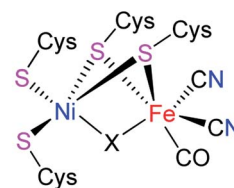
Received 15th May 2020
Accepted 20th August 2020
DOI: 10.1039/d0ra04344c
rsc.li/rsc-advances

Introduction

[NiFe]-hydrogenases ([NiFe]-H₂ases) are a class of biological enzymes that catalyze the reversible redox reaction between molecular H₂ and protons in a variety of microorganisms such as bacteria, archaea, and cyanobacteria.^{1–3} An X-ray crystallographic study revealed that the active site of [NiFe]-H₂ases features a Ni(tetracysteinate) moiety, in which the two cysteine atoms are connected to the Fe atom of an Fe(CN)₂(CO) moiety. In addition, the oxidized state of [NiFe]-H₂ases includes an additional bridging oxygenous ligand, while the reduced state of [NiFe]-H₂ases contains an additional bridging hydride ligand or a Ni–Fe metal–metal bond (Fig. 1).^{4–6}

In recent years, molecular H₂ has been considered as an ideal energy source to replace fossil fuels to solve the current energy shortage and environmental pollution problems. This is because dihydrogen is a high energy-dense, clean-burning, and sustainable energy.^{7–9} In addition, although H₂ production can be carried out in an industry scale from fossil fuel by petroleum

reforming, the utilized catalyst is the expensive and relatively scarce noble metal Pt. To develop the biomimetic chemistry of [NiFe]-H₂ases and to search for the water splitting H₂-producing catalysts that contain the earth-abundant and cheap metals such as iron and nickel present in the active site of [NiFe]-H₂ases, synthetic chemists have prepared a wide variety of biomimetic models for [NiFe]-H₂ases.^{9–29} However, among the prepared models no one bears a bridging azadithiolato ligand between their NiFe centers. In view of the important catalytic role played by the central N atom of the bridging azadithiolato ligand in the active site of [FeFe]-H₂ases,^{30–32} we recently launched a study on the preparation of the first azadithiolato ligand-containing [NiFe]-H₂ase models. Fortunately, we have successfully prepared and characterized such a type of azadithiolato-bridged models in which each contains an azadithiolato RN(CH₂S)₂ ligand bridged between their CpNi and (diphos)(CO)Fe units (Fig. 2). In addition, the two representative



X = O²⁻, HO⁻, H₂O, or H⁻

Fig. 1 Active site of [NiFe]-H₂ases.

^aDepartment of Chemistry, State Key Laboratory of Elemento-Organic Chemistry, College of Chemistry, Nankai University, Tianjin 300071, China. E-mail: lcsong@nankai.edu.cn

^bCollaborative Innovation Center of Chemical Science and Engineering (Tianjin), Tianjin 300072, China

† Electronic supplementary information (ESI) available: Some electrocatalytic plots/data (Fig. S1–S3/Table S1) and some IR/NMR spectra (Fig. S4–S23). CCDC 2001290–2001292. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ra04344c



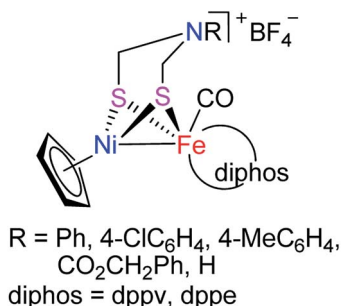


Fig. 2 Models of [NiFe]-H₂ases reported in this article.

models have been found to be catalysts for proton reduction to hydrogen under electrochemical conditions. Herein, we report the interesting results obtained by this study.

Results and discussion

Synthesis and characterization of precursor complexes

[RN(CH₂S)₂Fe(CO)₂(diphos)] (1–4)

The N-substituted azadithiolato ligand-chelated mononuclear Fe complexes 1–4 utilized for the preparation of the targeted [NiFe]-H₂ase models could be prepared by one-pot reactions of the bis(thioester) compounds RN[CH₂S(O)CMe]₂ (R = Ph, 4-ClC₆H₄, 4-MeC₆H₄, CO₂CH₂Ph) with *tert*-butoxysodium in THF at –78 °C followed by treatment of the resulting disodium mercaptides RN(CH₂SNa)₂ with (dppv)Fe(CO)₂Cl₂ or (dppe)Fe(CO)₂Cl₂ generated *in situ* by reaction of an acetone solution of FeCl₂ with the corresponding diphosphine and bubbling CO gas^{33,34} in 18–35% yields (Scheme 1).

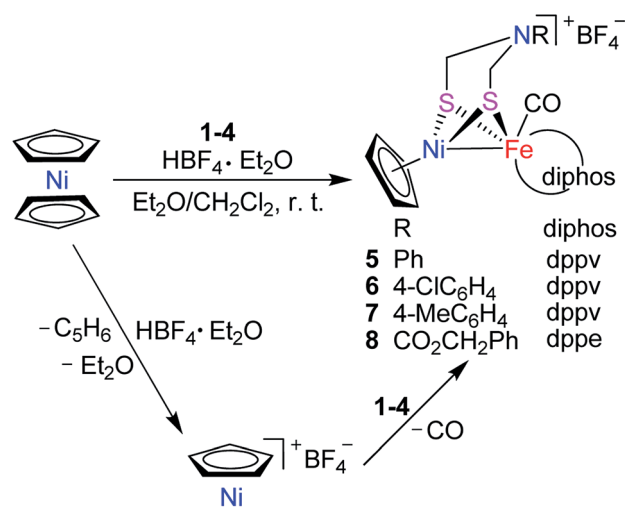
Precursor complexes 1–4 are air-stable, brown-red solids and have been characterized by elemental analysis and various spectroscopic methods. For example, the ¹H NMR spectra of 1–3 each displayed one singlet in the region 4.56–4.63 ppm for the four H atoms in their azadithiolato methylene groups, whereas the ¹H NMR spectrum of 4 displayed a multiplet in the range 4.89–5.30 ppm for the four H atoms in its azadithiolato methylene groups. The ¹³C{¹H} NMR spectra of 1–4 exhibited one or two singlets in the range 200–214 ppm for their two terminal carbonyl C atoms, whereas the ³¹P{¹H} NMR spectra of 1–3 displayed one singlet at about 79 ppm for the two P atoms in their dppv ligands and the ³¹P{¹H} NMR spectrum of 4 exhibited one singlet at 44.3 ppm for the two P atoms in its dppe ligand. In

addition, the IR spectra of 1–4 showed two absorption bands in the range 2035–1968 cm^{–1} for their two terminal carbonyls.

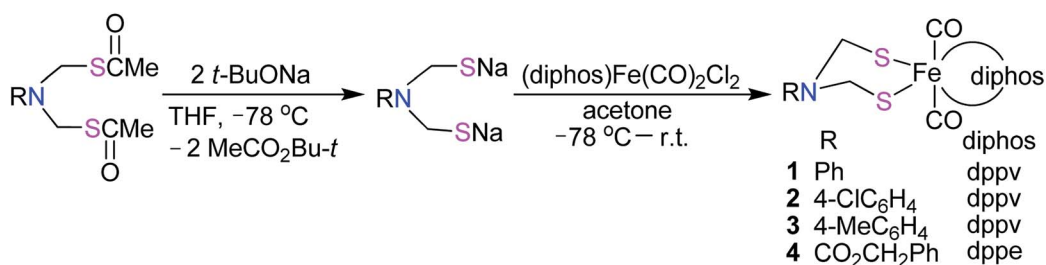
Synthesis and characterization of targeted [NiFe]-H₂ase models [CpNi{(μ-SCH₂)₂NR}Fe(CO)(diphos)]BF₄ (5–9)

Interestingly, the targeted N-substituted dithiolato-bridged NiFe-based models 5–8 were found to be prepared by treatment of the mononuclear Fe complexes 1–4 with nickelocene/HBF₄·Et₂O in a mixed solvent CH₂Cl₂/Et₂O at room temperature in 21%–62% yields (Scheme 2). Further treatment of the N-substituted dithiolato-bridged model complex 8 with HBF₄·Et₂O in a mixed solvent CH₂Cl₂/PhOMe at room temperature gave rise to the parent azadithiolato-bridged model 9 in 42% yield (Scheme 3). As shown in Schemes 2 and 3, while model 9 can be formally viewed as produced through the selective acidolysis of the N–C(=O) bond rather than O–C(=O) bond of model 8, model complexes 5–8 might be regarded as produced *via* two elementary reaction steps. The first step involves protonation of one anionic cyclopentadienyl ligand in Cp₂Ni to give cyclopentadiene and cyclopentadienylnickel tetrafluoroborate.^{35,36} The second step involves coordination of the two S atoms in complexes 1–4 to Ni atom of the *in situ* formed intermediate CpNi(BF₄) followed by loss of one CO ligand from 1–4 to give the final products.

Model complexes 5–9 are also air-stable, brown-red solids. They were characterized by the IR, ¹H NMR, ¹³C{¹H} NMR, and

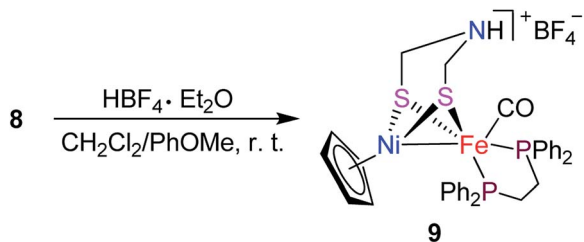


Scheme 2 Synthesis of targeted models 5–8.



Scheme 1 Synthesis of precursor complexes 1–4.





Scheme 3 Synthesis of targeted model 9.

$^{31}\text{P}\{^1\text{H}\}$ NMR spectroscopic techniques and elemental analysis. For instance, the IR spectra of 5–9 showed one strong absorption band in the region $1956\text{--}1939\text{ cm}^{-1}$ for their terminal carbonyls, which are close to that (1944 cm^{-1}) displayed by the $[\text{NiFe}]\text{-H}_2\text{ase}$ from *D. Vulgaris* Miyazaki F.³⁷ The ^1H NMR spectra of 5–9 displayed one singlet in the range 4.42–4.61 ppm for the five H atoms in their Cp rings. In addition, the $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of 5–9 exhibited one singlet in the region 206–220 ppm for their terminal carbonyl C atoms, whereas the $^{31}\text{P}\{^1\text{H}\}$ NMR spectra of 5–7 displayed one singlet in the region 82.7–83.9 ppm for the two P atoms in their dppv ligands and those of 8 and 9 exhibited the corresponding signals in the range 78.1–79.1 ppm for the two P atoms in their dppe ligands.

The molecular structures of the azadithiolato-bridged complexes 6–8 were confirmed by X-ray crystallographic analysis. While Fig. 3–5 show their molecular structures, Table 1 lists their selected bond lengths and angles. As shown in Fig. 3–5, the three biomimetic models are indeed the azadithiolato ligand-containing complexes that are composed of one monocation $[\text{CpNi}\{(\mu\text{-SCH}_2)_2\text{NC}_6\text{H}_4\text{Cl-4}\}\text{Fe}(\text{CO})(\text{dppv})]^+$, $[\text{CpNi}\{(\mu\text{-SCH}_2)_2\text{NC}_6\text{H}_4\text{Me-4}\}\text{Fe}(\text{CO})(\text{dppv})]^+$, or $[\text{CpNi}\{(\mu\text{-SCH}_2)_2\text{-NCO}_2\text{CH}_2\text{Ph}\}\text{Fe}(\text{CO})(\text{dppe})]^+$ and one monoanion BF_4^- . Since the molecular structures of 6 and 7 are virtually the same as that of 8 (except that the $4\text{-ClC}_6\text{H}_4/\text{dppv}$ in 6 and $4\text{-MeC}_6\text{H}_4/\text{dppv}$ in 7

Table 1 Selected bond lengths (Å) and angles (°) for 6–8

6			
Ni(1)–Fe(1)	2.5033(9)	Fe(1)–S(1)	2.2472(12)
Ni(1)–S(1)	2.1783(13)	Fe(1)–S(2)	2.2361(11)
Ni(1)–S(2)	2.1770(11)	Fe(1)–P(1)	2.2198(11)
N(1)–C(28)	1.433(5)	Fe(1)–P(2)	2.2215(12)
Ni(1)–S(1)–Fe(1)	68.87(4)	S(2)–Fe(1)–S(1)	87.09(4)
Ni(1)–S(2)–Fe(1)	69.10(4)	S(2)–Fe(1)–Ni(1)	54.34(3)
S(2)–Ni(1)–S(1)	90.34(4)	S(1)–Fe(1)–Ni(1)	54.26(3)
P(1)–Fe(1)–P(2)	86.10(4)	S(1)–Ni(1)–Fe(1)	56.86(4)
7			
Ni(1)–Fe(1)	2.5071(9)	Fe(1)–S(1)	2.2497(14)
Ni(1)–S(1)	2.1826(16)	Fe(1)–S(2)	2.2374(13)
Ni(1)–S(2)	2.1795(14)	Fe(1)–P(1)	2.2268(15)
N(1)–C(33)	1.429(7)	Fe(1)–P(2)	2.2249(14)
Ni(1)–S(1)–Fe(1)	68.88(5)	S(2)–Fe(1)–S(1)	87.15(5)
Ni(1)–S(2)–Fe(1)	69.15(4)	S(2)–Fe(1)–Ni(1)	54.33(4)
S(2)–Ni(1)–S(1)	90.31(5)	S(1)–Fe(1)–Ni(1)	54.30(4)
P(1)–Fe(1)–P(2)	86.13(5)	S(1)–Ni(1)–Fe(1)	56.83(4)
8			
Ni(1)–Fe(1)	2.5036(9)	Fe(1)–S(1)	2.2367(11)
Ni(1)–S(1)	2.1807(13)	Fe(1)–S(2)	2.2306(12)
Ni(1)–S(2)	2.1819(12)	Fe(1)–P(1)	2.2433(12)
N(1)–C(34)	1.434(5)	Fe(1)–P(2)	2.2242(11)
Ni(1)–S(1)–Fe(1)	69.04(4)	S(2)–Fe(1)–S(1)	87.28(4)
Ni(1)–S(2)–Fe(1)	69.13(4)	S(2)–Fe(1)–Ni(1)	54.52(4)
S(1)–Ni(1)–S(2)	89.93(4)	S(1)–Fe(1)–Ni(1)	54.43(3)
P(1)–Fe(1)–P(2)	87.08(4)	S(1)–Ni(1)–Fe(1)	56.54(4)

are replaced by the $\text{CO}_2\text{CH}_2\text{Ph}/\text{dppe}$ in 8), we just discuss the structure of 8. In the monocation of 8, the Ni–Fe distance (2.5036 Å) is shorter than that (2.57 Å) of the $[\text{NiFe}]\text{-H}_2\text{ase}$ from *D. Vulgaris* Miyazaki F.³⁸ and the sum (2.56 Å) of the Ni and Fe covalent radii,³⁹ indicative of the presence of a Ni–Fe metal–metal bond. In addition, the Ni–Fe distance of 8 is very close to

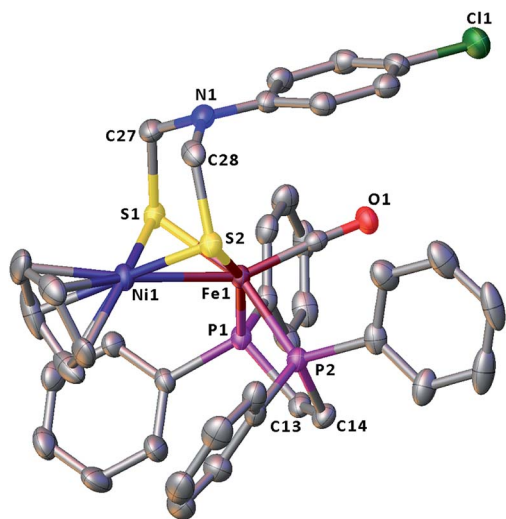


Fig. 3 Molecular structure of model complex 6 with ellipsoids drawn at a 50% probability level. All hydrogen atoms and the anion BF_4^- are omitted for clarity.

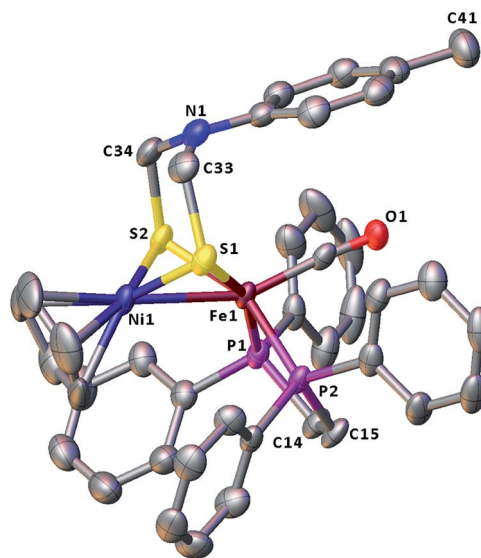


Fig. 4 Molecular structure of model complex 7 with ellipsoids drawn at a 50% probability level. All hydrogen atoms and the anion BF_4^- are omitted for clarity.



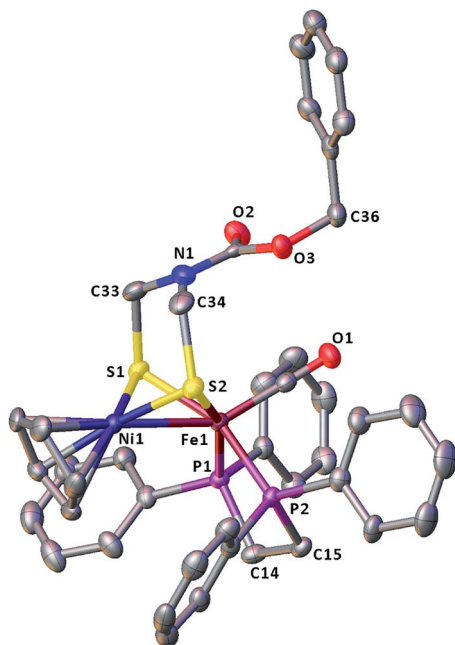


Fig. 5 Molecular structure of model complex **8** with ellipsoids drawn at a 50% probability level. All hydrogen atoms and the anion BF_4^- are omitted for clarity.

that (2.5145 Å) of the reported 1,3-propanedithiolato(pdt)-bridged analogue $[\text{CpNi}(\text{pdt})\text{Fe}(\text{dppe})(\text{CO})]\text{BF}_4$.⁴⁰ Owing to the presence of a Ni–Fe metal–metal bond in **8**, the Ni center could be regarded as taking a “piano stool” geometry and the Fe center adopts a pseudo-octahedral geometry. In addition, the azadithiolato ligand in **8** is bridged between Ni and Fe atoms to construct two fused six-membered rings. The six-membered ring Fe1S1C33N1C34S2 shown in Fig. 3 possesses a boat conformation, whereas the six-membered ring Ni1S1C33N1C34S2 possesses a chair conformation. While the N-substituted $\text{CO}_2\text{CH}_2\text{Ph}$ group is attached to the common N1 atom of the two fused six-membered rings with an axial type of bond, the terminal carbonyl ligand is bound to Fe1 atom in an axial position of the pseudo-octahedral geometry of the Fe center.

Electrochemical properties and electrocatalytic H_2 production catalyzed by model complexes **7** and **9**

Although the electrochemical properties and electrocatalytic H_2 production for some $[\text{NiFe}]\text{-H}_2\text{ase}$ models were previously

Table 2 Electrochemical data of **7/9**^a and **A/B**^b

Complex	$E_{\text{pc}1}/\text{V}$	$E_{\text{pc}2}/\text{V}$	E_{pa}/V
7	−1.07	−1.98	0.54
9	−1.15	−1.99	0.45
A ⁴⁰	−1.15	−2.18	0.72
B ⁴⁰	−1.16	−2.15	0.70

^a All potentials are versus Fc/Fc^+ in 0.1 M $n\text{-Bu}_4\text{NPF}_6/\text{MeCN}$ at a scan rate of 0.1 V s^{-1} . ^b All potentials are versus Fc/Fc^+ in 0.1 M $n\text{-Bu}_4\text{NPF}_6/\text{CH}_2\text{Cl}_2$ at a scan rate of 0.5 V s^{-1} .

studied by cyclic voltammetric techniques,^{40–46} there isn't any studied result regarding the electrochemical and electrocatalytic properties of the azadithiolato-bridged $[\text{NiFe}]\text{-H}_2\text{ase}$ models, so far. In order to see whether the prepared azadithiolato-bridged models **5–9** could have the electrocatalytic H_2 -producing ability, we chose models **7** and **9** (note that they have the same type of structures as **5**, **6** and **8**) as representatives to study their electrochemical properties by CV techniques using the supporting electrolyte $n\text{-Bu}_4\text{NPF}_6$ in MeCN at a scan rate of 100 mVs^{-1} . As shown in Table 2 and Fig. 6, complex **7** displayed one quasi-reversible reduction wave at -1.07 V (due to its $|E_{\text{pc}} - E_{\text{pa}}| = 82 \text{ mV}$ and $i_{\text{pc}}/i_{\text{pa}} = 1.05$),⁴⁷ one irreversible reduction wave at -1.98 V and one irreversible oxidation wave at 0.54 V , while complex **9** displayed one quasi-reversible reduction wave at -1.15 V (due to its $|E_{\text{pc}} - E_{\text{pa}}| = 103 \text{ mV}$ and $i_{\text{pc}}/i_{\text{pa}} = 1.33$),⁴⁷ one irreversible reduction wave at -1.99 V , and one irreversible oxidation wave at 0.45 V , respectively. Previously, the electrochemical properties of the two pdt-bridged analogues of **7** and **9**, namely complexes $[\text{CpNi}(\text{pdt})\text{Fe}(\text{dppv})(\text{CO})]\text{BF}_4$ (**A**) and $[\text{CpNi}(\text{pdt})\text{Fe}(\text{dppe})(\text{CO})]\text{BF}_4$ (**B**) (see Table 2) were studied by cyclic voltammetry.⁴⁰ Similar to this case, the three redox events of **7** and **9** could be assigned to the $\text{Ni}^{\text{II}}\text{Fe}^{\text{II}}/\text{Ni}^{\text{I}}\text{Fe}^{\text{II}}$, $\text{Ni}^{\text{I}}\text{Fe}^{\text{II}}/\text{Ni}^{\text{I}}\text{Fe}^{\text{I}}$ and $\text{Ni}^{\text{II}}\text{Fe}^{\text{II}}/\text{Ni}^{\text{III}}\text{Fe}^{\text{II}}$ couples, respectively.

Having determined the electrochemical properties of **7** and **9**, we further determined their cyclic voltammograms in the presence and absence (for comparison) of trifluoroacetic acid (TFA) ($\text{p}K_{\text{a}}^{\text{MeCN}} = 12.7$, $E_{\text{HA}}^{\circ} = -0.89 \text{ V}$),^{48,49} in order to see whether they could catalyze the proton reduction of TFA to give H_2 . As shown in Fig. 7 and 8, when TFA was sequentially added from 2 to 10 mM to MeCN solutions of **7** and **9**, the current heights of their original first reduction waves were basically unchanged. However, in contrast to this, the current heights of their original second reduction waves were remarkably increased. Apparently, such an observation features an electrocatalytic proton reduction process from TFA to hydrogen.^{41,43} Note that through comparison of the reduction potentials E_{pc} and peak currents i_{p} in the CVs of **7/9** with those in the CVs of TFA without **7/9** (see Fig. S1 and Table S1 in the ESI†), models **7/9**

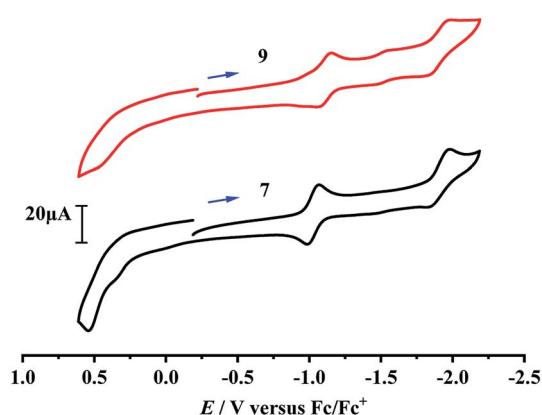


Fig. 6 Cyclic voltammograms of **7** and **9** (1.0 mM) in 0.1 M $n\text{-Bu}_4\text{NPF}_6/\text{MeCN}$ at a scan rate of 0.1 V s^{-1} . Arrows indicate the starting potential and scan direction.



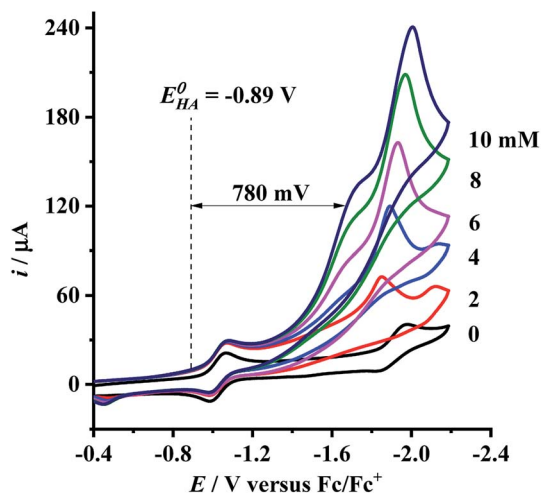


Fig. 7 Cyclic voltammograms of **7** (1.0 mM) with TFA (0–10 mM) in 0.1 M *n*-Bu₄NPF₆/MeCN at a scan rate of 0.1 V s⁻¹.

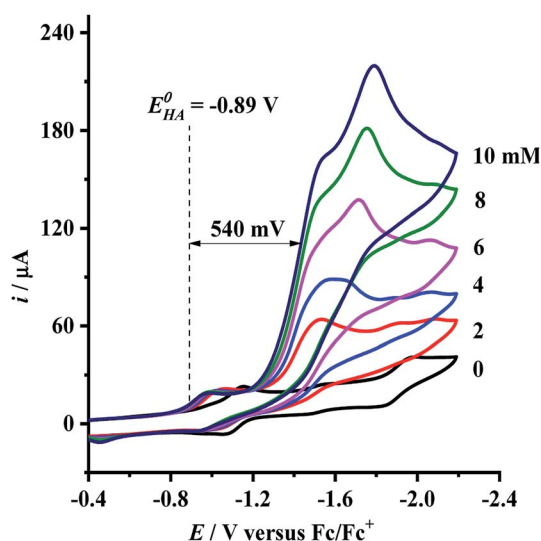


Fig. 8 Cyclic voltammograms of **9** (1.0 mM) with TFA (0–10 mM) in 0.1 M *n*-Bu₄NPF₆/MeCN at a scan rate of 0.1 V s⁻¹.

9 were further proved to be catalysts for proton reduction. In addition, during the proton reductions catalyzed by **7** and **9**, the original second reduction waves were positively shifted by *ca.* 15 and 45 mV, respectively; this is most likely due to protonation of the central N atoms in their bridged azadithiolato ligands.⁵⁰

Recently, we⁵¹ and others⁵² reported that some mononuclear Fe complexes can serve as catalysts for proton reduction to hydrogen. In order to observe whether the mononuclear Fe precursors of model complexes **5**–**8** (namely complexes **1**–**4**) could catalyze the proton reductions, we chose complexes **3** and **4** as representatives to determine their cyclic voltammograms in the presence of TFA under the same conditions as those used for determination of models **7** and **9**. We found that when TFA was sequentially added from 2 to 10 mM to MeCN solutions of **3** and **4**, the current heights of the acid-induced reduction waves were considerably increased near their only one original

reduction waves caused by their Fe(II)Fe(I) couples (see Fig. S2 and S3 in the ESI†). Obviously, such an observation demonstrates an electrocatalytic proton reduction process catalyzed by complexes **3** and **4**. So, we might conclude that for mononuclear Fe complexes **3** and **4**, only Fe makes contribution to their electrocatalytic proton reduction reactions caused by their Fe(II)/Fe(I) couples, while for dinuclear models **7** and **9**, both Ni and Fe make contribution to the electrocatalytic proton reduction reactions occurred near their original second reduction waves caused by their Ni(I)Fe(II)/Ni(I)Fe(I) couples.

To evaluate the electrocatalytic H₂-producing ability of **7** and **9**, we determined the ratio of the catalytic current (*i*_{cat}) to reductive peak current (*i*_p) in the absence of the added acid (note that a higher *i*_{cat}/*i*_p value indicates a faster catalysis).⁵³ At a concentration of 10 mM TFA, the *i*_{cat}/*i*_p values of **7** and **9** were calculated to be 6.0 and 5.3, respectively. It follows that the H₂-producing ability catalyzed by complex **7** (with a 4-MeC₆H₄N group and a dppv ligand) is higher than that catalyzed by **9** (with an NH group and a dppe ligand). However, in order to compare the electrocatalytic H₂-producing ability of the bridgehead N-substituted complex **8** with its parent complex **9**, we further determined its *i*_{cat}/*i*_p value by CV techniques to be 6.4 under the same conditions. Therefore, the H₂-producing ability of complex **8** with a stronger electron-attracting PhCH₂CO₂N group is higher than that of complex **9** with a weaker electron-attracting NH group. In addition, at a 10 mM TFA concentration, the *i*_{cat}/*i*_p values (5.3–6.4) of complexes **7**–**9** are higher than those (*ca.* 4) of the previously reported analogues [HNiFe(pdt)(dppe)(L)(CO)₂]₂BF₄ (L = P(OPh)₃, PPh₃).⁵⁴

Since overpotential is also an important parameter for evaluating the electrocatalytic ability for a catalyst (note that a better catalyst has a lower overpotential), we used Evans' method to calculate the overpotentials of **7** and **9** according to the following two equations:^{49,55}

$$\text{Overpotential} = \left| E_{\text{cat}/2}^{\circ} - E_{\text{HA}}^{\circ} \right| \quad (1)$$

$$E_{\text{HA}}^{\circ} = E_{\text{H}^{+}/\text{H}_2}^{\circ} - \left(\frac{2.303RT}{F} \right) \text{p}K_{\text{a,HA}} \quad (2)$$

where *E*_{cat/2} is the catalytic half-wave potential and *E*_{HA}[°] is the standard potential of the acid; *R* is the ideal gas constant 8.314 J mol⁻¹ K⁻¹; *T* is the absolute temperature 298 K; *F* is the Faraday's constant 9.65 × 10⁴ C mol⁻¹; in addition, the p*K*_a value of TFA in MeCN and the standard redox potential *E*_{H⁺/H₂}[°] have been previously determined to be 12.7 (ref. 48) and -0.14 V,⁴⁹ respectively. According to eqn (2), the standard potential *E*_{HA}[°] can be calculated to be -0.89 V and then according to eqn (1), the overpotentials of **7** and **9** can be calculated to be 0.78 and 0.54 V, respectively (see Fig. 7 and 8). At a 10 mM acid concentration, overpotentials of our azadithiolato-bridged dinuclear NiFe complexes **7** (0.78 V) and **9** (0.54 V) by using TFA as proton source are higher than those (*ca.* 0.5 V) of the previously reported analogues [HNiFe(pdt)(dppe)(L)(CO)₂]₂BF₄ (L = P(OPh)₃, PPh₃) using TFA as proton source.⁵⁴ Particularly worth noting is that in terms of the studied results regarding the structure and catalytic function described above, complexes **7**



Table 3 Crystal data and structure refinement details for 6–8

	6	7	8
Formula	C ₄₀ H ₃₅ BClF ₄ FeNNiO	C ₄₁ H ₃₈ BF ₄ FeNNiO	C ₄₂ H ₄₀ BF ₄ FeNNiO ₃
<i>M_w</i>	P ₂ S ₂	P ₂ S ₂ ·CH ₂ Cl ₂	P ₂ S ₂ ·CH ₂ Cl ₂
Cryst syst	908.57	973.08	1019.10
Space group	Monoclinic	Monoclinic	Triclinic
<i>a</i> (Å)	<i>P</i> 121/ <i>c</i> 1	<i>P</i> 121/ <i>c</i> 1	<i>P</i> 1̄
<i>b</i> (Å)	17.861(4)	17.8793(11)	10.113(2)
<i>c</i> (Å)	11.063(2)	11.0429(5)	12.365(3)
<i>α</i> (°)	22.252(5)	22.4895(10)	18.383(4)
<i>β</i> (°)	90	90	75.55(3)
<i>γ</i> (°)	108.15(3)	108.615(6)	87.96(3)
<i>V</i> (Å ³)	90	90	77.76(3)
<i>Z</i>	4178.2(16)	4208.0(4)	2175.1(8)
Crystal size (mm)	4	4	2
<i>D_c</i> (g cm ^{−3})	0.05 × 0.04 × 0.03	0.06 × 0.05 × 0.03	0.20 × 0.18 × 0.12
<i>μ</i> (mm ^{−1})	1.444	1.536	1.556
<i>F</i> (000)	1.088	1.148	1.117
Reflns collected	1856	1992	1044
Reflns unique	49 278	43 090	26 271
<i>θ</i> _{min} / <i>θ</i> _{max} (°)	9958	8586	10 291
Final <i>R</i>	2.078/27.863	1.905/26.372	1.144/27.834
Final <i>R_w</i>	0.0693	0.0712	0.0567
GOF on <i>F</i> ²	0.1648	0.1987	0.1575
<i>Δρ</i> _{max} / _{min} (e Å ^{−3})	1.127	1.061	1.060
	0.536/−0.576	1.000/−1.039	0.738/−1.255

and **9** might be regarded as not only the structural models but also the functional models for the active site of [NiFe]-H₂ases.

Summary and conclusion

We have synthesized the first azadithiolato-bridged dinuclear NiFe model complexes **5–8** and **9** by reactions of the N-substituted azadithiolato-chelated mononuclear Fe complexes **1–4** with Cp₂Ni/HBF₄·Et₂O and by reaction of the N-substituted azadithiolato-bridged complex **8** with HBF₄·Et₂O, respectively. Particularly worth noting is that the spectroscopic characterization of all the new model complexes **5–9** and the X-ray crystallographic study on their three representative complexes **6–8** have proved that models **5–9** indeed contain an azadithiolato ligand that is bridged between their NiFe centers to form a butterfly Ni(II)₂S₂Fe(II) cluster core with a Ni–Fe metal–metal bond. In addition, from the electrochemical and electrocatalytic studies, the two representative models **7** and **9** have been found to be catalysts for proton reduction to hydrogen using TFA as the proton source under electrochemical conditions. Therefore, according to the structural and functional similarity with the active site of [NiFe]-H₂ases, complexes **5–9** might be regarded as the structural and functional models of [NiFe]-H₂ases.

Experimental section

General comments

All reactions were carried out using standard Schlenk and vacuum-line techniques under an atmosphere of argon. While tetrahydrofuran (THF) and Et₂O were distilled under Ar from sodium/benzophenone ketyl, CH₂Cl₂ was distilled from CaH₂. *t*-BuONa, (η⁵-C₅H₅)₂Ni, HBF₄·Et₂O (50–55% w/w in Et₂O),

anhydrous FeCl₂ and other chemicals were commercially available and used as received. RN[CH₂S(O)CMe]₂ (R = Ph,⁵⁶ 4-ClC₆H₄,⁵⁷ CO₂CH₂Ph,⁵⁸ (dppv)Fe(CO)₂Cl₂ (dppv = 1,2-bis(diphenylphosphino)ethene)³⁵ and (dppe)Fe(CO)₂Cl₂ (dppe = 1,2-bis(diphenylphosphino)ethane)³⁶) were prepared according to the published procedures. While ¹H, ¹³C{¹H}, and ³¹P{¹H} NMR spectra were obtained on a Bruker Avance 400 NMR spectrometer, IR spectra were recorded on a Bio-Rad FTS 135 infrared spectrophotometer. Elemental analyses were performed on an Elementar Vario EL analyzer. Melting points were determined on a SGW X-4 microscopic melting point apparatus and were uncorrected.

Preparation of [PhN(CH₂S)₂]Fe(CO)₂(dppv) (1**).** A 250 mL Schlenk flask was charged with FeCl₂ (0.634 g, 5.0 mmol) in 100 mL acetone and dppv (1.98 g, 5.0 mmol) in 20 mL of THF. The mixture was stirred at room temperature and bubbled with CO gas for 4 h to give a (dppv)Fe(CO)₂Cl₂-containing brown-red solution. Additionally, a 100 mL Schlenk flask was charged with thioester PhN[CH₂S(O)CMe]₂ (2.02 g, 7.5 mmol), sodium *tert*-butoxide (1.44 g, 15.0 mmol), and 40 mL of THF. The mixture was stirred at −78 °C for 4 h to give a PhN(CH₂SNa)₂-containing pale yellow solution. To the *in situ* prepared (dppv)Fe(CO)₂Cl₂-containing solution cooled to −78 °C was slowly added the prepared PhN(CH₂SNa)₂-containing solution and then the mixture was stirred at −78 °C for 5 h. Solvent was removed at reduced pressure and the residue was subjected to column chromatography (silica gel). Elution with CH₂Cl₂ developed a red band, from which **1** (0.622 g, 18%) was obtained as a brown-red solid, mp 93–94 °C. Anal. calcd for C₃₆H₃₁FeNO₂·P₂S₂: C, 62.52; H, 4.52; N, 2.03. Found: C, 62.35; H, 4.65; N, 2.00. IR (KBr disk): ν_{C≡O} 2018 (w), 1973 (vs.) cm^{−1}. ¹H NMR (400



MHz, CDCl₃): 4.63 (s, 4H, CH₂NCH₂), 6.69–7.91 (m, 27H, CH=CH, 5C₆H₅) ppm. ¹³C{¹H} NMR (100 MHz, CDCl₃): 48.6 (s, CH₂NCH₂), 114.7–134.7 (m, C₆H₅), 146.3–150.8 (m, CH=CH), 211.8, 212.2 (2s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 79.9 (s, FeP) ppm.

Preparation of [4-ClC₆H₄N(CH₂S)₂]Fe(CO)₂(dppv) (2). The same procedure was followed as for the preparation of **1**, except that 4-ClC₆H₄N[CH₂S(O)CMe]₂ (2.28 g, 7.5 mmol) was used instead of PhN[CH₂S(O)CMe]₂. **2** (0.835 g, 23%) was obtained as a brown-red solid, mp 145–146 °C. Anal. calcd for C₃₆H₃₀ClFeNO₂P₂S₂: C, 59.56; H, 4.17; N, 1.93. Found: C, 59.47; H, 4.21; N, 1.69. IR (KBr disk): ν_{C=O} 2020 (w), 1973 (vs.) cm⁻¹. ¹H NMR (400 MHz, CDCl₃): 4.57 (s, 4H, CH₂NCH₂), 6.61–7.78 (m, 26H, CH=CH, 4C₆H₅, C₆H₄) ppm. ¹³C{¹H} NMR (100 MHz, DMSO-*d*₆): 48.0 (s, CH₂NCH₂), 115.5–144.1 (m, C₆H₅, C₆H₄), 150.2–150.6 (m, CH=CH), 212.2 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 80.0 (s, FeP) ppm.

Preparation of [4-MeC₆H₄N(CH₂S)₂]Fe(CO)₂(dppv) (3). In order to prepare **3**, we should first prepare its precursor, the new bis(thioester) 4-MeC₆H₄N[CH₂S(O)CMe]₂. (i) Preparation of 4-MeC₆H₄N[CH₂S(O)CMe]₂: a 250 mL three-necked flask was charged with 4-MeC₆H₄NH₂ (10.7 g, 100 mmol), 37% aqueous formaldehyde (24.2 mL, 320 mmol), thioacetic acid (14.6 mL, 200 mmol), and ethanol (100 mL). While stirring, the mixture was heated to about 60 °C and stirred at this temperature for 3 h. To the resulting mixture was added 100 mL of ice-water to give some precipitates. The precipitates were collected by filtration, washed with some cold ethanol, and finally dried under vacuum to give 4-MeC₆H₄N[CH₂S(O)CMe]₂ (25.6 g, 90%) as a white solid, mp 66–67 °C. Anal. calcd for C₁₃H₁₇NO₂S₂: C, 55.10; H, 6.05; N, 4.95. Found: C, 55.35; H, 6.30; N, 4.66. IR (KBr disk): ν_{C=O} 1688 (vs.) cm⁻¹. ¹H NMR (400 MHz, CDCl₃): 2.27 (s, 3H, CH₃C₆H₄), 2.36 (s, 6H, 2CH₃C=O), 5.10 (s, 4H, CH₂NCH₂), 6.68–7.10 (m, 4H, C₆H₄) ppm. ¹³C{¹H} NMR (100 MHz, CDCl₃): 20.5 (s, CH₃C₆H₄), 31.4 (s, CH₃C=O), 53.1 (s, CH₂NCH₂), 114.7–141.6 (m, C₆H₄), 196.1 (s, C=O) ppm. (ii) Preparation of **3**: the same procedure was followed as for the preparation of **1**, except that 4-MeC₆H₄N[CH₂S(O)CMe]₂ (2.13 g, 7.5 mmol) was employed in place of PhN[CH₂S(O)CMe]₂. **3** (0.741 g, 21%) was obtained as a brown-red solid, mp 113–114 °C. Anal. calcd for C₃₇H₃₃FeNO₂P₂S₂: C, 62.98; H, 4.71; N, 1.99. Found: C, 62.72; H, 4.95; N, 2.05. IR (KBr disk): ν_{C=O} 2035 (w), 1973 (vs.) cm⁻¹. ¹H NMR (400 MHz, acetone-*d*₆): 2.18 (s, 3H, CH₃), 4.56 (s, 4H, CH₂NCH₂), 6.92–7.80 (m, 24H, 4C₆H₅, C₆H₄), 8.17–8.31 (m, 2H, CH=CH) ppm. ¹³C{¹H} NMR (100 MHz, DMSO-*d*₆): 17.7 (s, CH₃), 45.9 (s, CH₂NCH₂), 113.1–131.6 (m, C₆H₅, C₆H₄), 144.7–145.8 (m, CH=CH), 200.1 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 77.9 (s, FeP) ppm.

Preparation of [PhCH₂O₂CN(CH₂S)₂]Fe(CO)₂(dppe) (4). The same procedure was followed as for the preparation of **1**, except that PhCH₂O₂CN[CH₂S(O)CMe]₂ (2.46 g, 7.5 mmol) was used in place of PhN[CH₂S(O)CMe]₂. **4** (1.32 g, 35%) was obtained as a brown-red solid, mp 91–92 °C. Anal. calcd for C₃₈H₃₅FeNO₄P₂S₂: C, 60.73; H, 4.69; N, 1.86. Found: C, 60.42; H, 4.94; N, 1.66. IR (KBr disk): ν_{C=O} 2013 (vs.), 1968 (vs.); ν_{C=O} 1700 (vs.) cm⁻¹. ¹H NMR (400 MHz, CDCl₃): 3.02–4.56 (m, 6H, PCH₂CH₂P, OCH₂); 4.89–5.30 (m, 4H, CH₂NCH₂); 7.30–7.76 (m, 25H,

5C₆H₅) ppm. ¹³C{¹H} NMR (100 MHz, CDCl₃): 24.6–57.0 (m, PCH₂CH₂P, OCH₂), 66.7–68.0 (m, CH₂NCH₂), 127.2–154.7 (m, C₆H₅), 210.8 (s, C=O), 213.0, 213.3 (2s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 44.3 (s, FeP) ppm.

Preparation of [CpNi{(μ-SCH₂)₂NPh}Fe(CO)(dppv)]BF₄ (5). While stirring, nickelocene (0.038 g, 0.2 mmol) was dissolved in diethyl ether (5 mL) and then HBF₄·Et₂O (27.2 μL, 0.2 mmol) was added. After the mixture was stirred at room temperature for 15 min, a brown-red suspension was formed. To this suspension was added a solution of **1** (0.138 g, 0.2 mmol) in CH₂Cl₂ (5 mL). The new mixture was stirred at room temperature for 2 h, solvent was removed at reduced pressure to give a residue, which was subjected to column chromatography (silica gel), elution with CH₂Cl₂/acetone (v/v = 10 : 1) developed a major red band, from which **5** (0.052 g, 30%) was obtained as brown-red solid. Mp 220–221 °C. Anal. calcd for C₄₀H₃₆BF₄FeNNiOP₂S₂: C, 54.96; H, 4.15; N, 1.60. Found: C, 54.86; H, 3.98; N, 1.62. IR (KBr disk): ν_{C=O} 1956 (m) cm⁻¹. ¹H NMR (400 MHz, acetone-*d*₆): 4.43 (s, 5H, C₅H₅), 4.68, 5.47 (dd, *J* = 14.6 Hz, 4H, CH₂NCH₂), 6.77–8.13 (m, 25H, 5C₆H₅), 8.71, 8.86 (2s, 2H, CH=CH) ppm. ¹³C{¹H} NMR (100 MHz, DMSO-*d*₆): 54.4 (s, CH₂NCH₂), 94.5, 95.9 (2s, C₅H₅), 115.4–142.2 (m, C₆H₅), 149.3–149.9 (m, CH=CH), 205.9 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 82.7 (s, FeP) ppm.

Preparation of [CpNi{(μ-SCH₂)₂NC₆H₄Cl-4}Fe(CO)(dppv)]BF₄ (6). The same procedure was followed as for the preparation of **5**, except that **2** (0.145 g, 0.2 mmol) was utilized instead of **1**. Complex **6** (0.113 g 62%) was obtained as a brown-red solid, mp 197–198 °C. Anal. calcd for C₄₀H₃₅BClF₄FeNNiOP₂S₂: C, 52.88; H, 3.88; N, 1.54. Found: C, 52.84; H, 4.01; N, 1.62. IR (KBr disk): ν_{C=O} 1952 (m) cm⁻¹. ¹H NMR (400 MHz, acetone-*d*₆): 4.47 (s, 5H, C₅H₅), 4.64, 5.49 (dd, 4H, *J* = 15.4 Hz, CH₂NCH₂), 7.32–8.13 (m, 24H, 4C₆H₅, C₆H₄), 8.73, 8.88 (2s, 2H, CH=CH) ppm. ¹³C{¹H} NMR (100 MHz, CD₂Cl₂): 52.6, 54.1 (2s, CH₂NCH₂), 94.8, 96.5 (2s, C₅H₅), 117.0–142.4 (m, C₆H₅, C₆H₄), 149.4–151.3 (m, CH=CH), 210.9 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 83.9 (s, FeP) ppm.

Preparation of [CpNi{(μ-SCH₂)₂NC₆H₄Me-4}Fe(CO)(dppv)]BF₄ (7). The same procedure was followed as for the preparation of **5**, except that **3** (0.141 g, 0.2 mmol) was employed in place of **1**. Complex **7** (0.038 g, 21%) was obtained as a brown-red solid, mp 210–211 °C. Anal. calcd for C₄₁H₃₈BF₄FeNNiOP₂S₂: C, 55.45; H, 4.31; N, 1.58. Found: C, 55.23; H, 4.28; N, 1.34. IR (KBr disk): ν_{C=O} 1955 (s) cm⁻¹. ¹H NMR (400 MHz, acetone-*d*₆): 2.12 (s, 3H, CH₃), 4.42 (s, 5H, C₅H₅), 4.67, 5.41 (dd, 4H, *J* = 13.0 Hz, CH₂NCH₂), 7.14–8.13 (m, 24H, 4C₆H₅, C₆H₄), 8.69–8.84 (m, 2H, CH=CH) ppm. ¹³C{¹H} NMR (100 MHz, CD₃CN): 20.1 (s, CH₃), 54.8 (s, CH₂NCH₂), 96.1 (s, C₅H₅), 117.0–141.5 (m, C₆H₅, C₆H₄), 150.5–151.2 (m, CH=CH), 219.5 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 83.7 (s, FeP) ppm.

Preparation of [CpNi{(μ-SCH₂)₂NCO₂CH₂Ph}Fe(CO)(dppe)]BF₄ (8). The same procedure was followed as for the preparation of **5**, except that **4** (0.150 g, 0.2 mmol) was used instead of **1**. Complex **8** (0.052 g, 28%) was obtained as a brown-red solid, mp 121–122 °C. Anal. calcd for C₄₂H₄₀BF₄FeNNiO₃P₂S₂: C, 54.00; H, 4.32; N, 1.50. Found: C, 54.27; H, 4.43; N, 1.58. IR (KBr disk): ν_{C=O} 1940 (vs.); ν_{C=O} 1704 (vs.) cm⁻¹. ¹H NMR (400 MHz,

CDCl₃): 3.03–4.05 (m, 6H, PCH₂CH₂P, OCH₂), 4.54 (s, 5H, C₅H₅), 4.98–5.31 (m, 4H, CH₂NCH₂), 7.12–7.86 (m, 25H, 5C₆H₅) ppm. ¹³C{¹H} NMR (100 MHz, CD₃CN): 29.5, 29.8 (2s, PCH₂CH₂P), 50.7 (s, CH₂NCH₂), 69.5 (s, OCH₂), 96.5 (s, C₅H₅), 128.9–154.1 (m, C₆H₅), 211.0 (s, C=O), 216.4 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, CDCl₃): 78.3, 79.1 (2s, FeP) ppm.

Preparation of [CpNi{(μ-SCH₂)₂NH}Fe(CO)(dppe)]BF₄ (9). To a solution of **8** (0.093 g, 0.1 mmol) in CH₂Cl₂ (5 mL) were added PhOMe (0.5 mL) and HBF₄·Et₂O (408 μL, 3.0 mmol). The mixture was stirred at room temperature for 4 h to give a brown-red solution. Solvents were removed to give a residue, which was subjected to column chromatography (silica gel). Elution with CH₂Cl₂/acetone developed a major red band, from which **9** (0.034 g, 42%) was obtained as a brown-red solid, mp 170 °C (dec). Anal. calcd for C₃₄H₃₄BF₄FeNNiOP₂S₂: C, 51.04; H, 4.28; N, 1.75. Found: C, 50.81; H, 4.56; N, 1.70. IR (KBr disk): ν_{C≡O} 1939 (vs.) cm⁻¹. ¹H NMR (400 MHz, CD₃CN): 2.26 (br.s, 1H, NH), 3.04–4.42 (m, 8H, PCH₂CH₂P, CH₂NCH₂), 4.61 (s, 5H, C₅H₅), 7.43–7.95 (m, 20H, 4C₆H₅) ppm. ¹³C{¹H} NMR (100 MHz, CD₃CN): 29.2, 29.6 (2s, PCH₂CH₂P), 52.4 (s, CH₂NCH₂), 95.7 (s, C₅H₅), 129.2–133.8 (m, C₆H₅), 218.3 (s, C≡O) ppm. ³¹P{¹H} NMR (162 MHz, acetone-d₆): 78.1 (s, FeP) ppm.

Electrochemical and electrocatalytic experiments

Acetonitrile (Amethyst Chemicals, HPLC grade) for electrochemical and electrocatalytic experiments was used directly without further purification. A solution of 0.1 M *n*-Bu₄NPF₆ in MeCN was used as the electrolyte in all CV experiments. The *n*-Bu₄NPF₆ electrolyte was dried in an oven at 110 °C for at least 24 h. Dry N₂ was purged through the solutions for at least 10 min before the measurements were taken. The measurements were made by using a BAS Epsilon potentiostat. All voltammograms were obtained in a three-electrode cell with a 3 mm diameter glass carbon working electrode, a platinum counter electrode, and a Ag/Ag⁺ (0.01 M AgNO₃/0.1 M *n*-Bu₄NPF₆ in MeCN) reference electrode under an atmosphere of nitrogen. The working electrode was polished with 0.05 μm alumina paste and sonicated in water for about 10 min prior to use. All potentials are quoted against Fc/Fc⁺ potential.

X-ray crystal structure determinations of 6–8

While single crystals of **6–8** for X-ray diffraction analysis were grown by slow diffusion of hexane into their CH₂Cl₂ solutions at –5 °C. A single crystal of **6–8** was mounted on a Rigaku MM-007 (rotating anode) diffractometer equipped with a Bruker P4 accessory. While the data of **6** and **8** were collected using a confocal monochromator with Mo Kα radiation (λ = 0.71073 Å) in the ω scanning mode at 113 K, data of **7** were collected using a confocal monochromator with Mo Kα radiation (λ = 0.71073 Å) in the ω scanning mode at 293 K, respectively. Data collection, reduction, and absorption correction were performed by the CRYSTALCLEAR program.⁵⁹ The structures were solved by direct methods using the SHELXS program⁶⁰ and refined by full-matrix least-squares techniques (SHELXL)⁶¹ on F². Hydrogen atoms were located by using the geometric

method. Details of crystal data, data collections, and structure refinements are summarized in Table 3.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

We are grateful to the National Natural Science Foundation of China (21772106) and the Ministry of Science and Technology of China (973 program 2014CB845604) for financial support.

References

- 1 R. Cammack, *Nature*, 1999, **397**, 214–215.
- 2 J. C. Fontecilla-Camps, A. Volbeda, C. Cavazza and Y. Nicolet, *Chem. Rev.*, 2007, **107**, 4273–4303.
- 3 W. Lubitz, H. Ogata, O. Rüdiger and E. Reijerse, *Chem. Rev.*, 2014, **114**, 4081–4148.
- 4 A. Volbeda, M.-H. Charon, C. Piras, E. C. Hatchikian, M. Frey and J. C. Fontecilla-Camps, *Nature*, 1995, **373**, 580–587.
- 5 A. Volbeda, E. Garcin, C. Piras, A. L. de Lacey, V. M. Fernandez, E. C. Hatchikian, M. Frey and J. C. Fontecilla-Camps, *J. Am. Chem. Soc.*, 1996, **118**, 12989–12996.
- 6 A. Volbeda, L. Martin, C. Cavazza, M. Matho, B. W. Faber, W. Roseboom, S. P. J. Albracht, E. Garcin, M. Rousset and J. C. Fontecilla-Camps, *J. Biol. Inorg. Chem.*, 2005, **10**, 239–249.
- 7 R. Cammack, M. Frey and R. Robson, *Hydrogen as a fuel: Learning from nature*, Taylor & Francis, London, 2001.
- 8 J. A. Turner, *Science*, 2004, **305**, 972–974.
- 9 W. Lubitz and W. Tumas, *Chem. Rev.*, 2007, **107**, 3900–3903.
- 10 T. R. Simmons, G. Berggren, M. Bacchi, M. Fontecave and V. Artero, *Coord. Chem. Rev.*, 2014, **270–271**, 127–150.
- 11 S. Ogo, *Coord. Chem. Rev.*, 2017, **334**, 43–53.
- 12 D. Schilter, J. M. Camara, M. T. Huynh, S. Hammes-Schiffer and T. B. Rauchfuss, *Chem. Rev.*, 2016, **116**, 8693–8749.
- 13 N. J. Lindenmaier, S. Wahlefeld, E. Bill, T. Szilvási, C. Eberle, S. Yao, P. Hildebrandt, M. Horch, I. Zebger and M. Driess, *Angew. Chem., Int. Ed.*, 2017, **56**, 2208–2211.
- 14 C. U. Perotto, G. Marshall, G. J. Jones, E. S. Davies, W. Lewis, J. McMaster and M. Schröder, *Chem. Commun.*, 2015, **51**, 16988–16991.
- 15 C. U. Perotto, C. L. Sodipo, G. J. Jones, J. P. Tidey, A. J. Blake, W. Lewis, E. S. Davies, J. McMaster and M. Schröder, *Inorg. Chem.*, 2018, **57**, 2558–2569.
- 16 F. Osterloh, W. Saak, D. Haase and S. Pohl, *Chem. Commun.*, 1997, 979–980.
- 17 L.-C. Song, Y. Lu, L. Zhu and Q.-L. Li, *Organometallics*, 2017, **36**, 750–760.
- 18 J. Jiang, M. Maruani, J. Solaimanzadeh, W. Lo, S. A. Koch and M. Millar, *Inorg. Chem.*, 2009, **48**, 6359–6361.
- 19 D. Sellmann, F. Geipel, F. Lauderbach and F. W. Heinemann, *Angew. Chem., Int. Ed.*, 2002, **41**, 632–634.



- 20 L.-C. Song, X.-Y. Gao, W.-B. Liu, H.-T. Zhang and M. Cao, *Organometallics*, 2018, **37**, 1050–1061.
- 21 M. C. Smith, J. E. Barclay, S. P. Cramer, S. C. Davies, W.-W. Gu, D. L. Hughes, S. Longhurst and D. J. Evans, *J. Chem. Soc., Dalton Trans.*, 2002, 2641–2647.
- 22 J. A. W. Verhagen, M. Lutz, A. L. Spek and E. Bouwman, *Eur. J. Inorg. Chem.*, 2003, 3968–3974.
- 23 Z. Li, Y. Ohki and K. Tatsumi, *J. Am. Chem. Soc.*, 2005, **127**, 8950–8951.
- 24 D. Sellmann, F. Lauderbach and F. W. Heinemann, *Eur. J. Inorg. Chem.*, 2005, 371–377.
- 25 S. Ogo, K. Ichikawa, T. Kishima, T. Matsumoto, H. Nakai, K. Kusaka and T. Ohhara, *Science*, 2013, **339**, 682–684.
- 26 Y. Ohki, K. Yasumura, M. Ando, S. Shimokata and K. Tatsumi, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 3994–3997.
- 27 D. Brazzolotto, L. Wang, H. Tang, M. Gennari, N. Queyriaux, C. Philouze, S. Demeshko, F. Meyer, M. Orio, V. Artero, M. B. Hall and C. Duboc, *ACS Catal.*, 2018, **8**, 10658–10667.
- 28 D. Brazzolotto, M. Gennari, N. Queyriaux, T. R. Simmons, J. Pécaut, S. Demeshko, F. Meyer, M. Orio, V. Artero and C. Duboc, *Nat. Chem.*, 2016, **8**, 1054–1060.
- 29 (a) L.-C. Song, X.-Y. Yang, M. Cao, X.-Y. Gao, B.-B. Liu, L. Zhu and F. Jiang, *Chem. Commun.*, 2017, **53**, 3818–3821; (b) L.-C. Song, X.-Y. Yang, X.-Y. Gao and M. Cao, *Inorg. Chem.*, 2019, **58**, 39–42; (c) L.-C. Song, X.-J. Sun, P.-H. Zhao, J.-P. Li and H.-B. Song, *Dalton Trans.*, 2012, **41**, 8941–8950.
- 30 H.-J. Fan and M. B. Hall, *J. Am. Chem. Soc.*, 2001, **123**, 3828–3829.
- 31 A. Silakov, B. Wenk, E. Reijerse and W. Lubitz, *Phys. Chem. Chem. Phys.*, 2009, **11**, 6592–6599.
- 32 G. Berggren, A. Adamska, C. Lambert, T. R. Simmons, J. Esselborn, M. Atta, S. Gambarelli, J.-M. Mouesca, E. Reijerse, W. Lubitz, T. Happe, V. Artero and M. Fontecave, *Nature*, 2013, **499**, 66–69.
- 33 L. P. Battaglia, G. P. Chiusoli, D. Delledonne, M. Nardelli, C. Pelizzi and G. Predieri, *Gazz. Chim. Ital.*, 1989, **119**, 345–348.
- 34 T. A. Manuel, *Inorg. Chem.*, 1963, **2**, 854–858.
- 35 A. Salzer and H. Werner, *Angew. Chem., Int. Ed.*, 1972, **11**, 930–932.
- 36 C. Elschenbroich, *Organometallics. Third, Completely Revised and Extended Edition*, WILEY-VCH Verlag GMBH & Co. KGdt, Weinheim Germany, 2006, p. 506.
- 37 C. Fichtner, C. Laurich, E. Bothe and W. Lubitz, *Biochemistry*, 2006, **45**, 9706–9716.
- 38 H. Ogata, K. Nishikawa and W. Lubitz, *Nature*, 2015, **520**, 571–574.
- 39 B. Cordero, V. Gómez, A. E. Platero-Prats, M. Revés, J. Echeverría, E. Cremades, F. Barragán and S. Alvarez, *Dalton Trans.*, 2008, 2832–2838.
- 40 G. M. Chambers, M. T. Huynh, Y. Li, S. Hammes-Schiffer, T. B. Rauchfuss, E. Reijerse and W. Lubitz, *Inorg. Chem.*, 2016, **55**, 419–431.
- 41 L.-C. Song, X.-F. Han, W. Chen, J.-P. Li and X.-Y. Wang, *Dalton Trans.*, 2017, **46**, 10003–10013.
- 42 M. E. Carroll, B. E. Barton, D. L. Gray, A. E. Mack and T. B. Rauchfuss, *Inorg. Chem.*, 2011, **50**, 9554–9563.
- 43 L.-C. Song, M. Cao and Y.-X. Wang, *Dalton Trans.*, 2015, **44**, 6797–6808.
- 44 S. Canaguier, L. Vaccaro, V. Artero, R. Ostermann, J. Pécaut, M. J. Field and M. Fontecave, *Chem.-Eur. J.*, 2009, **15**, 9350–9364.
- 45 M. Y. Darensbourg, I. Font, D. K. Mills, M. Pala and J. H. Reibenspies, *Inorg. Chem.*, 1992, **31**, 4965–4971.
- 46 B. Adhikary, S. Liu and C. R. Lucas, *Inorg. Chem.*, 1993, **32**, 5957–5962.
- 47 P. Zanello, *Inorganic Electrochemistry. Theory, Practice and Application*, Thomas Graham House, Cambridge, UK, 2003.
- 48 K. Izutsu, *Acid-Base Dissociation Constants in Dipolar Aprotic Solvents*, Blackwell Scientific Publications, Oxford, UK, 1990.
- 49 G. A. N. Felton, R. S. Glass, D. L. Lichtenberger and D. H. Evans, *Inorg. Chem.*, 2006, **45**, 9181–9184.
- 50 W. Dong, M. Wang, T. Liu, X. Liu, K. Jin and L. Sun, *J. Inorg. Biochem.*, 2007, **101**, 506–513.
- 51 (a) L.-C. Song, L. Zhu, F.-Q. Hu and Y.-X. Wang, *Inorg. Chem.*, 2017, **56**, 15216–15230; (b) L.-C. Song, F.-Q. Hu, M.-M. Wang, Z.-J. Xie, K.-K. Xu and H.-B. Song, *Dalton Trans.*, 2014, **43**, 8062–8071.
- 52 (a) S. C. Eady, T. Breault, L. Thompson and N. Lehnert, *Dalton Trans.*, 2016, **45**, 1138–1151; (b) S. Kaur-Ghumaan, L. Schwartz, R. Lomoth, M. Stein and S. Ott, *Angew. Chem., Int. Ed.*, 2010, **49**, 8033–8036.
- 53 G. A. N. Felton, C. A. Mebi, B. J. Petro, A. K. Vannucci, D. H. Evans, R. S. Glass and D. L. Lichtenberger, *J. Organomet. Chem.*, 2009, **694**, 2681–2699.
- 54 B. E. Barton and T. B. Rauchfuss, *J. Am. Chem. Soc.*, 2010, **132**, 14877–14885.
- 55 E. S. Rountree, B. D. McCarthy, T. T. Eisenhart and J. L. Dempsey, *Inorg. Chem.*, 2014, **53**, 9983–10002.
- 56 L.-C. Song, W.-B. Liu and B.-B. Liu, *Organometallics*, 2020, **39**, 1431–1439.
- 57 T. Izawa, Y. Terao and K. Suzuki, *Tetrahedron: Asymmetry*, 1997, **8**, 2645–2648.
- 58 C. Sommer, C. P. Richers, W. Lubitz, T. B. Rauchfuss and E. J. Reijerse, *Angew. Chem., Int. Ed.*, 2018, **57**, 5429–5432.
- 59 Rigaku Americas and Rigaku, *CrystalClear and CrystalStructure*, Rigaku and Rigaku Americas, The Woodlands, TX, 2007.
- 60 (a) G. M. Sheldrick, *SHELXS97, Program for Crystal Structure Solution*, University of Göttingen, Germany, 1997; (b) G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 61 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 3–8.

