

Cite this: *Chem. Sci.*, 2019, 10, 761

All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 30th September 2018  
Accepted 26th October 2018

DOI: 10.1039/c8sc04346a

rsc.li/chemical-science

# Bimetallic nickel–cobalt hydrides in H<sub>2</sub> activation and catalytic proton reduction†

Xiaoxiao Chu,<sup>ab</sup> Jihao Jin,<sup>a</sup> Bangrong Ming,<sup>a</sup> Maofu Pang,<sup>a</sup> Xin Yu,<sup>a</sup> Chen-Ho Tung<sup>a</sup> and Wenguang Wang<sup>id</sup>\*<sup>a</sup>

The synergism of the electronic properties of nickel and cobalt enables bimetallic NiCo complexes to process H<sub>2</sub>. The nickel–cobalt hydride [(dppe)Ni(pdt)(H)CoCp\*]<sup>+</sup> ([1H]<sup>+</sup>) arising from protonation of the reduced state **1** was found to be an efficient electrocatalyst for H<sub>2</sub> evolution with Cl<sub>2</sub>CHCOOH, and the oxidized [Ni(II)Co(III)]<sup>2+</sup> form is capable of activating H<sub>2</sub> to produce [1H]<sup>+</sup>. The features of stereodynamics, acid–base properties, redox chemistry and reactivity of these bimetallic NiCo complexes in processing H<sub>2</sub> are potentially related to the active site of [NiFe]-H<sub>2</sub>ases.

## Introduction

Bimetallic complexes have emerged as important components in inorganic chemistry, homogeneous catalysis and biocatalysis.<sup>1–5</sup> Compared to monometallic catalytic processes, bimetallic catalysis can promote the rate and selectivity of a reaction catalyzed by the first metal in the bimetallic catalyst by synergism with the electronic and steric properties of the second metal, or through a concerted activation process in which both metals participate in activation of the substrate, lowering the activation barrier.<sup>6,7</sup> Bimetallic catalysis is common in metalloenzymes that perform multi-electron redox reactions.<sup>8–10</sup> Examples are the active sites of [FeFe]-H<sub>2</sub>ases and [NiFe]-H<sub>2</sub>ases, which catalyze the production and uptake of dihydrogen.<sup>11,12</sup> Both of the catalysis invoke bimetallic hydride intermediates, but the [FeFe]-H<sub>2</sub>ases are thought to feature terminal-hydride intermediates while [NiFe]-H<sub>2</sub>ases are proposed to operate through bridging hydrides in catalytically significant states (Fig. 1).<sup>13–16</sup>

Reflecting interest in utilization of earth abundant metals for controlling energy storage and release, functional modeling of hydrogenases continues to attract considerable attention. Compared to [FeFe]-H<sub>2</sub>ase mimics, modeling the active site of [NiFe]-hydrogenases has proven more challenging<sup>17–23</sup> and very few of the models clarify the processing of H<sub>2</sub>.<sup>24–28</sup> Especially, the nickel–iron hydride models [Ni(II)HFe(II)]<sup>+</sup> remain rare although they are important in modeling research.<sup>29</sup> The

established [Ni(II)HFe(II)]<sup>+</sup> models are [(dppe)Ni(pdt)(H)Fe(PR<sub>3</sub>)<sub>x</sub>(CO)<sub>3–x</sub>]<sup>+</sup>,<sup>30</sup> [Ni(N<sub>2</sub>S<sub>2</sub>)Fe(H)(P(OEt)<sub>3</sub>)<sub>3</sub>]<sup>+</sup>,<sup>26</sup> [(dppe)Ni(pdt)(H)Fe(CNBAr<sup>F</sup><sub>3</sub>)<sub>2</sub>(CO)]<sup>–</sup> (ref. 27) and [(pnp)Ni(pdt)(H)Fe(CO)(dppv)]<sup>+</sup>.<sup>28</sup> They were derived from protonation of the corresponding reduced Ni(II)Fe(I) state or were generated from the activation of H<sub>2</sub> with an oxidized [Ni(II)Fe(II)]<sup>2+</sup> precursor (Fig. 2a).

Biological catalysis in a bimetallic manner offers guidance for the exploration of heteronuclear redox chemistry to precede small molecule activations under mild conditions. Four-coordinate nickel(II) complexes such as Ni(N<sub>2</sub>S<sub>2</sub>)<sup>24,31</sup> and (dppe)Ni(pdt)<sup>32,33</sup> are optimal modules to assemble diverse bimetallic complexes, which expand the synthetic NiFe models to the nickel-based heterobimetallic NiM platform and encourage the exploration of fast bioinspired catalysts for the production or activation of H<sub>2</sub>.<sup>31,34–37</sup> For example, [NiRu] complexes bearing the Ni(N<sub>2</sub>S<sub>2</sub>) metalloligand are active for H<sub>2</sub> heterolysis providing [Ni(II)HRu(II)]<sup>+</sup> hydrides with the assistance of an external base (Fig. 2b).<sup>33,38</sup>

Cobalt is earth-abundant and many cobalt complexes have been shown to be efficient catalysts for H<sub>2</sub> production<sup>11,39</sup> and hydrogenation reactions.<sup>40,41</sup> However, bimetallic NiCo complexes have been reported only infrequently<sup>42</sup> and the related heteronuclear hydride complexes have never been documented. In this work, we produced a heterobimetallic

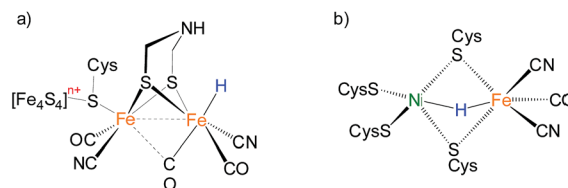


Fig. 1 Hypothesized "hydride structure" for the active site of (a) [FeFe]-H<sub>2</sub>ases and (b) [NiFe]-H<sub>2</sub>ases.

<sup>a</sup>Key Lab for Colloid and Interface Chemistry of Education Ministry, School of Chemistry and Chemical Engineering, Shandong University, 250100, China. E-mail: ww@sdzu.edu.cn

<sup>b</sup>School of Chemistry and Materials Science, Ludong University, Yantai, 264025, China

† Electronic supplementary information (ESI) available: Experimental results, procedures and characterization including NMR spectra and crystallographic data. CCDC 1841904–1841908. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8sc04346a



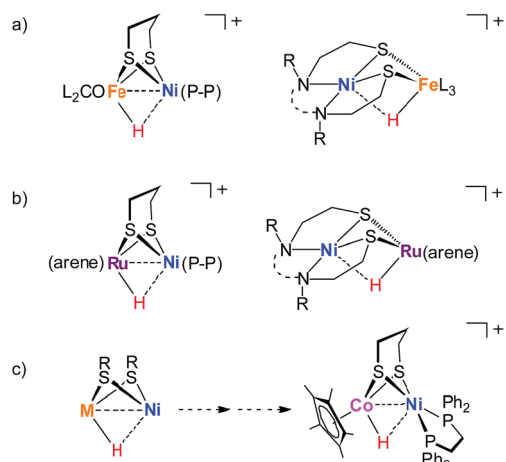


Fig. 2 Examples of nickel-based heterobimetallic hydride complexes: (a) reported Ni(II)–H–Fe(III), (b) reported Ni(II)–H–Ru(III), and (c) Ni(II)–H–Co(III) described in this work.

[Ni(II)Co(III)H]<sup>+</sup> hydride [(dppe)Ni(pdt)(H)CoCp\*]<sup>+</sup> (**[1H]**<sup>+</sup>, pdt<sup>2−</sup> = 1,3-(CH<sub>2</sub>)<sub>3</sub>S<sub>2</sub><sup>2−</sup>, Cp\*<sup>−</sup> = Me<sub>5</sub>C<sub>5</sub><sup>−</sup>, and dppe = Ph<sub>2</sub>PC<sub>2</sub>H<sub>4</sub>PPh<sub>2</sub>) by protonating the reduced state [(dppe)Ni(pdt)CoCp\*] (**1**) for a catalytic proton reduction (Fig. 2c). Processing of H<sub>2</sub> with a class of Ni–Co complexes, which are potentially related to the structure and properties of the [NiFe]–H<sub>2</sub>ase, has been studied.

## Results and discussion

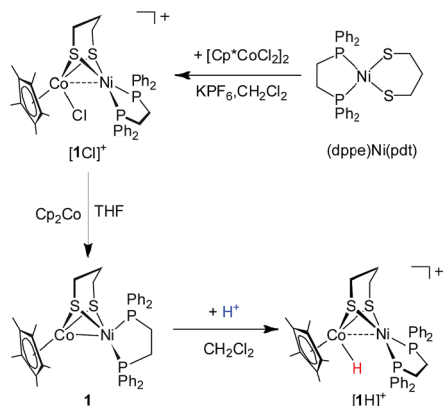
### [(dppe)Ni(pdt)(Cl)CoCp\*]<sup>+</sup> and (dppe)Ni(pdt)CoCp\*

The synthesis of a heteronuclear Ni(II)Co(III) dithiolate [**1Cl**]<sup>+</sup> entailed the assembly of (dppe)Ni(pdt) to [Cp\*CoCl<sub>2</sub>]<sub>2</sub>. Treatment of [Cp\*CoCl<sub>2</sub>]<sub>2</sub> with two equiv. of (dppe)Ni(pdt) and KPF<sub>6</sub> solids in CH<sub>2</sub>Cl<sub>2</sub> results in the formation of the product [**1Cl**]<sup>+</sup> in a high yield (Scheme 1). The <sup>31</sup>P NMR spectrum of [**1Cl**]<sup>+</sup> has a sharp singlet at δ 54.4 corresponding to dppe, indicating that the two phosphine groups are chemically equivalent. Crystallographic analysis of [**1Cl**]<sup>+</sup> reveals that the Ni center adopts a square-planar geometry (Fig. 3) and is linked to the Cp\*Co fragment through the pdt<sup>2−</sup> ligand. The Ni⋯Co distance of

2.926(2) Å far exceeds the sum of the covalent atomic radii of Ni (1.24 Å) and Co (1.26 Å, low-spin).<sup>43</sup>

Reduction of the cationic Ni(II)Co(III) complex [**1Cl**]<sup>+</sup> with two equivalents of Cp<sub>2</sub>Co affords the reduced compound **1**. It dissolves well in nonpolar solvents such as toluene or benzene to form deep red solutions, which is very air sensitive and decomposes into insoluble species. Crystallographic analysis of **1** confirms a neutral complex with the formula (dppe)Ni(pdt)CoCp\* (Fig. 3). The structure of **1** is more compact than that of [**1Cl**]<sup>+</sup> and this is reflected by the Ni–Co distance, which is 0.454 Å shorter after the 2e<sup>−</sup> reduction. The Ni–Co distance of 2.4722(5) Å in **1** is comparable to the Ni–Fe distance of 2.4666(6) Å in (dppe)Ni(pdt)Fe(CO)<sub>3</sub>.<sup>17</sup> Given the sum of the covalent atomic radii of Ni and Co, the Ni–Co separation suggests a metal–metal bond. The 2e<sup>−</sup> reduction causes the Cp\*–Co distance to decrease from 1.699 Å to 1.686 Å. Since the Ni center adopts an approximately tetrahedral geometry, compound **1** can be appropriately considered as a Ni(I)Co(II) species in the solid state.

Interestingly, the <sup>31</sup>P NMR spectroscopic analysis shows that compound **1** exhibits two isomers in solution at room temperature (eqn (1)). The phosphorus resonance signals appeared as two broad signals at δ 42.9 and 41.8 in a ratio of approximately 1 : 1 (Fig. 4). When the temperature was increased to 318 K, the broad resonance at δ 41.8 became resolved but the ratio of the two signals was maintained. Decreasing the temperature to 253



Scheme 1 Synthesis of Ni–Co bimetallic complexes.

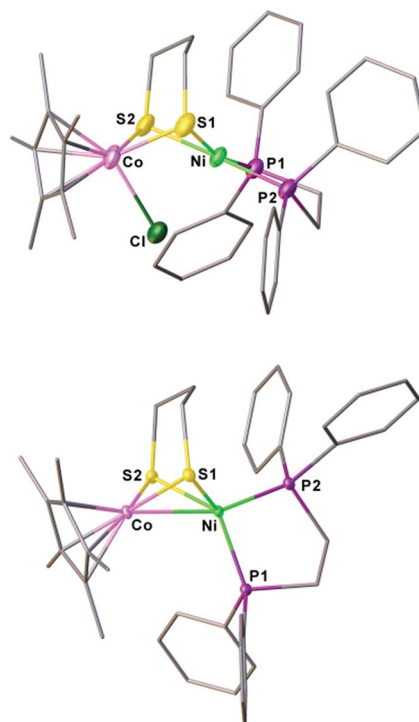


Fig. 3 Structures of [**1Cl**]<sup>+</sup> and **1** with thermal ellipsoids drawn at the 50% probability level. The counter anion PF<sub>6</sub><sup>−</sup> in [**1Cl**]<sup>+</sup> and all hydrogen atoms are omitted for clarity. Selected distances (Å): for [**1Cl**]<sup>+</sup>, Co–Ni, 2.926(2); Co–Cl, 2.316(2); Co–S1, 2.252(2); Co–S2, 2.259(2); Ni–S1, 2.258(2); Ni–S2, 2.247(2). For **1**, Co–Ni, 2.4722(5); Co–S1, 2.1947(7); Co–S2, 2.1820(7); Ni–S1, 2.2645(7); Ni–S2, 2.2268(7).

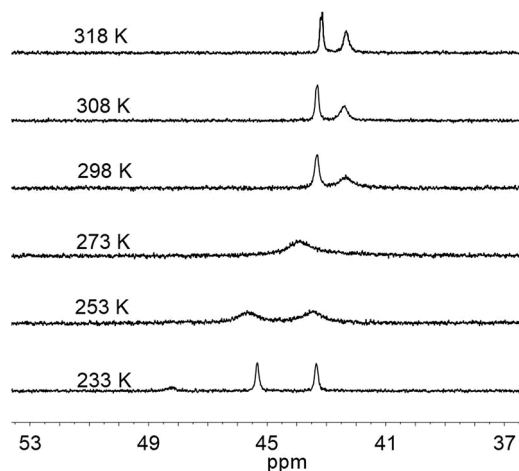
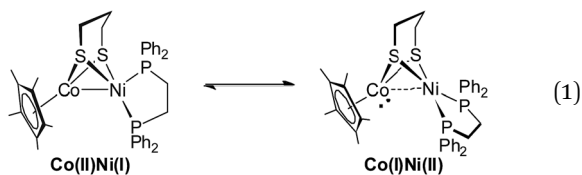


Fig. 4  $^{31}\text{P}$  NMR spectra of **1** recorded at various temperatures in toluene.

K, the two resonances coalesced into a single broad peak at  $\delta$  44. At this lower temperature, however, the broad signal decoalesced into two peaks at 45.7 and 43.6 ppm, which is thought to be consistent with the tetrahedral geometry of Ni that was observed in the solid state structure. Overall, the results of  $^{31}\text{P}$  NMR observations at temperatures ranging from 233 K to 298 K reveal a dynamic process in which the  $^{31}\text{P}$  sites in dppe are interchanged. The rotation of dppe at the Ni site could proceed through an intermediate or transition state with square-planar Ni.<sup>30b</sup> It is more likely that in solution at room temperature, **1** consists of two isomers with resonance states of Ni(I)Co(II) and Ni(II)Co(I). Such rotation-induced redox behavior has also been described by Rauchfuss *et al.*<sup>30a</sup> for  $(\text{CO})_3\text{Fe}(\text{pdt})\text{Ni}(\text{dppe})$ . In addition, the reduced NiRu compound (cymene)Ru(pdt)Ni(dppe) features a rigid tetrahedral Ni(0) center.<sup>33a</sup>



### Protonation of (dppe)Ni(pdt)CoCp\*

The reduced compound **1** undergoes protonation, affording the  $[\text{Ni}(\text{II})\text{Co}(\text{III})\text{H}]^+$  hydride complex  $[\mathbf{1H}]^+$ . Addition of an equivalent amount of an acid such as  $[\text{HPPH}_3]\text{BF}_4$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 8.0$ )<sup>44</sup> or  $\text{Cl}_2\text{CHCOOH}$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 13.2$ )<sup>45</sup> to solutions of **1** in  $\text{CH}_2\text{Cl}_2$  immediately leads to the color changing from brown to dark. The resulting complex  $[\mathbf{1H}]^+$  was isolated by filtration after dilution of the reaction solution with  $\text{Et}_2\text{O}$ . The  $^1\text{H}$  NMR spectrum of  $[\mathbf{1H}]^+$  features a hydride signal at  $\delta$   $-9.77$  as a singlet (Fig. S9†). No  $^{31}\text{P}$  coupling was resolved, indicating that the hydride is located at the Co(III) center rather than being bound to Ni.<sup>46</sup> The  $^{31}\text{P}$  signal of the dppe group appears as a singlet at  $\delta$  67.5, about 13.1 ppm down-field shifted relative to that of  $[\mathbf{1Cl}]^+$  (Fig. S8†). An alternative approach to synthesize  $[\mathbf{1H}]^+$  is

by displacement of the  $\text{Cl}^-$  ligand of  $[\mathbf{1Cl}]^+$  by a hydride ligand donated from  $\text{NaBH}_4$ .

Crystallographic analysis agrees with the NMR spectroscopic assignments of the nickel-cobalt dithiolates as a hydride. The framework of  $[(\text{dppe})\text{Ni}(\text{pdt})(\text{X})\text{CoCp}^*]^+$  is very similar to that of  $[\mathbf{1Cl}]^+$  (Fig. 5). The striking difference is the Ni...Co separation (2.556(1) Å), which is 0.37 Å shorter than that in  $[\mathbf{1Cl}]^+$ . Protonation causes the Ni–Co distance to increase only by 0.1 Å, which is consistent with the metal–metal bond changes associated with the protonation of reduced NiFe models.<sup>30a</sup> The hydride ligand was located and refined and the Co–H bond length of 1.45(5) Å is much shorter than the Ni–H bond length (1.91(5) Å), suggesting that the hydride is strongly coordinated at the Co center instead of adopting a bridging position between the bimetallic centers.

Given the rotation-induced redox behavior of the reduced compound, we propose that the protonation reaction proceeds *via* the mixed-valent conformer of Ni(II)Co(I). With dppe rotation, nickel is stabilized in the oxidized Ni(II) form with concomitant cleavage of the metal–metal bond resulting in cobalt being in a strongly reduced Co(I) state.<sup>29,30b</sup> The formation of the  $[\text{Ni}(\text{II})\text{Co}(\text{III})\text{H}]^+$  hydride arises from protonation of the  $\text{Cp}^*\text{Co}$  site rather than the metal–metal bond. This “redox isomerization–protonation” mechanism is close to that proposed for  $(\text{CO})_3\text{Fe}(\text{pdt})\text{Ni}(\text{dppe})$  and reflects the redox-flexibility of the (dppe)Ni(pdt) module.<sup>30</sup>

### Acidity of $[\mathbf{1H}]^+$

The acidic character of the hydride ligand in  $[\mathbf{1H}]^+$  was manifested by the H/D exchange reaction with  $\text{D}_2\text{O}$  giving  $[\mathbf{1D}]^+$  solely and with no liberation of HD or  $\text{H}_2$ . After addition of excess  $\text{D}_2\text{O}$  (20 equiv.) to the  $\text{CD}_3\text{CN}$  solution (0.6 mL) of  $[\mathbf{1H}]^+$  in a J. Young tube, the reaction process was monitored by NMR spectroscopy.  $^1\text{H}$  NMR spectroscopic analysis showed that the hydride signal disappears slowly, while the  $^{31}\text{P}$  resonance remains as a singlet at  $\delta$  67.4. The appearance of a Co–D signal was observed at  $\delta$   $-9.66$  in the  $^2\text{H}$  NMR spectrum (Fig. S14†). In agreement with the calculated isotopic distribution of  $[\mathbf{1D}]^+$ , ESI-MS spectral analysis featured a characteristic peak at  $m/z = 758.1216$  for  $[\mathbf{1D}]^+$  vs.  $m/z = 757.1175$  for  $[\mathbf{1H}]^+$ . The exchange reaction between  $[\mathbf{1H}]^+$  and  $\text{D}_2\text{O}$  is relatively slow. A plot of the

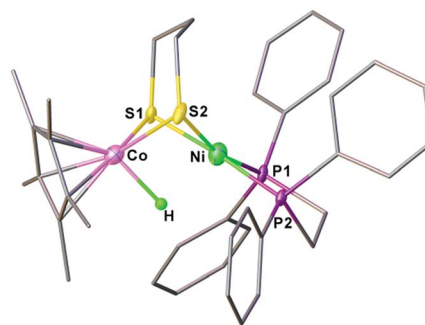
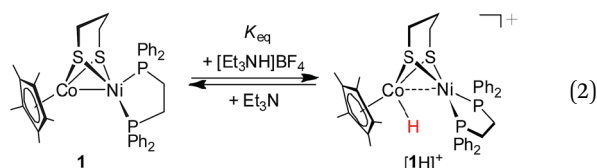


Fig. 5 Structure of  $[\mathbf{1H}]^+$ . Selected bond distances (Å): Co–Ni, 2.556(1); Co–H, 1.45(5); Co–S1, 2.22(1); Co–S2, 2.225(1); Ni–S1, 2.222(1); Ni–S2, 2.247(1).



integrated hydride signal vs. time suggests that the reaction follows unimolecular kinetics and has a rate constant of  $5.023 \times 10^{-5} \text{ s}^{-1}$  at  $25^\circ \text{C}$  ( $t_{1/2} = 3.83 \text{ h}$ , Fig. S16†).



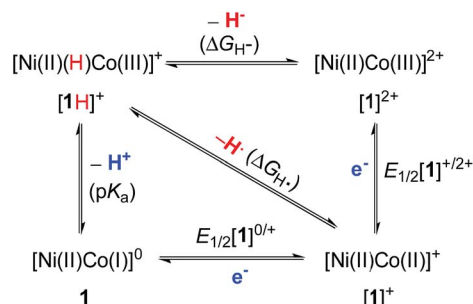
This bimetallic  $[\text{Ni(II)Co(III)H}]^+$  hydride was found to be much less acidic than the reported  $[\text{Ni(II)(H)Fe(II)}]^+$  hydrides such as  $(\text{CO})_3\text{Fe}(\text{pdt})\text{Ni}(\text{dppe})$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 10.7$ ) and  $(\text{CO})_2(\text{PPh}_3)\text{Fe}(\text{pdt})\text{Ni}(\text{dppe})$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 14.9$ ),<sup>30a</sup> and the diiron bridging hydride  $[(\mu\text{-H})\text{Fe}(\text{pdt})(\text{PMe}_3)_2(\text{CO})_4]^+$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 12$ ).<sup>47</sup> The acidity of  $[\text{1H}]^+$  was further evaluated by the protonation of **1** with  $[\text{Et}_3\text{NH}]\text{BF}_4$  ( $\text{p}K_{\text{a}}^{\text{MeCN}} = 18.8$ ).<sup>44</sup> An equilibrium was established between **1** and 1.0 equiv. of  $[\text{Et}_3\text{NH}]\text{BF}_4$  in PhCN (eqn (2)), which provided an equilibrium mixture of  $[\text{1H}]^+$  and **1** in a ratio of 0.22 : 1 as determined from the  $^{31}\text{P}$  NMR spectrum. A  $\text{p}K_{\text{a}}$  of 17.5 was estimated for  $[\text{1H}]^+$  on the basis of the calculated equilibrium constant  $K_{\text{eq}}$  (0.05) (Fig. S17†). The results indicate that the acidity of  $[\text{1H}]^+$  is weak, while its conjugated base, the  $\text{Ni(II)Co(I)}$  complex, exhibits a strong basic character.

### Homolytic and heterolytic bond energies for NiCo–H

Because of the centrality of metal hydrides in catalytic  $\text{H}_2$  production, we determined the homolytic and heterolytic bond energies for NiCo–H in an effort to learn more about the thermodynamic properties of the  $[\text{Ni(II)(H)Co(III)}]^+$  hydride. The cleavage of the M–H bond can involve transfer of electrons, a proton, a hydrogen atom or a hydride<sup>48–51</sup> and is very dependent on the bond energy. Scheme 2 illuminates the relationship between acidity ( $\text{p}K_{\text{a}}$ ), homolytic bond dissociation energy (BDFE,  $\Delta G_{\text{H}^\bullet}$ ), thermodynamic hydricity ( $\Delta G_{\text{H}^-}$ ), and redox potentials ( $E_{1/2}$ ) for the NiCo–H cleavage.

$$\Delta G_{\text{H}^\bullet} = 1.37 \text{ p}K_{\text{a}} + 23.06 E_{1/2}[\text{1}]^{0/+} + 54.9 \quad (3)$$

$$\Delta G_{\text{H}^-} = 1.37 \text{ p}K_{\text{a}} + 23.06 E_{1/2}[\text{1}]^{0/+} + 23.06 E_{1/2}[\text{1}]^{+/2+} + 79.6 \quad (4)$$



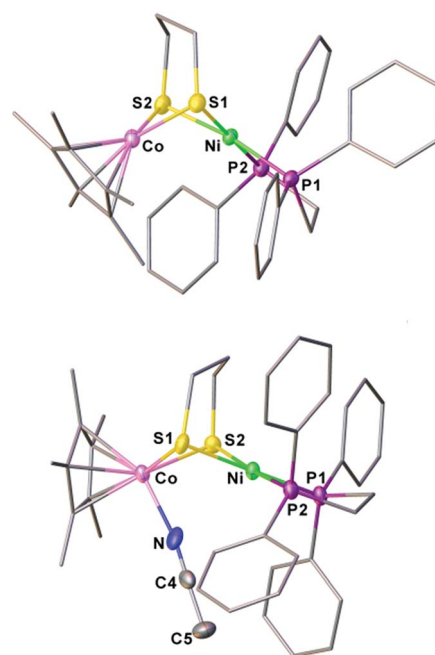
**Scheme 2** Schematic pathways to the cleavage of the NiCo–H bond in  $[\text{1H}]^+$ .

According to the thermodynamic cycle, the redox potentials of the  $[\text{1}]^{0/+}$  and  $[\text{1}]^{+/2+}$  couples are required to calculate  $\Delta G_{\text{H}^\bullet}$  and  $\Delta G_{\text{H}^-}$ . Cyclic voltammetry on PhCN solutions of **1** indicates oxidations at  $E_{1/2} = -0.45 \text{ V}$  ( $i_{\text{pa}}/i_{\text{pc}} = 1$ ) and  $E_{1/2} = -0.01 \text{ V}$  (quasi-reversible, Fig. S19†). These couples are assigned as one-electron processes for  $[\text{1}]^{0/+}$  and  $[\text{1}]^{+/2+}$ , respectively. The direct evidence is derived from oxidations conducted on a preparative scale (see below). Using the thermodynamic equations derived from eqn (3) and (4),<sup>30a,48,49</sup>  $\Delta G_{\text{H}^\bullet}$  and  $\Delta G_{\text{H}^-}$  are calculated to be 69 and 93  $\text{kcal mol}^{-1}$ , respectively.

### Oxidizing **1** for $\text{H}_2$ activation

Oxidation of **1** with one equiv. of  $\text{FcBF}_4$  provided a cationic complex  $[\text{1}]^+$ , while using  $\text{AgBF}_4$  ( $E_{1/2} = 0.65 \text{ V}$ ,  $\text{CH}_2\text{Cl}_2$ )<sup>52</sup> allowed for the further oxidation of  $[\text{1}]^+$  to  $[\text{1}]^{2+}$ . The formation of a new diamagnetic species was signaled by a  $^{31}\text{P}$  peak at  $\delta 59.5$  (s) (Fig. S20†). As a  $32e^-$  bimetallic species,  $[\text{1}]^{2+}$  is unstable and cannot be isolated. When the reaction was conducted in the presence of MeCN, the dicationic complex  $[\text{Ni(II)Co(III)}]^{2+}$  was stabilized in the form of  $[(\text{dppe})\text{Ni}(\text{pdt})(\text{MeCN})\text{CoCp}^*]^{2+}$  ( $[\text{1(NCMe)}]^{2+}$ ).

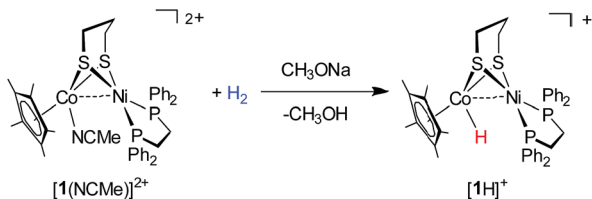
The structures of  $[\text{1}]^+$  and  $[\text{1(NCMe)}]^{2+}$  were characterized crystallographically. In both structures, the Ni center adopts a square-planar coordination geometry (Fig. 6). In  $[\text{1(NCMe)}]^{2+}$ , the MeCN is bound to the Co, consistent with electrochemical assignment that the second oxidation occurs at the  $\text{Cp}^*\text{Co}$  site. The  $1e^-$  oxidation increases the Ni...Co distance from 2.6572(10) Å for  $[\text{1}]^+$  to 3.005 Å for  $[\text{1(NCMe)}]^{2+}$ . The principal



**Fig. 6** Structures of  $[\text{1}]^+$  and  $[\text{1(NCMe)}]^{2+}$  with thermal ellipsoids drawn at the 50% probability level.  $\text{BF}_4^-$  anions, hydrogen atoms and solvent are omitted for clarity. Selected bond distances (Å): for  $[\text{1}]^+$ , Co–Ni, 2.657(1); Co–S1, 2.210(1); Co–S2, 2.215(1); Ni–S1, 2.239(1); Ni–S2, 2.229(1); for  $[\text{1(NCMe)}]^{2+}$ , Co–Ni, 3.005; Co–S1, 2.246(1); Co–S2, 2.263(1); Ni–S1, 2.237(1); Ni–S2, 2.239(1).





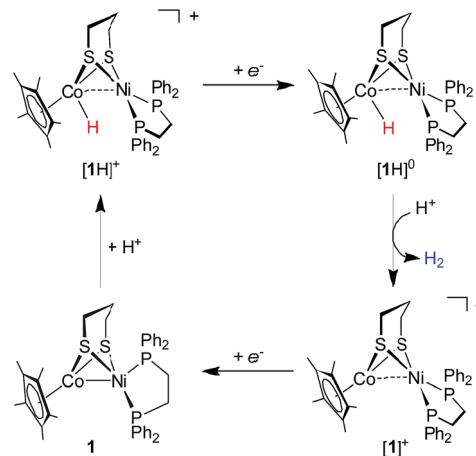
Scheme 3 Hydrogen activation by  $[1(\text{NCMe})]^{2+}$ .

change in the oxidation of **1** is the coordination sphere of nickel, which turns from roughly tetrahedral to a strictly square-planar geometry. According to the structures of **1** and  $[1]^+$ , the first oxidation event occurs at the Ni(dppe) site. Related to **1**, the Ni...Co distance in  $[1]^+$  is longer by 0.18 Å, meanwhile the Cp\*-Co distance increases slightly from 1.686 Å to 1.693 Å.

The high affinity of the dicationic  $[\text{Ni}(\text{II})\text{Co}(\text{III})]^{2+}$  complex toward the hydride was further demonstrated by the reaction of activating  $\text{H}_2$  (Scheme 3). The MeCN solution of  $[1(\text{NCMe})]^{2+}$  contained in a Schlenk flask was bubbled with  $\text{H}_2$  (10 psi) for 3 min, and then a solution of  $\text{CH}_3\text{ONa}$  in MeOH was added (Fig. S24†). Over the course of 10 min at room temperature, the reaction mixture turns from red to dark brown. As confirmed by  $^1\text{H}$  NMR and  $^{31}\text{P}$  NMR spectra, the organometallic product is  $[1\text{H}]^+$ . Interestingly, in addition to  $\text{CH}_3\text{ONa}$ , alkali metal carbonates such as  $\text{Na}_2\text{CO}_3$  can be employed as the base to assist  $[1(\text{NCMe})]^{2+}$  to activate  $\text{H}_2$ .

### Electrocatalytic $\text{H}_2$ production

Cyclic voltammetry indicates that  $[1\text{H}]^+$  only undergoes a reversible reduction process at  $-1.36$  V ( $i_{\text{pc}}/i_{\text{pa}} = 0.98$ ), which is tentatively assigned to the  $[\text{Ni}(\text{II})\text{Co}(\text{III})\text{H}]^+ / [\text{Ni}(\text{II})\text{Co}(\text{II})\text{H}]^0$  couple (Fig. S26†). To evaluate the catalytic activity of  $[1\text{H}]^+$  as an electrocatalyst for  $\text{H}_2$  production,  $\text{Cl}_2\text{CHCOOH}$  ( $E^0 = -0.92$  V)<sup>45</sup> was selected as the proton source. Upon the addition of

Scheme 4 Proposed catalytic cycle for proton reduction by  $[1\text{H}]^+$ .

$\text{Cl}_2\text{CHCOOH}$  to the  $\text{CH}_2\text{Cl}_2$  solution of  $[1\text{H}]^+$ , the reduction events for  $[1\text{H}]^{+/0}$  became irreversible, and the cathodic current intensity increased linearly with sequential increasing of the acid concentration (Fig. 7, inset). These observations are consistent with the aspects of proton reduction catalysis.<sup>53,54</sup> Plots of  $i_{\text{c}}/i_{\text{p}}$  vs.  $[\text{Cl}_2\text{CHCOOH}]$  are linear up to 218 equiv. of acid, indicating that the catalysis is second order with respect to the acid. The turnover frequency was estimated to be  $244 \text{ s}^{-1}$  (Table S1†). In the controlled experiment, proton reduction with a glassy carbon electrode was performed at a potential nearly 0.37 V more negative than that in the catalysis performed with  $[1\text{H}]^+$  (Fig. S25†).

To gain insight into the catalytic mechanism, we examined the chemical reduction of  $[1\text{H}]^+$  with  $\text{Cp}^*\text{Co}$  ( $E_{1/2} = -1.94$  V in  $\text{CH}_2\text{Cl}_2$ ). The reaction was indicated by an immediate change of the color of the solution from brown to black. Efforts to isolate and characterize the reduced  $[1\text{H}]^0$  were unsuccessful since it is converted to **1** and loses  $\text{H}_2$  in a matter of minutes. Addition of protic acids such as  $\text{Cl}_2\text{CHCOOH}$  or  $\text{H}(\text{OEt})_2\text{BAr}^{\text{F}}_4$  into a solution of  $[1\text{H}]^0$  in THF, generated *in situ* by reaction of  $[1\text{H}]^+$  with  $\text{Cp}^*\text{Co}$ , resulted in the formation of  $[1]^+$  and release of  $\text{H}_2$ . The yield of  $\text{H}_2$ , quantified by GC analysis, is  $91\% \pm 5$  (in three experiments) and is close to the stoichiometric value. We propose that the catalytic  $\text{H}_2$  evolution is based on the reduction of the  $[\text{Ni}(\text{II})\text{Co}(\text{III})\text{H}]^+$  hydride, operating through an ECEC mechanism (Scheme 4).

### Conclusions

A class of new NiCo complexes, relevant to the active site models of  $[\text{NiFe}]\text{-H}_2\text{ases}$ , have been studied and their features of stereodynamics, acid-base properties, redox chemistry and reactivity are disclosed. Incorporating the (dppe)Ni(pdt) module with a  $\text{Cp}^*\text{Co}$  fragment enables the bimetallic NiCo complexes to achieve Co-centered  $\text{H}_2$  evolution and uptake. Such a bimetallic way can disperse the strong effects of redox over the two metal sites.<sup>29</sup> Owing to the redox flexibility of Ni(dppe), distortion at the Ni center of the reduced state affects the oxidation state of the bimetallic centers, notably interconversion of Ni(II)

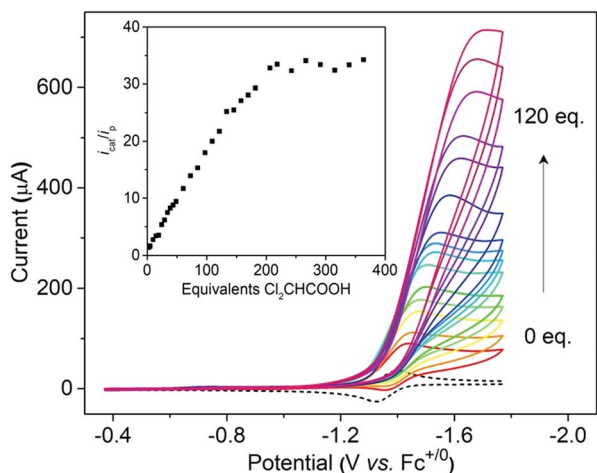


Fig. 7 Cyclic voltammograms of  $[1\text{H}]^+$  with increments of  $\text{Cl}_2\text{CHCOOH}$  (from 0 to 120 equiv.), inset: plot of catalytic current  $i_{\text{c}}/i_{\text{p}}$  for  $[1\text{H}]^+$  vs. equivalent of  $\text{Cl}_2\text{CHCOOH}$ . Conditions: 1 mM  $[1\text{H}]^+$  in MeCN, 0.1 M  $n\text{-NBu}_4\text{PF}_6$ , scan rate =  $100 \text{ mV s}^{-1}$ , and V vs.  $\text{Fc}^{+/0}$ .



Co(II) and Ni(II)Co(I). The protonation reaction can proceed *via* the mixed valence isomer Ni(II)Co(I) with enhanced basicity to afford  $[\text{Ni(II)Co(III)H}]^+$ . The bimetallic hydride is catalytically active and was demonstrated to be an efficient electrocatalyst for the reduction of weak protic acid at a mild potential. Despite a stoichiometric reaction, the oxidized state  $[\text{Ni(II)Co(III)}]^{2+}$  is capable of activating  $\text{H}_2$  to give the hydride.

Bimetallic hydride species are commonly encountered in hydrogenase modeling. Bimetallic active sites in hydrogenases provide an elegant means of softening the effect of redox reactions on the acid–base properties of the hydride.<sup>30a</sup> In addition to biocatalysis, metal hydrides play a central role in a variety of chemical transformations.<sup>55–59</sup> Studies of metal hydride chemistry are essential to control the elementary steps for catalysis.<sup>60–62</sup> However, the related thermodynamic properties of bimetallic systems have been inadequately examined to date.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We gratefully acknowledge the “Thousand Plan” Youth Program and the National Natural Science Foundation of China (21871166 and 91427303) for the financial support. We also thank Prof. Di Sun for assistance with the X-ray crystallography.

## Notes and references

- M. Iglesias, E. Sola and L. A. Oro, in *Homo- and Heterobimetallic Complexes in Catalysis: Cooperative Catalysis*, ed. P. Kalck, Springer International Publishing, Cham, Switzerland, 2016, ch. 2, pp. 31–58.
- F. A. Cotton, *Acc. Chem. Res.*, 1978, **11**, 225–232.
- L. H. Gade, *Angew. Chem., Int. Ed.*, 2000, **39**, 2658–2678.
- D. G. Huang and R. H. Holm, *J. Am. Chem. Soc.*, 2010, **132**, 4693–4701.
- I. G. Powers and C. Uyeda, *ACS Catal.*, 2017, **7**, 936–958.
- M. J. Page, D. B. Walker and B. A. Messerle, in *Homo- and Heterobimetallic Complexes in Catalysis: Cooperative Catalysis*, ed. P. Kalck, Springer International Publishing, Cham, Switzerland, 2016, ch. 4, pp. 103–138.
- J. I. van der Vlugt, *Eur. J. Inorg. Chem.*, 2012, **3**, 363–375.
- A. M. Appel, J. E. Bercaw, A. B. Bocarsly, H. Dobbek, D. L. DuBois, M. Dupuis, J. G. Ferry, E. Fujita, R. Hille, P. J. A. Kenis, C. A. Kerfeld, R. H. Morris, C. H. F. Peden, A. R. Portis, S. W. Ragsdale, T. B. Rauchfuss, J. N. H. Reek, L. C. Seefeldt, R. K. Thauer and G. L. Waldrop, *Chem. Rev.*, 2013, **113**, 6621–6658.
- R. Pulukkody and M. Y. Darensbourg, *Acc. Chem. Res.*, 2015, **48**, 2049–2058.
- P. V. Rao, S. Bhaduri, J. F. Jiang and R. H. Holm, *Inorg. Chem.*, 2004, **43**, 5833–5849.
- (a) V. Artero, G. Berggren, M. Atta, G. Caserta, S. Roy, L. Pecqueur and M. Fontecave, *Acc. Chem. Res.*, 2015, **48**, 2380–2387; (b) C. H. Hsieh, S. Ding, Ö. F. Erdem, D. J. Crouthers, T. Liu, C. C. Mccrory, W. Lubitz, C. V. Popescu, J. H. Reibenspies, M. B. Hall and M. Y. Darensbourg, *Nat. Commun.*, 2014, **5**, 3684.
- M. Can, F. A. Armstrong and S. W. Ragsdale, *Chem. Rev.*, 2014, **114**, 4149–4174.
- P. P. Liebgott, A. L. de Lacey, B. Burlat, L. Cournac, P. Richaud, M. Brugna, V. M. Fernandez, B. Guigliarelli, M. Rousset, C. Leger and S. Dementin, *J. Am. Chem. Soc.*, 2011, **133**, 986–997.
- W. Lubitz, H. Ogata, O. Rudiger and E. Reijerse, *Chem. Rev.*, 2014, **114**, 4081–4148.
- B. L. Greene, C. H. Wu, G. E. Vansuch, M. W. W. Adams and R. B. Dyer, *Biochemistry*, 2016, **55**, 1813–1825.
- (a) P. A. Ash, R. Hidalgo and K. A. Vincent, *ACS Catal.*, 2017, **7**, 2471–2485; (b) B. J. Murphy, R. Hidalgo, M. M. Roessler, R. M. Evans, P. A. Ash, W. K. Myers, K. A. Vincent and F. A. Armstrong, *J. Am. Chem. Soc.*, 2015, **137**, 8484–8489; (c) R. Hidalgo, P. A. Ash, A. J. Healy and K. A. Vincent, *Angew. Chem., Int. Ed.*, 2015, **54**, 7110–7113.
- W. Zhu, A. C. Marr, Q. Wang, F. Neese, D. J. E. Spencer, A. J. Blake, P. A. Cooke, C. Wilson and M. Schroder, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 18280–18285.
- (a) S. Canaguier, V. Fourmond, C. U. Perotto, J. Fize, J. Pecaut, M. Fontecave, M. J. Field and V. Artero, *Chem. Commun.*, 2013, **49**, 5004–5006; (b) Y. Oudart, V. Artero, J. Pecaut and M. Fontecave, *Inorg. Chem.*, 2006, **45**, 4334–4336; (c) S. Canaguier, M. Field, Y. Oudart, J. Pecaut, M. Fontecave and V. Artero, *Chem. Commun.*, 2010, **46**, 5876–5878; (d) T. R. Simmons, G. Berggren, M. Bacchi, M. Fontecave and V. Artero, *Coord. Chem. Rev.*, 2014, **270**, 127–150.
- C. Tard and C. J. Pickett, *Chem. Rev.*, 2009, **109**, 2245–2274.
- Y. Ohki, K. Yasumura, M. Ando, S. Shimokata and K. Tatsumi, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 3994–3997.
- K. Weber, O. F. Erdem, E. Bill, T. Weyhermuller and W. Lubitz, *Inorg. Chem.*, 2014, **53**, 6329–6337.
- (a) D. Schilter, T. B. Rauchfuss and M. Stein, *Inorg. Chem.*, 2012, **51**, 8931–8941; (b) D. Schilter, M. J. Nilges, M. Chakrabarti, P. A. Lindahl, T. B. Rauchfuss and M. Stein, *Inorg. Chem.*, 2012, **51**, 2338–2348.
- X. Chu, X. Yu, S. Raje, R. Angamuthu, J. Ma, C.-H. Tung and W. Wang, *Dalton Trans.*, 2017, **46**, 13681–13685.
- (a) P. Ghosh, S. D. Ding, R. B. Chupik, M. Quiroz, C. H. Hsieh, N. Bhuvanesh, M. B. Hall and M. Y. Darensbourg, *Chem. Sci.*, 2017, **8**, 8291–8300; (b) S. D. Ding, P. Ghosh, M. Y. Darensbourg and M. B. Hall, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, E9775–E9782; (c) S. Ding, P. Ghosh, A. M. Lunsford, N. Wang, N. Bhuvanesh, M. B. Hall and M. Y. Darensbourg, *J. Am. Chem. Soc.*, 2016, **138**, 12920–12927.
- D. Brazzolotto, M. Gennari, N. Queyriaux, T. R. Simmons, J. Pecaut, S. Demeshko, F. Meyer, M. Orio, V. Artero and C. Duboc, *Nat. Chem.*, 2016, **8**, 1054–1060.
- S. Ogo, K. Ichikawa, T. Kishima, T. Matsumoto, H. Nakai, K. Kusaka and T. Ohhara, *Science*, 2013, **339**, 682–684.



- 27 B. C. Manor and T. B. Rauchfuss, *J. Am. Chem. Soc.*, 2013, **135**, 11895–11900.
- 28 L. Song, X. Yang, M. Cao, X. Gao, B. Liu, L. Zhu and F. Jiang, *Chem. Commun.*, 2017, **53**, 3818–3821.
- 29 D. Schilter, J. M. Camara, M. T. Huynh, S. Hammes-Schiffer and T. B. Rauchfuss, *Chem. Rev.*, 2016, **116**, 8693–8749.
- 30 (a) B. E. Barton and T. B. Rauchfuss, *J. Am. Chem. Soc.*, 2010, **132**, 14877–14885; (b) M. T. Huynh, D. Schilter, S. Hammes-Schiffer and T. B. Rauchfuss, *J. Am. Chem. Soc.*, 2014, **136**, 12385–12395; (c) O. A. Ulloa, M. T. Huynh, C. P. Richers, J. A. Bertke, M. J. Nilges, S. Hammes-Schiffer and T. B. Rauchfuss, *J. Am. Chem. Soc.*, 2016, **138**, 9234–9245.
- 31 (a) J. A. Denny and M. Y. Darensbourg, *Chem. Rev.*, 2015, **115**, 5248–5273; (b) M. V. Rampersad, S. P. Jeffery, J. H. Reibenspies, C. G. Ortiz, D. J. Darensbourg and M. Y. Darensbourg, *Angew. Chem.*, 2005, **117**, 1243–1246; (c) K. N. Green, S. P. Jeffery, J. H. Reibenspies and M. Y. Darensbourg, *J. Am. Chem. Soc.*, 2006, **128**, 6493–6498.
- 32 (a) M. Schmidt and G. G. Hoffmann, *Eur. J. Inorg. Chem.*, 1979, **112**, 2190–2196; (b) M. Schmidt and G. G. Hoffmann, *J. Organomet. Chem.*, 1977, **124**, 5–8.
- 33 (a) G. M. Chambers, J. Mitra, T. B. Rauchfuss and M. Stein, *Inorg. Chem.*, 2014, **53**, 4243–4249; (b) 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.
- 34 V. Fourmond, S. Canaguier, B. Golly, M. J. Field, M. Fontecave and V. Artero, *Energy Environ. Sci.*, 2011, **4**, 2417–2427.
- 35 D. Schilter, A. L. Fuller and D. L. Gray, *Eur. J. Inorg. Chem.*, 2015, **28**, 4638–4642.
- 36 X. Chu, X. Xu, H. Su, S. Raje, R. Angamuthu, C.-H. Tung and W. Wang, *Inorg. Chem. Front.*, 2017, **4**, 706–711.
- 37 A. M. Lunsford, K. F. Goldstein, M. A. Cohan, J. A. Denny, N. Bhuvanesh, S. D. Ding, M. B. Hall and M. Y. Darensbourg, *Dalton Trans.*, 2017, **46**, 5175–5182.
- 38 (a) S. Ogo, R. Kabe, K. Uehara, B. Kure, T. Nishimura, S. C. Menon, R. Harada, S. Fukuzumi, Y. Higuchi, T. Ohhara, T. Tamada and R. Kuroki, *Science*, 2007, **316**, 585–587; (b) T. Matsumoto, B. Kure and S. Ogo, *Chem. Lett.*, 2008, **37**, 970–971.
- 39 J. L. Dempsey, B. S. Brunschwig, J. R. Winkler and H. B. Gray, *Acc. Chem. Res.*, 2009, **42**, 1995–2004.
- 40 W. Liu, B. Sahoo, K. Junge and M. Beller, *Acc. Chem. Res.*, 2018, **51**, 1858–1869.
- 41 P. J. Chirik, *Acc. Chem. Res.*, 2015, **48**, 1687–1695.
- 42 H. Han, Z. Wei, M. C. Barry, J. C. Carozza, M. Alkan, A. Y. Rogachev, A. S. Filatov, A. M. Abakumov and E. V. Dikarev, *Chem. Sci.*, 2018, **9**, 4736–4745.
- 43 B. Cordero, V. Gomez, A. E. Platero-Prats, M. Reves, J. Echeverria, E. Cremades, F. Barragan and S. Alvarez, *Dalton Trans.*, 2008, 2832–2838.
- 44 I. Kaljurand, A. Kutt, L. Soovali, T. Rodima, V. Maemets, I. Leito and I. A. Koppel, *J. Org. Chem.*, 2005, **70**, 1019–1028.
- 45 G. A. N. Felton, R. S. Glass, D. L. Lichtenberger and D. H. Evans, *Inorg. Chem.*, 2006, **45**, 9181–9184.
- 46 (a) F. Zhang, X. Xu, Y. Zhao, J. Jia, C.-H. Tung and W. Wang, *Organometallics*, 2017, **36**, 1238–1244; (b) J. Liu, F. Zhang, A. Zhang, Q. Tong, C.-H. Tung and W. Wang, *Chem.-Asian J.*, 2016, **11**, 2271–2277; (c) F. Zhang, J. Jia, S. Dong, W. Wang and C.-H. Tung, *Organometallics*, 2016, **35**, 1151–1159.
- 47 G. Eilers, L. Schwartz, M. Stein, G. Zampella, L. de Gioia, S. Ott and R. Lomoth, *Chem.-Eur. J.*, 2007, **13**, 7075–7084.
- 48 Y. Hu, A. P. Shaw, D. P. Estes and J. R. Norton, *Chem. Rev.*, 2016, **116**, 8427–8462.
- 49 E. S. Wiedner, M. B. Chambers, C. L. Pitman, R. M. Bullock, A. J. M. Miller and A. M. Appel, *Chem. Rev.*, 2016, **116**, 8655–8692.
- 50 M. Bourrez, R. Steinmetz, S. Ott, F. Gloaguen and L. Hammarstrom, *Nat. Chem.*, 2015, **7**, 140–145.
- 51 Y. Matsubara, E. Fujita, M. D. Doherty, J. T. Muckerman and C. Creutz, *J. Am. Chem. Soc.*, 2012, **134**, 15743–15757.
- 52 N. G. Connelly and W. E. Geiger, *Chem. Rev.*, 1996, **96**, 877–910.
- 53 A. D. Wilson, R. H. Newell, M. J. McNevin, J. T. Muckerman, M. R. DuBois and D. L. DuBois, *J. Am. Chem. Soc.*, 2006, **128**, 358–366.
- 54 C. P. Andrieux, C. Blocman, J. M. Dumas-Bouchiat, F. M'Halla and J. M. Savéant, *J. Electroanal. Chem.*, 1980, **113**, 19–40.
- 55 M. Peruzzini and R. Poli, *Recent Advances in Hydride Chemistry*, Elsevier, Amsterdam, 2001.
- 56 R. Noyori and S. Hashiguchi, *Acc. Chem. Res.*, 1997, **30**, 97–102.
- 57 H. Nakazawa and M. Itazaki, *Iron Catalysis: Fundamentals and Applications*, 2011, vol. 33, pp. 27–81.
- 58 (a) R. M. Bullock, *Catalysis with precious metals*, Wiley-VCH, Weinheim, 2010; (b) R. M. Bullock, *Science*, 2013, **342**, 1054–1055.
- 59 S. Chakraborty, P. Bhattacharya, H. G. Dai and H. R. Guan, *Acc. Chem. Res.*, 2015, **48**, 1995–2003.
- 60 C. L. Pitman, K. R. Brereton and A. J. M. Miller, *J. Am. Chem. Soc.*, 2016, **138**, 2252–2260.
- 61 C. M. Zall, J. C. Linehan and A. M. Appel, *J. Am. Chem. Soc.*, 2016, **138**, 9968–9977.
- 62 (a) R. M. Henry, R. K. Shoemaker, D. L. DuBois and M. R. DuBois, *J. Am. Chem. Soc.*, 2006, **128**, 3002–3010; (b) T. B. Liu, D. L. DuBois and R. M. Bullock, *Nat. Chem.*, 2013, **5**, 228–233.

