

Scheme 1 Hypotheses for inhibition of [FeFe]-hydrogenases by  $\text{CH}_2\text{O}$ .

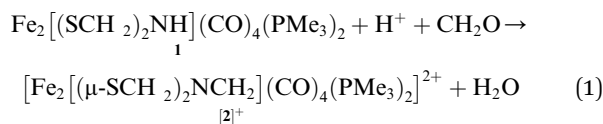
The amine pathways, which involve standard organic reactions, might be relevant to what Bachmeier *et al.* refer to as “matrix” formaldehyde, *i.e.*, unselective binding of formaldehyde in the vicinity of the active site. The Fe-centered reactions are preceded in organometallic chemistry, although not necessarily with iron. Bachmeier *et al.* favor this hypothesis. The third pathway, for which we provide evidence, involves covalent linking the amine and distal iron with a methylene bridge, locking up the  $[\text{2Fe}]_{\text{H}}$  active site.

The model complexes used in this paper are  $\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NH}](\text{CO})_4\text{L}_2$ , where  $\text{L} = \text{PMe}_3$  and  $\text{CN}^-$ . These models feature the authentic azadithiolate cofactor bound to a pair of  $\text{Fe}(\text{CO})_2\text{L}$  centers. Such complexes are functional models in that they undergo protonation to give hydrides and are redox-active.<sup>17</sup>

## Results and discussion



Solutions of  $\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NH}](\text{CO})_4(\text{PMe}_3)_2$  (**1**) were found to react with a mixture of  $\text{HBF}_4$  and paraformaldehyde to give  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NCH}_2](\text{CO})_4(\text{PMe}_3)_2]^+$  (**[2]<sup>+</sup>**). Using stoichiometric amounts of the three reagents, the conversion proceeds rapidly and in good yields at room temperature. These conditions are compatible with those reported for the enzyme. The formula of **[2]<sup>+</sup>** was initially determined by ESI-MS, which showed a strong parent ion (eqn (1)).



In a control experiment, the reaction of the propane-dithiolate  $\text{Fe}_2(\mu\text{-S}_2\text{C}_3\text{H}_6)(\text{CO})_4(\text{PMe}_3)_2$  with  $\text{HBF}_4$  and paraformaldehyde afforded only the well-known hydride  $[\text{HFe}_2(\mu\text{-S}_2\text{C}_3\text{H}_6)(\text{CO})_4(\text{PMe}_3)_2]^+$ ,<sup>18</sup> the formaldehyde had no effect.

The structure of **[2]<sup>+</sup>** was determined by an X-ray crystallographic study of its  $\text{BAR}^{\text{F}}_4^-$  salt ( $\text{Ar}^{\text{F}} = \text{C}_6\text{H}_3\text{-3,5-(CF}_3)_2$ ) (Fig. 2). The two  $\text{PMe}_3$  ligands are *trans*-dibasal. Fe1 has an  $\text{S}_2(\text{CO})_2(\text{-PMe}_3)(\text{alkyl})$  coordination sphere. The ligand-Fe1-ligand angles are suitable for octahedral coordination, as appropriate for Fe(II). The proximal Fe center Fe2 occupies a  $\text{S}_2(\text{CO})_2(\text{PMe}_3)$  coordination sphere. Its coordination number is ambiguous

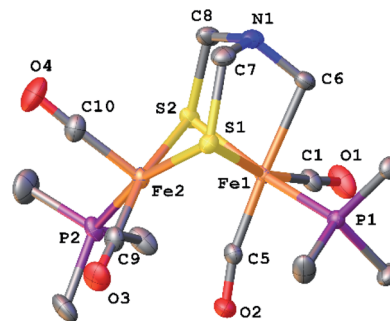


Fig. 2 Structure of  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NCH}_2](\text{CO})_4(\text{PMe}_3)_2]\text{BAR}^{\text{F}}_4^-$  with thermal ellipsoids shown at 50% probability. H atoms and  $\text{BAR}^{\text{F}}_4^-$  have been omitted for clarity. Selected distances and angles ( $\text{\AA}$  and  $^\circ$ ): Fe1–Fe2, 2.5934(3); Fe1–C5, 1.796(2); Fe2–C5, 2.559(2); Fe1–C6, 2.168(2); N1–C6, 1.441(2); N1–C7, 1.431(3); N1–C8, 1.432(2); Fe1–C5–O2, 167.8(2); Fe1–C1–O1, 177.9(2).

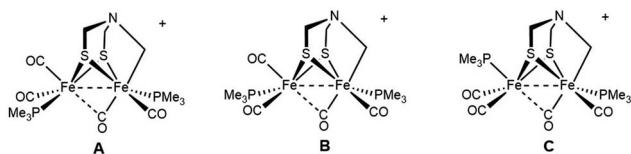
because one CO, primarily bound to Fe1, is semi-bridging: Fe1–CO = 1.796(2) and Fe2–CO = 2.559(2)  $\text{\AA}$ . The Fe1–C5–O2 angle for the semi-bridging CO ligand is 167.8(2) $^\circ$ , suggesting that Fe2 is weakly Lewis acidic. Analogous to terminal hydride derivatives of  $\text{Fe}_2^{\text{II}}(\mu\text{-SR})_2$  complexes,<sup>19,20</sup> the ligand *trans* to alkyl is CO. In a related thioaldehyde complexes<sup>21</sup>  $\text{Fe}_2(\mu\text{-SR})(\mu\text{-}\eta^2\text{-SCHR}^{\prime})(\text{diphosphine})(\text{CO})_4$ , CO is also *trans* to alkyl.<sup>22</sup>

The NMR data for **[2]<sup>+</sup>** are consistent with a stereo-rigid, chiral structure. For example, the  $^{13}\text{C}$  NMR spectrum shows four CO signals, three signals in the  $\delta$  211.45–210.70 region assigned to terminal CO groups, and one signal at  $\delta$  201.8 assigned to the semi-bridging CO. These  $^{13}\text{C}$  CO signals are all coupled to  $^{31}\text{P}$  ( $J_{\text{PC}} = 19.7$  Hz), characteristic of CO *cis* to  $\text{PMe}_3$ .<sup>23</sup> The  $^{13}\text{C}$  NMR signal for the formaldehyde-derived methylene appears at  $\delta$  75.77. Its  $^1\text{H}$  NMR spectrum shows multiplets at  $\delta$  5.52 and 4.79, assigned to the diastereotopic  $\text{CH}_2$  protons. The four  $\text{SCH}_2\text{N}$  protons are nonequivalent, also consistent with the low symmetry of the complex. Spin-saturation transfer experiments, which probes the exchange of signals at rates faster than  $1/T_1$ , were conducted on **[2]<sup>+</sup>**. Saturation of one of the  $\text{SCH}_2$  signals centered at  $\delta$  3.84 ( $T_1 = 1.17$  s for  $\text{SCH}_2$ ) or one of the  $\text{NCH}_2\text{Fe}$  signals at  $\delta$  4.79 ( $T_1 = 1.31$  s for  $\text{NCH}_2\text{Fe}$ ) revealed that these sites do not exchange on the seconds time scale (Fig. S14 and S15<sup>†</sup>). As discussed below,  $^{13}\text{C}$ -labeling reveals that all three methylene groups do in fact exchange over the course of several minutes.

The  $^{31}\text{P}$  NMR data for **[2]<sup>+</sup>** reveal the presence of a mixture of three (chiral) diastereoisomers in a 5 : 1 : 1 ratio. The minor diastereoisomers are not evident in the above-discussed  $^{13}\text{C}$  NMR data. If we assume that Fe1 center has CO *trans* to the alkyl ligand, as mentioned above, the three isomers result from the three diastereomeric sites on Fe2:

We assume that the main isomer (A) has *trans*-dibasal phosphine ligands as established by X-ray crystallography. This dominant and one minor isomer (B) both show  $^{31}\text{P}$ – $^{31}\text{P}$  coupling (respectively,  $J = 7.4, 7.6$  Hz). The two  $^{31}\text{P}$  NMR signals for the third isomer (minor, C) show no  $^{31}\text{P}$ ,  $^{31}\text{P}$  coupling. Its unique (non)coupling is consistent with a unique structure, *i.e.*, apical–basal disposition of the  $\text{PMe}_3$  ligands.



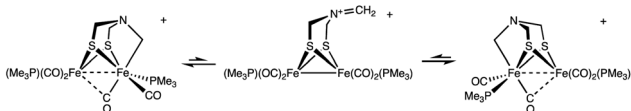


The entirety of the NMR data is accommodated by an exchange process involving reversible scission of the Fe–C bond, concomitant with regeneration of an Fe(I)Fe(I) species. Scission of the CH<sub>2</sub>–Fe bond introduces an effective plane of symmetry such that the two Fe(I) centers become equivalent (Scheme 2). Further relevant to stereodynamics, the exchange for the FeL<sub>3</sub> sites is rapid in Fe(I)Fe(I) complexes, whereas bioctahedral Fe(II)Fe(II) complexes are more rigid.<sup>24</sup>

Evidence of the process shown in Scheme 2 is provided by <sup>31</sup>P NMR spin saturation experiments. The *T*<sub>1</sub> of the signal at  $\delta$  9.45 was determined to be 8.2 s, and the exchange rate was *k* = 0.85 s<sup>-1</sup>. Saturation of either of the signals at  $\delta$  22.45 or 9.45 resulted in collapse of the other five <sup>31</sup>P NMR signals (Fig. 3).

### Mechanistic studies

Since Brønsted acids are required for the conversion of **1** to [2]<sup>+</sup>, we examined the ammonium complex [Fe<sub>2</sub>[( $\mu$ -SCH<sub>2</sub>)<sub>2</sub>-NH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]<sup>+</sup> ([1H]<sup>+</sup>). These results, which overlap with those reported earlier by Pickett,<sup>25</sup> are rather fundamental and merit thorough analysis. <sup>1</sup>H and <sup>31</sup>P NMR spectra confirm that [1H]<sup>+</sup> is stable in solution for days. Treating a CH<sub>2</sub>Cl<sub>2</sub> solution of [1H]BF<sub>4</sub> with NaBAR<sub>4</sub><sup>F</sup> induced tautomerization to the hydride [HFe<sub>2</sub>[( $\mu$ -SCH<sub>2</sub>)<sub>2</sub>NH](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]<sup>+</sup> ([H1]<sup>+</sup>). According to IR measurements, the tautomerization is complete after 3 h at room temperature (eqn (2)).



Scheme 2 Proposed stereodynamics for [2]<sup>+</sup>. The process would proceed with racemization.

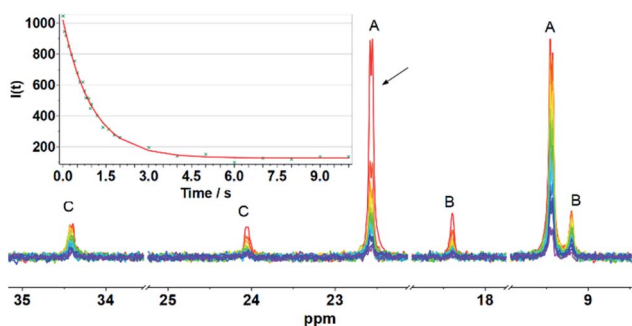
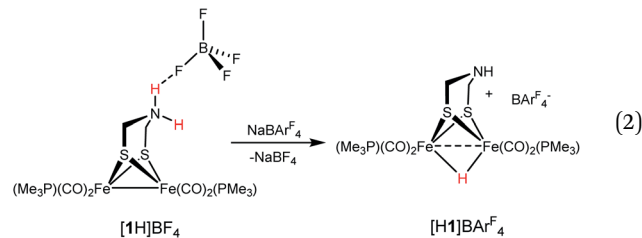


Fig. 3 <sup>31</sup>P NMR spin saturation transfer spectra of [2]BAR<sub>4</sub><sup>F</sup> at 298 K in CD<sub>2</sub>Cl<sub>2</sub>. Irradiation of the signal at  $\delta$  22.59. *T*<sub>1</sub> = 8.2 s for the resonance at  $\delta$  9.45. Inset: graph of intensity (*I*) of the  $\delta$  9.45 peak vs. irradiation time at  $\delta$  22.59. Fitting:  $I_t = I_0 \times \{1/(1 + \tau/8.2)\} \times \exp[-t \times (1/8.2 + 1/\tau)] + 1/(1 + 8.2/\tau)$ , where  $\tau = 1.17$  s, *k* = 1/ $\tau$  = 0.85 s<sup>-1</sup>.



The <sup>1</sup>H NMR spectrum for [H1]<sup>+</sup> matches published data for related salts.<sup>18</sup> The anion-dependent tautomerization reflects the stabilization of ammonium centers by hydrogen-bonding to BF<sub>4</sub><sup>-</sup>, which persists in solution.<sup>26</sup> One consequence of the ion-pairing (or its absence in the case of BAR<sub>4</sub><sup>F-</sup>) is that the proton-induced reaction of **1** with CH<sub>2</sub>O is sensitive to the identity of the acid: H(OEt<sub>2</sub>)BF<sub>4</sub> cleanly gives [2]<sup>+</sup> but H(OEt<sub>2</sub>)<sub>2</sub>BAR<sub>4</sub><sup>F</sup>, depending on the specific conditions, can afford significant quantities of the hydride [H1]<sup>+</sup>.

X-ray crystallography verified the extensive hydrogen-bonding in solid [H1]<sup>+</sup> (Fig. 4). The asymmetric unit consists of three ion pairs; two cations have *trans*-dibasal phosphine ligands, one is apical-basal. All NH centers are hydrogen bonded to BF<sub>4</sub><sup>-</sup>. The F...N distances range from 1.97–2.54 Å with the average distance of 2.22 Å.<sup>27</sup>

The hydroxymethylation of secondary amines by formaldehyde is well studied.<sup>28</sup> When a solution of **1** was treated with CH<sub>2</sub>O in the absence of acid, only subtle shifts (<5 cm<sup>-1</sup>) were observed in the IR spectrum in the  $\nu_{CO}$  region. It is known, however, that  $\nu_{CO}$  is relatively insensitive to substituents on nitrogen of the amine. For example, in this work we found that the  $\nu_{CO}$  bands for Fe<sub>2</sub>[( $\mu$ -SCH<sub>2</sub>)<sub>2</sub>NR](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub> are almost identical for R = H (1983, 1943, 1899 cm<sup>-1</sup>) and R = Me (1983, 1945, 1909, 1894 cm<sup>-1</sup>). <sup>1</sup>H NMR spectroscopy proved to be a more sensitive indicator of the interaction of **1** and CH<sub>2</sub>O. A 1 : 0.25 mixture of these reactants generates ~25% of a new species that we assign to the hydroxymethyl derivative Fe<sub>2</sub>[( $\mu$ -

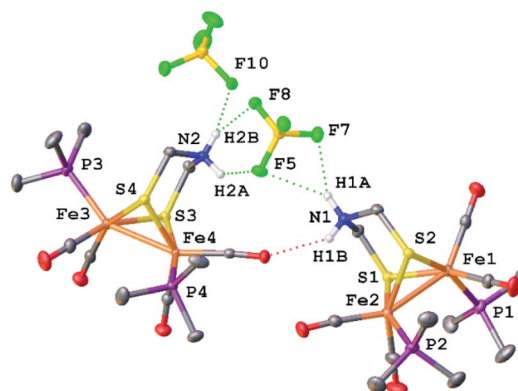


Fig. 4 Structure of [Fe<sub>2</sub>[( $\mu$ -SCH<sub>2</sub>)<sub>2</sub>NH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BF<sub>4</sub> with thermal ellipsoids shown at 50% probability. H atoms except for the NH<sub>2</sub> centres have been omitted for clarity. Two of the three ion pairs in the asymmetric unit are shown. Notice the presence of stereoisomers of [H1]<sup>+</sup>. Selected distances (Å): Fe1–Fe2, 2.5689(3); Fe3–Fe4, 2.5444(3); H1A–F5, 2.54(2); H1A–F7, 2.10(2); H2A–F5, 2.30(2); H2B–F8, 2.41(2); H2B–F10, 2.00(2).

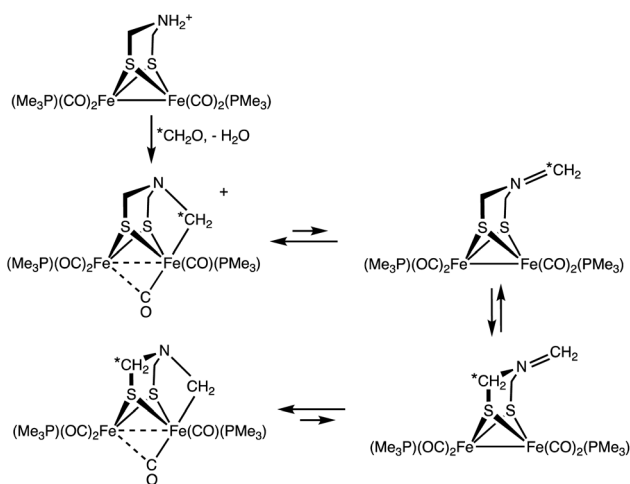


$\text{SCH}_2)_2\text{NCH}_2\text{OH}](\text{CO})_4(\text{PMe}_3)_2$ . The same species is observed with  $^{13}\text{CH}_2\text{O}$  under otherwise identical conditions. In that experiment, the  $\text{SCH}_2\text{N}$  groups did not show any enrichment. When **1** and  $\text{CH}_2\text{O}$  were mixed in a 1 : 1 ratio, several species are observed,  $\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NCH}_2\text{OH}](\text{CO})_4(\text{PMe}_3)_2$ , some unreacted **1**, and what appears to be  $\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-N}(\text{CH}_2\text{O})_n\text{CH}_2\text{OH}](\text{CO})_4(\text{PMe}_3)_2$ . Equilibration of these species is rapid, since the mixture reacts with 1 equiv. of  $\text{HBF}_4$  to cleanly give  $[2]^+$ . No reaction was evident when **1** was treated with  $\text{PhCHO}$ , in the presence or absence of  $\text{HBF}_4$ .

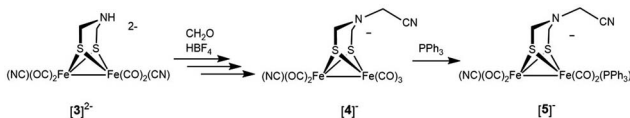
Treatment of a solution of  $^{13}\text{CH}_2\text{O}$  and **1** with  $\text{H}(\text{Et}_2\text{O})_2\text{BAR}^{\text{F}}_4$  gave  $[^{13}2]^+$  with selective formation of the  $\text{Fe-}^{13}\text{CH}_2$  isotopomer. Interestingly, this label exchanges with the other methylene groups in the complex over the course of hours (Scheme 3). The kinetics of exchange are first order in  $[2]^+$  up to about 90% conversion, which points to an intramolecular process. The NMR data show that this  $^{13}\text{CH}_2/^{12}\text{CH}_2$  exchange affects the diastereotopic  $\text{SCH}_2$  groups equally. We suggest that exchange occurs for the  $\text{Fe}(\text{I})\text{Fe}(\text{I})$  species where the diastereomerization is rapid. The ESI-MS of the product of the labelling shows only singly labelled  $[2]^+$ . Intermolecular processes would be expected to yield detectable levels of doubly labeled product.

#### Reaction of $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NH}](\text{CN})_2(\text{CO})_4]^{2-}$ with $\text{CH}_2\text{O}/\text{HBF}_4$

The reaction of paraformaldehyde with  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NH}](\text{CN})_2(\text{CO})_4]^{2-}$  ( $[3]^{2-}$ ) was investigated because this complex resembles the  $[2\text{Fe}]_{\text{H}}$  active site, which is also a dicyanide. In the presence of one equiv. of  $\text{HBF}_4$ ,  $[3]^{2-}$  converts to the ammonium derivative, which is stable in MeCN solution for several minutes



Scheme 3 Synthesis of  $[2]^+$  using  $^{13}\text{CH}_2\text{O}$ , showing site exchange.



Scheme 4 Pathway for formation of  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NCH}_2\text{CN}](\text{CN})(\text{CO})_4(\text{PPh}_3)]^-$  ( $[5]^-$ ).

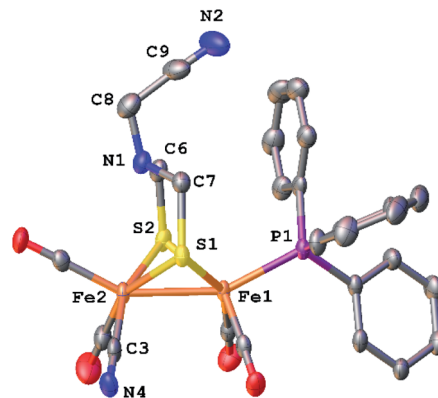


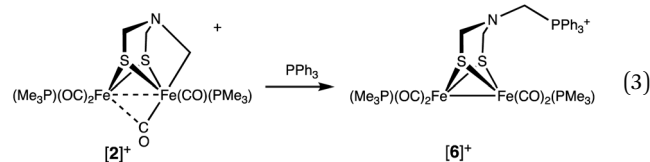
Fig. 5 Structure of  $\text{Et}_4\text{N}[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NCH}_2\text{CN}](\text{CN})(\text{CO})_4(\text{PPh}_3)]^-$  ( $\text{Et}_4\text{N}[5]$ ) with thermal ellipsoids shown at 50% probability. H atoms and the  $\text{Et}_4\text{N}^+$  cation have been omitted for clarity. Selected distances (Å):  $\text{Fe1-Fe2}$ , 2.5263(8);  $\text{Fe2-C3}$ , 1.928(5);  $\text{C3-N4}$ , 1.148(6);  $\text{C9-N2}$ , 1.144(8).

prior to irreversible tautomerization to the hydride (Fig. S39<sup>†</sup>). Addition of paraformaldehyde to  $[3\text{H}]^-$  gives one main product, which we assign as the pentacarbonyl  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NCH}_2\text{CN}](\text{CN})(\text{CO})_5]^-$  ( $[4]^-$ ). This formula is supported by ESI-MS analysis. Attempted purification of  $[4]^-$  was unsuccessful, however its FT-IR spectrum is very similar to that for  $[\text{Fe}_2[(\mu\text{-S}_2\text{-C}_6\text{H}_6)](\text{CN})(\text{CO})_5]^-$ . When  $^{13}\text{CH}_2\text{O}$  was used, the singly labeled product was generated according to ESI-MS. We propose that  $[4]^-$  arises by reductive elimination of the nitrile from  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NCH}_2](\text{CN})_2(\text{CO})_4]^-$  followed by CO-scavenging (Scheme 4).

The phosphine derivative of  $[4]^-$  was obtained when the reaction of  $[3]^{2-}$  with  $\text{CH}_2\text{O}$  was conducted in the presence of  $\text{PPh}_3$  (Scheme 4). The  $^{31}\text{P}$  NMR signal of this product at  $\delta$  60.1 indicates coordination of  $\text{PPh}_3$ , leading to the formula  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NCH}_2\text{CN}](\text{CN})(\text{CO})_4(\text{PPh}_3)]^-$  ( $[5]^-$ ). In the FT-IR spectrum of  $[5]^-$ , the  $\nu_{\text{CO}}$  bands are shifted by  $21\text{ cm}^{-1}$  toward high energy compared to dicyanide complex  $[3]^{2-}$ . The structure of  $[5]^-$  was verified by X-ray crystallography (Fig. 5), which confirms the presence of a conventional  $[\text{Fe}_2(\mu\text{-SR})_2(\text{CN})(\text{CO})_4(\text{PPh}_3)]^-$  complex,<sup>29</sup> and, most importantly, the presence of the cyanomethyl substituent.

#### Reactions of methylenated FeFe complex with nucleophiles

The  $\text{Fe-CH}_2\text{N}$  bond in  $[2]^+$  is electrophilic. For example, treating  $[2]^+$  with  $\text{PPh}_3$  cleanly gave  $[\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{-NCH}_2\text{PPh}_3](\text{CO})_4(\text{PMe}_3)_2]\text{BF}_4$  ( $[6]^+$ ), the result of C-P bond formation. Dealkylation of Fe involves reduction to an  $[\text{Fe}(\text{I})]_2$  complex (eqn (3)).



The  $\text{PMe}_3$  ligands in  $[6]^+$  appear equivalent, as is typical for related  $[\text{Fe}(\text{I})]_2$  complexes. The presence of a phosphonium



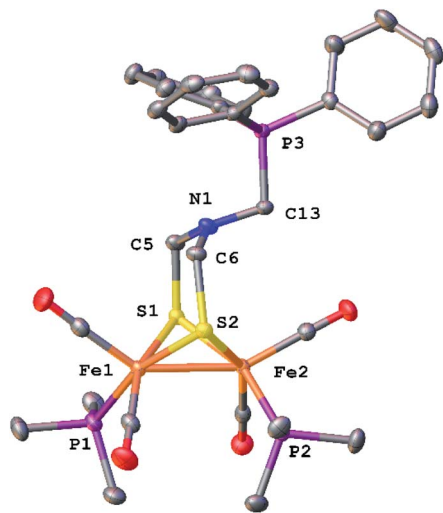
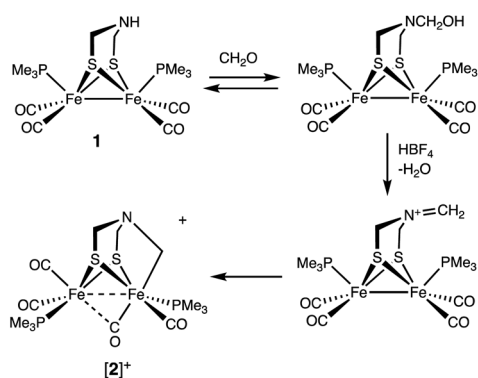
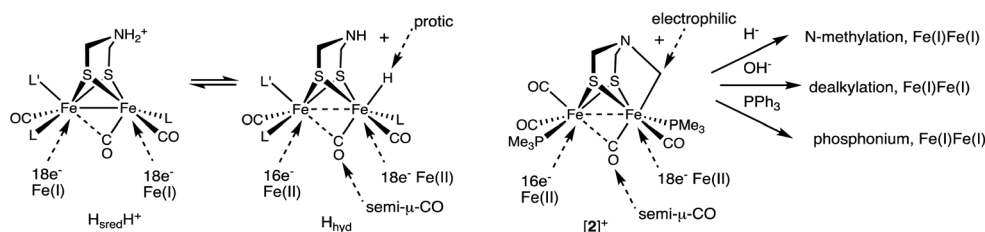


Fig. 6 Structure of  $[\text{Fe}_2(\mu\text{-SCH}_2)_2\text{NCH}_2\text{PPh}_3](\text{CO})_4(\text{PMe}_3)_2\text{BF}_4$  (**6**) with thermal ellipsoids shown at 50% probability. H atoms and  $\text{BF}_4^-$  anion have been omitted for clarity. Selected distances (Å): Fe1–Fe2, 2.5706(4); N1–C5, 1.436(3); N1–C6, 1.445(3); N1–C13, 1.449(3).



Scheme 5 Proposed pathway for the methylenation of **1**.

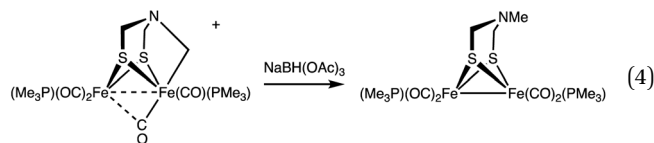
center is indicated by the  $^{31}\text{P}$  NMR singlet at  $\delta$  9.17, much higher field than  $\delta$  65.7 for  $\text{Fe}_2(\mu\text{-S}_2\text{C}_3\text{H}_6)(\text{CO})_5(\text{PPh}_3)$ .<sup>30</sup> In the region assigned to  $\text{NCH}_2\text{P}$ , the  $^1\text{H}$  NMR spectrum features a broad signal at  $\delta$  4.90. The broadness is associated with the nonequivalent protons, each of which is coupled to  $^{31}\text{P}$ . The IR spectrum of **6** also agrees with reduction of  $\text{Fe(II)Fe(II)}$  to  $\text{Fe(I)Fe(I)}$ :  $\nu_{\text{CO}}$  shifts to lower frequency by  $55\text{ cm}^{-1}$  (1978, 1948,  $1903\text{ cm}^{-1}$ ). These frequencies are comparable to those in **1**.



Scheme 6 Comparison of the  $[\text{2Fe}]_{\text{H}}$  centers in the  $\text{H}_{\text{sred}}\text{H}^+$  and  $\text{H}_{\text{hyd}}$  states ( $\text{L} = \text{CN}^-$ ) and  $[\text{2}]^+$ , including its reactions.

The structure of **6** was verified by X-ray crystallography (Fig. 6). The complex is a conventional  $\text{Fe}_2(\mu\text{-SR})_2(\text{CO})_{6-x}\text{L}_x$  butterfly. The bulky phosphonium substituent is distant from the  $\text{Fe}_2$  core.

Conversion of  $[\text{2}]^+$  back to **1** was induced upon treatment with  $\text{Et}_4\text{NOH}$ . From this reaction, **1** was recovered in 40% yield after purification by column chromatography. The electrophilic nature of the  $\text{Fe-CH}_2$  bond is also supported by the reaction of  $[\text{2}]^+$  with  $\text{BH}(\text{OAc})_3^-$ , a mild hydride donor. In this case,  $\text{Fe}_2[(\mu\text{-SCH}_2)_2\text{NMe}](\text{CO})_4(\text{PMe}_3)_2$  was obtained in good yield (eqn (4)).



## Conclusions

As a specific conclusion, this work provides a plausible model for the inhibition of  $[\text{FeFe}]$ -hydrogenases by formaldehyde. The methylene group donated by formaldehyde occupies both substrate binding sites, amine and the distal Fe. The methylation proceeds by addition of the aldehyde to the secondary amine followed by generation of the iminium cation, which oxidatively adds to one of the  $\text{Fe(I)}$  centers, oxidizing the diiron site by  $2e^-$  (Scheme 5). Many examples exist for the addition of iminium cations to low-valent metals.<sup>31</sup> The methylation reaction does not proceed from the diiron  $\mu$ -hydride. In accord with the results of Bachmeier *et al.*,<sup>13</sup> the methylation is selective for reduced state(s) of the diiron center, as required for oxidative addition.

Complex  $[\text{2}]^+$  represents a rare mimic of a terminal hydride for the  $[\text{FeFe}]$ -hydrogenases. Normally terminal hydrides of synthetic diiron dithiolates rapidly isomerize,<sup>20</sup> which precludes extensive characterization of this key intermediate.<sup>32</sup> In the Mulheim mechanism, the diferrous terminal hydride corresponds to  $\text{H}_{\text{hyd}}$  state, defined as  $[\text{4Fe-4S}]^+ - \text{Fe}_p(\text{II}) - \text{amine} - \text{Fe}_d(\text{II}) - \text{H}$ . In this state, the hydride is protic, being reversibly deprotonated by the amine cofactor. Consistent with this model, the alkyl ligand in  $[\text{2}]^+$  is electrophilic. Furthermore, analogous to the reversible deprotonation of  $\text{H}_{\text{hyd}}$ ,  $[\text{2}]^+$  reversibly dealkylates (see Scheme 2) to give a  $\text{Fe(I)Fe(I)}$  species as proposed for the  $\text{H}_{\text{sred}}\text{H}^+$  state. Complex  $[\text{2}]^+$  exists as an equilibrium mixture of three isomers. The finding that these isomers are separated by



less than  $\sim 1$  kcal mol $^{-1}$ , shows that stereochemistry of the other ligands on the diiron dithiolate has little influence on the Fe–alkyl bond (and by inference Fe–hydride bond). Also like H<sub>hyd</sub> state, [2]<sup>+</sup> has a highly unsymmetrical semi-bridging CO *trans* to R (= alkyl, hydride) on the distal Fe is persistent. In a future paper we plan to describe the redox chemistry of this electrophilic Fe(II)Fe(II) in our quest to further probe analogues of the elusive H<sub>hyd</sub>H<sup>+</sup> state.

As established by its reactions with a range of nucleophiles (Scheme 6), [2]<sup>+</sup> presents opportunities for appending the Fe<sub>2</sub>(μ-SR)<sub>2</sub> center to other scaffolds.

The susceptibility of [2]<sup>+</sup> to nucleophilic attack is reminiscent of Co(III)-alkyls as represented by vitamin B<sub>12</sub> and its derivatives and models.<sup>33,34</sup> Given the vast chemistry of B<sub>12</sub> mimics, it is possible that a wide range of diiron alkyl chemistry awaits discovery and development.

## Experimental

### Materials and methods

Reactions were conducted in stirred solutions or slurries under nitrogen at room temperature unless otherwise indicated. Sample work-up routinely included rinsing solids with Et<sub>2</sub>O or pentane and storage under vacuum to remove traces of solvent. All reactions and purifications were conducted using standard Schlenk techniques or in an MBraun glovebox under N<sub>2</sub>. Solvents were purified using solvent purification system equipped with alumina filtration column. CD<sub>2</sub>Cl<sub>2</sub> was degassed by freeze–pump–thaw cycles and dried using 4 Å molecular sieves. <sup>1</sup>H, <sup>31</sup>P{<sup>1</sup>H}, and <sup>13</sup>C NMR spectra were recorded on Varian 500, Varian 600, Bruker 500, or Bruker Ascend 600 MHz spectrometers. Chemical shifts (δ/ppm) are referenced to residual solvent peak (5.32 ppm for <sup>1</sup>H and 53.84 ppm for <sup>13</sup>C in CD<sub>2</sub>Cl<sub>2</sub>). Chemical shifts (δ/ppm) for <sup>31</sup>P{<sup>1</sup>H} NMR were calibrated using 85% H<sub>3</sub>PO<sub>4</sub> as an external reference (0 ppm). Solution IR spectra were recorded on a PerkinElmer Spectrum 100 FTIR spectrometer. Elemental analysis was performed utilizing an Exeter CE-440 elemental Analyzer. A Waters Micromass Quattro II spectrometer was used to acquire ESI-MS data. Crystallographic data were collected on a Bruker D8 Venture kappa diffractometer equipped with a Photon II CPAD detector. An Iμs Microfocus Mo source (λ = 0.71073 Å) coupled with a multi-layer mirror monochromator provided the incident beam. Literature procedures were followed for the synthesis of (Et<sub>4</sub>N)<sub>2</sub>[Fe<sub>2</sub>(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CN)<sub>2</sub>(CO)<sub>4</sub>,<sup>35</sup> Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>-NH](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>,<sup>25</sup> and Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NMe](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>.<sup>23,25</sup> Other chemicals were purchased from commercial sources and used without further purification.

[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BF<sub>4</sub> ([1H]BF<sub>4</sub>). To a solution of Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub> (**1**) (100 mg, 0.21 mmol) in 10 mL of CH<sub>2</sub>Cl<sub>2</sub> was added HBF<sub>4</sub>·Et<sub>2</sub>O (34 mg, 0.21 mmol). After 10 min, the solution was concentrated to  $\sim 1$  mL, and a red solid was precipitated upon addition of 10 mL of Et<sub>2</sub>O. Yield: 95 mg (81%). <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 6.36 (s, 2H, NH<sub>2</sub>), 3.49 (s, 4H, (SCH<sub>2</sub>)<sub>2</sub>N), 1.57 (d, 18H, P(CH<sub>3</sub>)<sub>3</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (202 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 26.24. IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 2000, 1963,

1923 cm $^{-1}$ . Anal. calcd for C<sub>12</sub>H<sub>25</sub>BF<sub>4</sub>Fe<sub>2</sub>NO<sub>4</sub>P<sub>2</sub>S<sub>2</sub>: C, 25.2; H, 4.41; N, 2.45. Found, C, 25.06; H, 4.05; N, 2.32.

[HFe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BAR<sup>F</sup><sub>4</sub> ([1H]BAR<sup>F</sup><sub>4</sub>). To a solution of [1H]BF<sub>4</sub> (40 mg, 0.070 mmol) in 4 mL of CH<sub>2</sub>Cl<sub>2</sub> was added solid NaBAR<sup>F</sup><sub>4</sub> (73 mg, 0.070 mmol). IR spectra showed full conversion to [1H]BAR<sup>F</sup><sub>4</sub> after 3 h. The reaction mixture was filtered through Celite and evaporated. The residue was recrystallized using CH<sub>2</sub>Cl<sub>2</sub>/pentane at  $-30$  °C to give a red crystalline solid. Yield: 85% (79.9 mg). <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 7.72 (s, 8H, ArH), 7.57 (s, 4H, ArH), 4.13 (s, 4H, SCH<sub>2</sub>), 2.32 (p, *J* = 8.8 Hz, 1H, NH), 1.55 (d, *J*<sub>PH</sub> = 10.2 Hz, 18H, P(CH<sub>3</sub>)<sub>3</sub>),  $-14.52$  (t, *J*<sub>PH</sub> = 21.5 Hz, 1H). <sup>31</sup>P{<sup>1</sup>H} NMR (202 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 21.81. IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 2033, 1992 cm $^{-1}$ . ESI-MS *m/z* calcd for [M<sup>+</sup>], 483.9. Found, 484.0. Anal. calcd for C<sub>44</sub>H<sub>35</sub>-BF<sub>24</sub>Fe<sub>2</sub>NO<sub>4</sub>P<sub>2</sub>S<sub>2</sub>: C, 39.25; H, 2.62; N, 1.04. Found, 39.23; H, 2.57; N, 1.27.

[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NCH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BF<sub>4</sub> ([2]BF<sub>4</sub>). A solution of Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub> (**1**) (200 mg, 0.41 mmol) in 20 mL of CH<sub>2</sub>Cl<sub>2</sub> was treated with CH<sub>2</sub>O (25 mg, 0.83 mmol). After stirring this mixture for 1 h, HBF<sub>4</sub>·Et<sub>2</sub>O (67 mg, 0.41 mmol) was added by syringe. The color of the reaction mixture immediately changed from red to dark brown. After a further 30 min, the mixture was analyzed by IR spectroscopy, which showed that the bands for **1** were replaced by new bands at higher energy. The solution was concentrated to 5 mL under vacuum. Addition of 40 mL of Et<sub>2</sub>O precipitated the black solid. Yield: 207 mg (86%). <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 5.51 (br, 1H, NCH<sub>2</sub>Fe), 5.14 (br, 1H, NCH<sub>2</sub>Fe), 3.95 (br, 2H, SCH<sub>2</sub>), 3.31 (br, 2H, SCH<sub>2</sub>), 1.69 (br, 18H, P(CH<sub>3</sub>)<sub>3</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (203 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 35.00 (br), 24.67, 24.28, 22.70 (br), 8.94 (br). IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 2044, 2017, 1990, 1942 cm $^{-1}$ . HR-MS (ESI) *m/z* calcd for [M<sup>+</sup>], 495.9321. Found, 495.9312. Elemental analysis was obtained on the BAR<sup>F</sup><sub>4</sub><sup>-</sup> salt (see next procedure).

[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NCH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BAR<sup>F</sup><sub>4</sub> ([2]BAR<sup>F</sup><sub>4</sub>). A 20 mL vial was loaded with [2]BF<sub>4</sub> (50 mg, 0.086 mmol), 1.0 equiv. of NaBAR<sup>F</sup><sub>4</sub> (88 mg, 0.086 mmol), followed by 5 mL of CH<sub>2</sub>Cl<sub>2</sub>. After stirring the mixture for 1 h, solvent was removed *in vacuo*. The residue was extracted into 2 mL of CH<sub>2</sub>Cl<sub>2</sub>. This extract was filtered through Celite and layered with hexane. Dark-brown microcrystals were obtained after 2 days. Yield: 110 mg (95%). <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 7.73 (m, 8H, ArH), 7.57 (s, 4H, ArH), 5.52 (m, 1H, NCH<sub>2</sub>Fe), 4.79 (m, 1H, NCH<sub>2</sub>Fe), 4.17 (dd, *J* = 11.5, 3.8 Hz, 1H, SCH<sub>2</sub>), 3.84 (dd, *J* = 11.8, 8.7 Hz, 1H, SCH<sub>2</sub>), 3.25 (dd, *J* = 11.8, 3.4 Hz, 2H, SCH<sub>2</sub>), 1.66 (d, *J*<sub>PH</sub> = 10.3 Hz, 9H, P(CH<sub>3</sub>)<sub>3</sub>), 1.53 (d, *J*<sub>PH</sub> = 10.4 Hz, 9H, P(CH<sub>3</sub>)<sub>3</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (243 MHz, CD<sub>2</sub>Cl<sub>2</sub>), three isomers were detected: δ 22.59 (d, *J*<sub>PP</sub> = 7.3 Hz), 9.45 (d, *J*<sub>PP</sub> = 7.4 Hz), *trans*-dibasal; 34.46 (d, *J*<sub>PP</sub> = 7.7 Hz), 24.10 (d, *J*<sub>PP</sub> = 7.6 Hz), *cis*-dibasal; 18.41 (s), 9.21 (s), apical-basal. <sup>13</sup>C NMR (126 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 212.37–210.12 (m, *t*-CO), 201.90 (d, *J*<sub>PC</sub> = 19.7 Hz, μ-CO), 162.18 (q, <sup>1</sup>*J*<sub>BC</sub> = 49.8 Hz), 135.22, 131.08–128.70 (qq, <sup>2</sup>*J*<sub>CF</sub> = 31.4 Hz, <sup>4</sup>*J*<sub>CF</sub> = 2.9 Hz), 125.03 (q, <sup>1</sup>*J*<sub>CF</sub> = 272.4 Hz), 118.55–116.81 (m), 75.77 (d, <sup>2</sup>*J*<sub>PC</sub> = 12.0 Hz, NCH<sub>2</sub>Fe), 58.45 (s, SCH<sub>2</sub>), 58.15 (d, <sup>3</sup>*J*<sub>PC</sub> = 5.5 Hz, SCH<sub>2</sub>), 18.74 (d, <sup>1</sup>*J*<sub>PC</sub> = 33.4 Hz, P(CH<sub>3</sub>)<sub>3</sub>), 16.69 (d, <sup>1</sup>*J*<sub>PC</sub> = 32.0 Hz, P(CH<sub>3</sub>)<sub>3</sub>). IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 2046, 2020, 1992, 1942 (μ-CO). Anal. calcd for C<sub>45</sub>H<sub>36</sub>BF<sub>24</sub>Fe<sub>2</sub>NO<sub>4</sub>P<sub>2</sub>S<sub>2</sub>: C, 39.76; H, 2.67; N, 1.03. Found, C,



39.33; H, 2.59; N, 1.09. Single crystals were grown by diffusion of hexane into a CH<sub>2</sub>Cl<sub>2</sub> solution.

**Reaction of [2]BF<sub>4</sub> with NaBH(OAc)<sub>3</sub>.** To a solution of [2]BF<sub>4</sub> (15 mg, 0.026 mmol) in 2 mL of MeCN was added NaBH(OAc)<sub>3</sub> (5.5 mg, 0.026 mmol, 1 equiv.). The reaction solution changed from dark brown to red immediately. After a further 10 min, solvent was removed, and the residue was extracted into pentane. Removing the solvent under vacuum gave Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NMe](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub> as a red solid. Yield: 86% (11 mg). The NMR spectrum of this product matches that of authentic Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NMe](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>.<sup>25</sup> <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 2.91 (s, 4H, SCH<sub>2</sub>), 2.10 (s, 3H, NCH<sub>3</sub>), 1.49 (d, *J*<sub>PH</sub> = 9.1 Hz, 18H, P(CH<sub>3</sub>)<sub>3</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (202 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 22.81. ESI-MS *m/z* calcd for [M + H<sup>+</sup>], 497.9. Found, 498.2. IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 1983, 1945, 1909, 1894 sh cm<sup>-1</sup>.

**[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NCH<sub>2</sub>PPh<sub>3</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BF<sub>4</sub> ([6]BF<sub>4</sub>).** A solution of PPh<sub>3</sub> (23 mg, 0.086 mmol) in 2 mL of CH<sub>2</sub>Cl<sub>2</sub> was added dropwise to a solution of [2]BF<sub>4</sub> (50 mg, 0.086 mmol) in 2 mL of CH<sub>2</sub>Cl<sub>2</sub>. The color changed from purple to red immediately. After stirring for 10 min, the solution was concentrated to 1 mL. The concentrate was layered with 10 mL of Et<sub>2</sub>O, and this biphasic mixture was stored at -30 °C. Red crystals appeared after 24 h. Yield: 66 mg (91%). <sup>1</sup>H NMR (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 8.49–7.29 (m, 5H, ArH), 4.90 (s, 2H, NCH<sub>2</sub>P), 3.45 (s, 4H, SCH<sub>2</sub>), 1.50 (d, *J*<sub>PH</sub> = 9.2 Hz, 18H, P(CH<sub>3</sub>)<sub>3</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (203 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 24.15, 9.18. IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 1978, 1948, 1903. ESI-MS: *m/z* calcd for [M<sup>+</sup> - CO], 730.4. Found, 730.0. Anal. calcd for C<sub>31</sub>H<sub>39</sub>BF<sub>4</sub>Fe<sub>2</sub>NO<sub>4</sub>P<sub>3</sub>S<sub>2</sub>: C, 44.05; H, 4.65; N, 1.66. Found, C, 43.63; H, 4.96; N, 1.80. Single crystals were grown by diffusion of Et<sub>2</sub>O into a CH<sub>2</sub>Cl<sub>2</sub> solution at -30 °C.

**Reaction of [Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NCH<sub>2</sub>](CO)<sub>4</sub>(PMe<sub>3</sub>)<sub>2</sub>]BF<sub>4</sub> ([2]BF<sub>4</sub>) with Et<sub>4</sub>NOH.** To a solution of [2]BF<sub>4</sub> (40 mg, 0.069 mmol) in 2 mL of THF was added Et<sub>4</sub>NOH (10.1 mg, 0.069 mmol, 1.0 equiv., 20% wt aqueous solution). After 2 h, the reaction mixture was evaporated to dryness. The residue was extracted into 1 mL of CH<sub>2</sub>Cl<sub>2</sub>. This extract was filtered through Celite to remove Et<sub>4</sub>NBF<sub>4</sub>. A concentrate of this filtrate was purified by column chromatography on silica gel eluting with Et<sub>2</sub>O/pentane. Yield of **1**: 13 mg (40%). Product as **1** was identified by FT-IR and <sup>1</sup>H NMR spectroscopy, as well as TLC.

**Et<sub>4</sub>N[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NCH<sub>2</sub>CN](CN)(CO)<sub>4</sub>(L)] ((Et<sub>4</sub>N[4] (L = CO) and Et<sub>4</sub>N[5] (L = PPh<sub>3</sub>))**

**Et<sub>4</sub>N[4].** A solution of (Et<sub>4</sub>N)<sub>2</sub>[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CN)<sub>2</sub>(CO)<sub>4</sub>] (50 mg, 0.078 mmol) in MeCN was treated with paraformaldehyde (4.7 mg, 0.016 mmol). After stirring this mixture for 2 h, a solution of H(OEt<sub>2</sub>)BF<sub>4</sub> (13 mg, 0.078 mmol) in 2 mL of MeCN was added dropwise. The color of the reaction mixture changed from deep red to dark brown immediately. FT-IR: ν<sub>CO</sub> = 2038, 2000, 1980, 1945, 1935 (sh), 1912 cm<sup>-1</sup>; ν<sub>CN</sub> = 2108 cm<sup>-1</sup>. ESI-MS: *m/z* calcd for [M<sup>-</sup>], 423.8. Found, 423.8. When the experiment was conducted in the presence of <sup>13</sup>CH<sub>2</sub>O, the FT-IR spectrum was the same. ESI-MS: *m/z* calcd for [M<sup>-</sup>], 424.8. Found, 424.8.

**Et<sub>4</sub>N[5].** The experiment above was repeated using [HPPPh<sub>3</sub>]BF<sub>4</sub> (27 mg, 0.078 mmol) in place of H(OEt<sub>2</sub>)BF<sub>4</sub>. A solution of (Et<sub>4</sub>N)<sub>2</sub>[Fe<sub>2</sub>[(μ-SCH<sub>2</sub>)<sub>2</sub>NH](CN)<sub>2</sub>(CO)<sub>4</sub>] (50 mg, 0.078 mmol) in MeCN was treated with paraformaldehyde (4.7 mg, 0.016

mmol). After 2 h, the IR spectrum showed no change in the CO region. A solution of [HPPPh<sub>3</sub>]BF<sub>4</sub> (27 mg, 0.078 mmol) in 2 mL of MeCN was then added dropwise. The color of the reaction mixture changed from red to dark brown immediately. After 12 h, the color turned to red again. The mixture was then concentrated to 2 mL, and the concentrate was filtered through Celite and layered with 20 mL of Et<sub>2</sub>O. After 2 days at -30 °C, the layered solution yielded a red solid. Yield: 75% (45 mg). <sup>1</sup>H NMR (600 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 7.68–7.66 (m, 6H, ArH) 7.40–7.39 (m, 9H, ArH), 3.17–3.15 (q, 8H, <sup>1</sup>N(CH<sub>2</sub>CH<sub>3</sub>)<sub>4</sub>), 2.57–2.52 (br, 4H, SCH<sub>2</sub>), 2.43 (s, 2H, NCH<sub>2</sub>CN), 1.25 (t, 12H, <sup>1</sup>N(CH<sub>2</sub>CH<sub>3</sub>)<sub>4</sub>). <sup>31</sup>P{<sup>1</sup>H} NMR (203 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 60.06. <sup>13</sup>C NMR (151 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ 218.28 (CO), 138.69 (d, CN), 133.71 (d, *J*<sub>PC</sub> = 11.5 Hz, P(C<sub>6</sub>H<sub>5</sub>)<sub>3</sub>), 129.67 (d, *J*<sub>PC</sub> = 2 Hz, P(C<sub>6</sub>H<sub>5</sub>)<sub>3</sub>), 128.44 (d, *J*<sub>PC</sub> = 9.0 Hz, P(C<sub>6</sub>H<sub>5</sub>)<sub>3</sub>), 114.82 (NCH<sub>2</sub>CN), 53.23 (<sup>1</sup>N(CH<sub>2</sub>CH<sub>3</sub>)<sub>4</sub>), 50.64 (SCH<sub>2</sub>), 46.77 (NCH<sub>2</sub>CN), 8.00 (<sup>1</sup>N(CH<sub>2</sub>CH<sub>3</sub>)<sub>4</sub>). IR (CH<sub>2</sub>Cl<sub>2</sub>): ν<sub>CO</sub> = 1988, 1950, 1916 cm<sup>-1</sup>, ν<sub>CN</sub> = 2081 cm<sup>-1</sup>. Anal. calcd for C<sub>35</sub>H<sub>41</sub>Fe<sub>2</sub>N<sub>4</sub>O<sub>4</sub>PS<sub>2</sub>·0.2CH<sub>2</sub>Cl<sub>2</sub>: C, 52.49; H, 5.18; N, 6.96. Found, C, 52.48; H, 5.42; N, 7.15. ESI-MS: *m/z* calcd for [M<sup>-</sup>], 657.9. Found, 657.9. Single crystals were grown by diffusion of Et<sub>2</sub>O into a CH<sub>2</sub>Cl<sub>2</sub> solution at room temperature.

## Data availability

All experimental and crystallographic data are available in the ESI.†

## Author contributions

Methodology, investigation, writing: F. Z.; conceptualisation and writing: T. B. R. Investigation: L. Z. and T. J. W. All authors have given approval to the final version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by GM-61153 from the National Institutes of Health.

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