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## Bioassembly of complex iron–sulfur enzymes: hydrogenases and nitrogenases

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### Abstract

Nature uses multinuclear metal clusters to catalyse a number of important multielectron redox reactions. Examples that employ complex Fe–S clusters in catalysis include the Fe–Mo cofactor (FeMoco) of nitrogenase and its V and all-Fe variants, and the [FeFe] and [NiFe] hydrogenases. This Perspective begins with a focus on the catalytic H-cluster of [FeFe] hydrogenase, which is highly active in producing molecular H<sub>2</sub>. There has been much recent progress in characterizing the enzyme-catalysed assembly of the H-cluster, including information gleaned from spectroscopy combined with in vitro isotopic labelling of this cluster using chemical synthesis. We then compare the lessons learned from H-cluster biosynthesis to what is known about the bioassembly of the binuclear active site of [NiFe] hydrogenase and the nitrogenase active site cluster FeMoco.

Several complex enzymes based on Fe–S clusters play crucial roles in biological redox processes and serve as aspirational paradigms for renewable energy catalysis<sup>1</sup>. Nitrogenases are expressed in a specific group of microorganisms called diazotrophs (of which *Azotobacter vinelandii* is the best studied organism) and carry out the reduction of atmospheric N<sub>2</sub> to NH<sub>3</sub>. In this way, the enzymes supply the vast majority of natural bioavailable nitrogen in the biosphere, which is supplemented by anthropogenic processes such as the Haber–Bosch process<sup>2</sup>. Hydrogenases catalyse the interconversion of H<sup>+</sup> and H<sub>2</sub>, a process crucial for metabolism of their host organisms. These highly active enzymes also serve as aspirational examples for H<sub>2</sub> production and utilization in renewable energy applications<sup>3,4</sup>. Hydrogenases are found in many microorganisms from archaea and bacteria to some eukaryotes and can be categorized into three subgroups based on the metal content of their catalytic sites: [FeFe] hydrogenases, [NiFe] hydrogenases and [Fe] hydrogenases. This Perspective covers the bioassembly of the active sites in [FeFe] and [NiFe] hydrogenases, which catalyse the redox interconversion  $2\text{H}^+ + 2\text{e}^- \rightleftharpoons \text{H}_2$ . Both active sites feature Fe atoms ligated by CO and CN<sup>-</sup> ligands and the question of how these toxic inorganic diatomics are safely formed and vectored into the assembly of complex Fe–S

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clusters is of great interest. [NiFe] hydrogenases in all organisms feature, aside from their active sites, three additional Fe–S clusters or connect to additional Fe–S-cluster-containing subunits that serve as an electron-transport chain<sup>3</sup>. By contrast, [FeFe] hydrogenase enzymes all feature a catalytic 6Fe ‘H-cluster’ but have varying arrangements of accessory Fe–S clusters. For example, [FeFe] hydrogenase from *Clostridium pasteurianum* (CpI) contains three [4Fe–4S] clusters and one [2Fe–2S] cluster in its electron-transport chain<sup>5</sup>. By contrast, [FeFe] hydrogenase from *Desulfovibrio desulfuricans* (DdH) uses two additional [4Fe–4S] clusters<sup>6</sup>, while the enzyme from green algae *Chlamydomonas reinhardtii* (CrHydA1) features the H-cluster as its only Fe–S centre<sup>7</sup>. Thus, the H-cluster in the CrHydA1 enzyme is the most easily studied because it can be probed without interferences from other Fe–S clusters. For other enzymes, electron paramagnetic resonance (EPR) spectra feature overlapping signals for the H-cluster and accessory clusters.

## The [FeFe] hydrogenase H-cluster

We begin this Perspective with a focus on the bioassembly of the catalytic 6Fe H-cluster of [FeFe] hydrogenase, which produces H<sub>2</sub> at a very high rate ( $\sim 10^4$  s<sup>-1</sup>)<sup>8</sup>. The H-cluster consists of a canonical [4Fe–4S]<sub>H</sub> cluster linked by a Cys thiolate to a unique binuclear [2Fe]<sub>H</sub> cluster (FIG. 1; the subscript H denotes part of the H-cluster). Each Fe in the organometallic [2Fe]<sub>H</sub> cluster has terminal CN<sup>-</sup> and CO ligands, with an additional CO bridging the two Fe centres<sup>5,6,9</sup>. Lastly, there is an additional bridging azadithiolate (adt<sup>2-</sup>, HN(CH<sub>2</sub>S<sup>-</sup>)<sub>2</sub>) ligand, of which the secondary amine is thought to relay H<sup>+</sup> to/from a coordination site on the Fe that is distal to the [4Fe–4S]<sub>H</sub> cluster<sup>10–13</sup>. In active states of the enzyme, this site can either be vacant or be occupied by H<sup>-</sup> or H<sub>2</sub> ligands. For example, the site is vacant in the oxidized H<sub>ox</sub> state and occupied by H<sup>-</sup> in the H<sub>hyd</sub> state. As we see, these two prominent EPR-active forms have spectroscopic signatures that allow us to rapidly compare native and semi-synthetic enzymes to deduce the enzyme’s structure and biosynthesis.

Nature uses a set of three Fe–S proteins — HydE, HydF and HydG — to construct the unusual [2Fe]<sub>H</sub> cluster and link it to the [4Fe–4S]<sub>H</sub> cluster, which is installed by the regular Fe–S cluster assembly machinery encoded by, for example, the *isc* operon<sup>7,14–16</sup>. In recent years, we have combined spectroscopic methods with in vitro maturation of the H-cluster, employing the purified apo-hydrogenase CrHydA1 (as expressed in *Escherichia coli* with only the [4Fe–4S]<sub>H</sub> subcluster), along with cell lysates enriched in the three maturase proteins as well as with a cocktail of low-molecular-weight cofactors and precursors, all mixed together in a test tube under anaerobic conditions<sup>17–19</sup>. With the in vitro-maturation approach, we can conveniently introduce isotopic labels into the maturation pathway by using labelled substrates and cofactors during enzymatic assembly of the H-cluster or, alternatively, in kinetically-resolved studies of individual maturation enzymes. This approach has proven powerful and has allowed us to identify the molecular sources of each atom in the [2Fe]<sub>H</sub> cluster and to isotopically label the H-cluster to enable further spectroscopic and mechanistic studies (FIG. 1a). Many of the intermediates in the maturation pathway are paramagnetic, such that magnetically-active nuclei near unpaired spin density are readily detected because they give rise to electron–nuclear hyperfine interactions. We have used both continuous-wave (CW) and pulse EPR techniques in

these studies, with pulse methods such as hyperfine sublevel correlation (HYSCORE) and electron–nuclear double resonance (ENDOR) allowing us to probe weaker hyperfine interactions than are evident in conventional CW EPR (see REFS<sup>20–22</sup> for the principles and applications of pulse EPR techniques). The information gleaned from the spectroscopy is then translated into chemical insights into the maturation pathway.

## Initial Fe–CO and Fe–CN bonding

Of the three H-cluster maturases, HydG has been the most studied and best understood enzyme. It is a member of the radical *S*-adenosylmethionine (rSAM) superfamily of enzymes<sup>23–25</sup> that homolytically cleave an *S*-adenosylmethionine (SAM) coenzyme bound to a [4Fe–4S] cluster. The resulting C-centred 5'-deoxyadenosyl radical<sup>26,27</sup> typically initiates transformation of a substrate through stereoselective and regioselective H-atom abstraction. The substrate of HydG is free tyrosine<sup>28</sup>, the amino group of which undergoes H-atom abstraction by the 5'-deoxyadenosyl radical<sup>29</sup> (FIG. 2a), leading to homolysis of the tyrosine C $\alpha$ –C $\beta$  bond and a transient EPR-active 4-hydroxybenzyl (4-OB $\cdot$ ) radical, along with dehydroglycine (DHG)<sup>30</sup>. The DHG further fragments to form CN $^-$  and CO (REFS<sup>31–34</sup>), which are observable in transient Fourier transform infrared (FTIR) spectra<sup>35</sup> (FIG. 2b). We note that, although toxic, at low levels, CO plays an important signalling role. In humans, this signalling provides protective stress and immune responses to injuries, preventing the development of high-fat-diet-induced obesity and regulating metabolic states in various cancer cells or tumour microenvironments<sup>36</sup>. However, the CO used in signalling is produced by the oxidative degradation of haem by haem oxygenase<sup>37</sup>, a very different process to the rSAM HydG generation of CO/CN $^-$  stoichiometrically through tyrosine radical fission.

In HydG, the potentially toxic CO and CN $^-$  fragments are immediately passivated by rapid binding to an Fe site in a unique 5Fe auxiliary Fe–S cluster, which, in the X-ray structure of HydG, is located at the other end of a 24 Å channel from the SAM cluster<sup>38,39</sup>. The CO and CN $^-$  molecules produced by the first tyrosine scission both bind a ‘dangler’ Fe (REF.<sup>40</sup>), which is additionally ligated by a conserved His residue and a cysteine cofactor. The thiolate in the latter non-peptidic ligand bridges this unique Fe to a [4Fe–4S] cluster<sup>41,42</sup> (FIG. 2a). Scission of a second tyrosine molecule again affords CO and CN $^-$ , but whereas the former binds the dangler Fe, the latter binds the [4Fe–4S] cluster to give an EPR-detectable [4Fe–4S]-CN species<sup>41</sup>. This binding releases the HydG product, an organometallic {Fe(CO)<sub>2</sub>(CN)(cysteinato)} synthon and precursor in the assembly of the [2Fe]<sub>H</sub> cluster (FIG. 2c). By introducing <sup>57</sup>Fe into HydG and using <sup>13</sup>C-labelled and <sup>15</sup>N-labelled tyrosine, one can use CW and pulse EPR to track the incorporation of these isotopes into the paramagnetic H<sub>ox</sub> and H<sub>hyd</sub> forms of the H-cluster (FIG. 1a) assembled in vitro<sup>35,43,44</sup>. This magnetic-isotope editing allows us to measure the specific nuclear hyperfine interactions between the <sup>13</sup>C, <sup>15</sup>N and <sup>57</sup>Fe centres and the unpaired electron spin density residing on the H-cluster in these two paramagnetic states. Moreover, use of <sup>13</sup>C-labelled and <sup>15</sup>N-labelled tyrosine as the substrate produces clear isotopic shifts in time-resolved FTIR bands assigned to the Fe-bound CO and CN $^-$  ligands<sup>35</sup>.

It is notable that cysteine is a component of the organometallic HydG product  $\{\text{Fe}(\text{CO})_2(\text{CN})(\text{cysteinato})\}$ . The  $[\text{4Fe-4S}]\text{-CN}$  co-product converts to  $[\text{4Fe-4S}]\text{-cysteinato}$ , of which the pendant amino acid and the aforementioned proximal His residue make up a binding site for a fifth Fe. The introduction of this Fe completes the catalytic cycle. Aside from the cysteinato's role in binding Fe, could it also be possible that this ligand is the source of the S atoms in the  $\text{adt}^{2-}$  cofactor? And, if so, does the rest of the  $\text{adt}^{2-}$  bridge also originate from cysteine? This has recently been addressed in a collaboration between our group and the Rauchfuss group. In this work, we performed in vitro maturation without HydG and tyrosine, instead using a synthetic  $\{\text{Fe}(\text{CO})_2(\text{CN})(\text{cysteinato})\}$  source, termed syn-B, to test whether the proposed final product of HydG can be effectively introduced as a synthetic analogue<sup>45</sup> (FIG. 3a). This approach is reminiscent of prior work in which the synthetic 2Fe complex  $[\text{Fe}_2(\text{CO})_4(\text{CN})_2(\text{adt})]^{2-}$  was linked to  $[\text{4Fe-4S}]\text{-containing}$  apo-hydrogenase to form fully active hydrogenase in the absence of maturase enzymes (or with only HydF in the first publication)<sup>9,46-48</sup>. Our distinct semi-synthesis approach takes this a step further back in the biosynthesis to the point where the mononuclear Fe complex is assembled. This approach affords a  $[\text{FeFe}]$  hydrogenase, whose  $\text{H}_{\text{ox}}$  EPR signal and high activity for  $\text{H}_2$  evolution are comparable to that of the enzyme matured in vitro with HydE, HydF and HydG present. All these measurements are, reassuringly, also similar to those for the native enzyme. If one uses a source of  $\{\text{Fe}(\text{CO})_2(^{13}\text{CN})(\text{cysteinato})\}$ , the final product gives rise to the same  $^{13}\text{C}$  pulse EPR spectrum as that obtained earlier using  $^{13}\text{C}$ -labelled tyrosine and HydG as part of the standard in vitro method. The precursor syn-B does not only offer possibilities for isotope substitution at the diatomic ligands but also at the cysteinato ligand. However,  $^{13}\text{C}$  and/or  $^{15}\text{N}$  labels in this ligand are not incorporated into the  $\text{adt}^{2-}$  bridge, with the expected hyperfine coupling being absent from pulse EPR signals. Parallel mass spectrometry experiments revealed the presence of  $^{13}\text{C}_3\text{-pyruvate}$  in the maturation mixture, indicating S- $^{13}\text{C}$  bond cleavage of the  $^{13}\text{C}_3,^{15}\text{N}\text{-cysteinato}$  ligand within labelled syn-B. One could, in principle, explore possible couplings to a magnetically active  $^{33}\text{S}$  nucleus introduced by using  $^{33}\text{S}\text{-cysteine}$ , but  $^{33}\text{S}$  is difficult to obtain (it is 0.7% abundant) and is an  $I = 3/2$  nucleus with a relatively low magnetic moment, along with quadrupolar broadening. An alternative approach is to incorporate selenocysteine into a synthetic carrier and to use extended X-ray absorption fine structure spectroscopy to characterize the structure of the resultant H-cluster (FIG. 3b). Here, we observe two Se-Fe interactions, measured at both the Se and Fe X-ray absorption edges, and the consensus Se-Fe distance of 2.43 Å is consistent with two Se atoms being incorporated into  $\text{adt}^{2-}$  at the locations usually occupied by S (FIG. 3). We, thus, concluded that each of the two chalcogen atoms in  $\text{adt}^{2-}$  are from cysteine/selenocysteine in the  $\{\text{Fe}(\text{CO})_2(\text{CN})(\text{cysteinato}/\text{selenocysteinato})\}$  carrier, with the rest of cysteine being clipped off and converted into pyruvate ( $^{13}\text{C}_3\text{-pyruvate}$  when using  $^{13}\text{C}_3,^{15}\text{N}\text{-cysteine}$ ). This demonstrates that the S-Fe bond, first formed when cysteine binds the dangler Fe to give the resting state of the HydG, remains intact throughout the assembly all the way to the mature H-cluster<sup>45</sup>. But these results leave open another question: what are the molecular sources of C and N components of the  $\text{adt}^{2-}$  bridge?

## The CNC moiety of the azadithiolate

Given the conclusion that only the S atom of cysteine ends up incorporated into the  $\text{adt}^{2-}$  bridge, we set out to find the source of the CNC atoms in the centre of the bridge. A distinct possibility is one or more amino acids, so we first added a fully  $^{13}\text{C}$ ,  $^{15}\text{N}$ -labelled algal amino acid mixture to the maturation media and observed a set of  $^{13}\text{C}$  and  $^{15}\text{N}$  peaks in the HYSCORE data of the  $\text{H}_{\text{ox}}$  state<sup>49</sup>. This motivated a systematic survey of subsets of labelled amino acids with their possible incorporation assayed by HYSCORE, which told us that labelled serine affords an enzyme with a fully  $^{15}\text{N}$ -labelled and  $^{13}\text{C}$ -labelled  $\text{adt}^{2-}$  bridge. Using  $^{13}\text{C}_3, ^{15}\text{N}$ -serine affords an enzyme whose  $\text{H}_{\text{ox}}$  state gives rise to an ENDOR spectrum with a well-defined pair of  $^{15}\text{N}$  transitions and two distinct  $^{13}\text{C}$  couplings, consistent with a central  $^{15}\text{N}$  and two magnetically inequivalent  $^{13}\text{C}$  sites originating from the labelled amino acid (FIG. 4a). Moreover, these  $^{15}\text{N}$  and  $^{13}\text{C}$  couplings closely match those observed from semisynthesis of the enzyme using synthetic  $[\text{Fe}_2(\text{CO})_4(\text{CN})_2(\text{adt})]^{2-}$  featuring a  $^{13}\text{C}_2, ^{15}\text{N}$ - $\text{adt}^{2-}$  bridge<sup>50,51</sup>. Furthermore, by separately using the three singly  $^{13}\text{C}$ -labelled serine isotopologues and comparing  $^{13}\text{C}$  ENDOR data of the resulting product to that of the  $^{13}\text{C}_3, ^{15}\text{N}$ -serine sample, we showed that the C3 atom of serine is the source of the two C atoms of the  $\text{adt}^{2-}$  bridge (FIG. 4b). We also observed a  $^2\text{H}$  ENDOR signal in another  $\text{H}_{\text{ox}}$  sample prepared with  $\text{C}3\text{-}^2\text{H}_2$  serine in the maturation media. Given that only C3 of serine is utilized, it is clear that two such serine molecules are needed for the synthesis of each  $\text{adt}^{2-}$  bridge<sup>49</sup>.

## The radical SAM enzyme HydE

We have, so far, built up a detailed picture of the reaction mechanism and final product of HydG, yet, the other rSAM enzyme, HydE, has remained relatively uncharacterized. It has been suggested that HydE is involved in building the  $\text{adt}^{2-}$  ligand, and SAM turnover studies have been interpreted to support a thiol-containing compound as its substrate<sup>52</sup>. More recently, other groups have shown that HydE reacts with 1,3-thiazolidine to form a new  $\text{C}5'\text{-S}$  bond<sup>53</sup>. Although the relevance of this reaction to H-cluster synthesis seemed unclear, our recent work using the synthetic HydG product surrogate syn-B reveals a parallel HydE reaction that is on-pathway, where HydE generates the 5'-deoxyadenosyl radical that specifically attacks the cysteine S of the organometallic synthon formed by HydG, which we described putatively as  $\{\text{Fe}^{\text{II}}(\text{CO})_2(\text{CN})(\text{cysteinato})\}$ . The  $\text{C}5'\text{-S}$  bond forms concomitant with reduction of the low-spin  $\text{Fe}^{\text{II}}$  to  $\text{Fe}^{\text{I}}$  (REF.<sup>54</sup>), which, in turn, leads to cleavage of the internal cysteine  $\text{C}3\text{-S}$  bond, showing that HydE is responsible for the specific cysteine cleavage revealed earlier by maturation with the HydG-free system augmented with syn-B<sup>45</sup>. HydE's production of a mononuclear  $\{\text{Fe}^{\text{I}}(\text{CO})_2(\text{CN})\text{S}\}$  species is proposed to enable the subsequent formation of a dinuclear complex  $\{\text{Fe}_2^{\text{I}}(\text{CO})_4(\text{CN})_2(\mu\text{-S})_2\}$ , which then only needs the addition of the serine-derived CNC moiety to complete the  $[2\text{Fe}]_{\text{H}}$  subcluster. A remaining question is: does the final maturase HydF have a role in this last step of  $[2\text{Fe}]_{\text{H}}$  subcluster synthesis and/or its final linkage to  $[4\text{Fe-}4\text{S}]_{\text{H}}$  subcluster to complete the active H-cluster? We conclude our discussion of  $[\text{FeFe}]$  hydrogenase with this open question and now describe what is known about the assembly of  $[\text{NiFe}]$  hydrogenase enzymes.

## The [NiFe] hydrogenase cluster

The [NiFe] class of hydrogenase harbours a heterobimetallic  $\{(Cys)_2Ni(\mu-Cys)_2Fe(CO)(CN)_2\}$  active site that, relative to [FeFe] hydrogenase, is more tightly anchored to its polypeptide backbone. Instead of featuring just one Cys ligand, [NiFe] hydrogenases have four: two Cys thiolates bind Ni terminally (one of these is replaced by selenocysteinate in the related enzyme [NiFeSe] hydrogenase), while another two bridge it to Fe. Thus, the elaborate machinery responsible for the installation of the non-peptidic dithiolate  $adt^{2-}$  in [FeFe] hydrogenase is not required here. Nevertheless, both enzymes feature CO and  $CN^-$  ligands bound to a low-spin  $Fe^{II}$  centre<sup>4</sup>. It is, therefore, of interest to compare the bioassembly of these two clusters, not least the way in which the diatomic ligands are synthesized. The present model of [NiFe] binuclear cluster bioassembly involves a few sequential steps, including the biogenesis of the CO and  $CN^-$  ligands, formation of the  $\{Fe(CO)(CN)_2\}$  synthon and the insertion of Ni (REFS<sup>4,55</sup>) (FIG. 5a). In *E. coli*, the necessary machinery is encoded by the *hyp* operon (HypABCDE and HypF). Although the  $\{Fe(CO)(CN)_2\}$  unit in [NiFe] hydrogenases is similar to the  $\{Fe(CO)_2(CN)\}$  unit in the [FeFe] congener, little similarity can be found in their bioassembly pathways. The  $CN^-$  ligand in [NiFe] hydrogenase is derived from carbamoyl phosphate (an intermediate metabolite in the urea cycle) through HypEF-complex-catalysed *S*-carbamoylation and subsequent ATP-dependent dehydration<sup>56</sup> (FIG. 5b). The biogenesis of the CO ligand has long been elusive but has recently been unravelled. However, the new mechanism is only for the aerobic bioassembly of [NiFe] hydrogenase; the source of CO in anaerobic organisms still remains enigmatic. The central  $C_1$  pool is the aerobic source of the CO ligand<sup>57</sup> and the purified protein HypX, an additional maturase required in the aerobic assembly, acts on  $N^{10}$ -formyltetrahydrofolate to release CO, which is proposed to occur through a formyl-coenzyme A intermediate<sup>58</sup> (FIG. 5c). Thus, the mechanism of CO/ $CN^-$  synthesis is dramatically different from that used in the assembly of the [FeFe] hydrogenase H-cluster. Notably, the chemistry leading to both CO and  $CN^-$  ligands is mediated by enzymes that do not contain any metal cofactors, in sharp contrast to the case of HydG-catalysed tyrosine cleavage, in which the nascent CO and  $CN^-$  ligands are trapped by a labile Fe site inside this Fe-S protein to directly generate the organometallic synthon. This requires a different shuffle mechanism of these toxic ligands in the assembly of the [NiFe] centre. Indeed, HypE serves as the  $CN^-$  carrier by converting a cysteine thiol into a thiocyanate group. This process occurs twice and, in each case, the  $CN^-$  ligand is transferred from the HypE-SCN adduct to an Fe centre coordinated by the HypCD protein complex<sup>59,60</sup>. The CO ligand is proposed to be attached to the same Fe, although the delivery of CO to finish the  $\{Fe(CO)(CN)_2\}$  synthon is not fully understood. In any case, the  $\{Fe(CO)(CN)_2\}$  synthon is then transferred into the apo-form of the [NiFe] hydrogenase large subunit, which triggers the selective insertion of Ni into the active site, as facilitated by metallochaperone proteins HypAB and SlyD<sup>61,62</sup>. On completion, the holo-protein undergoes proteolysis, conformational changes and translocation to form the fully functional [NiFe] hydrogenase.

## The nitrogenase FeMoco cluster

The nitrogenases are another important class of Fe-S enzyme, catalysing the crucial  $N_2$  reduction reaction:  $N_2 + 8e^- + 16ATP + 8H^+ \rightarrow 2NH_3 + H_2 + 16ADP + 16P_i$  (REF.<sup>63</sup>). As

with the hydrogenases, nitrogenases evolve  $H_2$ , albeit as part of a larger overall reduction reaction that consumes ATP to afford  $NH_3$ . The Mo nitrogenase enzyme NifDK has two types of Fe–S cluster, the 8Fe–7S P-cluster and the 8Fe–C–9S–Mo M-cluster that serves as the catalytic centre; alternative forms have V (VnfDGK) or Fe (AnfDGK) atoms instead of Mo (REFS<sup>64,65</sup>). There is also a specific protein bearing a 4Fe–4S cluster that serves as the obligatory electron source for the  $N_2$  reduction reaction. There has been much recent progress in characterizing the biosynthesis of the M-cluster, which relies on the chemistry of a key radical SAM enzyme NifB (REFS<sup>66–68</sup>) (FIG. 6). In addition to the rSAM [4Fe–4S] cluster, NifB houses two auxiliary 4Fe–4S clusters, K1 and K2, that are transformed into an 8Fe precursor to the M-cluster. The SAM cofactor plays two distinct roles here. First, there is a SAM-sourced transfer of a methyl group to one of the 4Fe–4S clusters. Second, rSAM chemistry initiated at the NifB SAM cluster gives a 5'-deoxyadenosyl radical from a second SAM, which then abstracts a H atom from a methyl group, ultimately forming the 8Fe–8S–C 'L\*-cluster', wherein the  $\mu^6$ -coordinated interstitial carbide is from the methyl group<sup>69–71</sup>. Insertion of an additional (ninth) S atom completes the central belt<sup>72</sup>, forming the 8Fe–9S–C 'L-cluster', which is then transferred from NifB to the scaffold protein NifEN (or VnfEN), where Mo (or V) and homocitrate are inserted by the NifH protein to form the final M-cluster<sup>67,73</sup>. All that remains is to transfer the M-cluster to the specific catalytic nitrogenase protein NifDK, VnfDGK or AnfDGK.

## Closing summary

Although the [FeFe] and [NiFe] hydrogenases catalyse the same reaction and share some organometallic structural elements, their assembly processes are quite different. Specifically, the formation and handling of the toxic CO and  $CN^-$  species are very different in the two biosyntheses. The construction of [FeFe] hydrogenase sees CO and  $CN^-$  formed from amino acids through an rSAM reaction in HydG, with these diatomics then being stabilized by binding to a specific Fe site at a second Fe–S cluster of the same enzyme to give a  $\{Fe(CO)_2(CN)(cysteinato)\}$  synthon. In [NiFe] hydrogenase, a comparable  $\{Fe(CO)(CN)_2\}$  synthon is formed from the interplay of multiple protein complexes that are essentially metal-free. It is, therefore, interesting to see how nature has evolved two different solutions to one problem. Circumstances often lead to nature forming the same product in different ways, as we highlighted earlier in the case of biosynthesis of signalling CO through haem oxidation.

In comparing the complexity of the biosynthesis of the two redox-active hydrogenases, the reader will note the many steps required to prepare the  $adt^{2-}$  cofactor, a ligand unique to the [FeFe] class. The biosynthesis of this cofactor, which is unstable in metal-free form, is challenging and may require additional enzymes in a process that is not yet fully understood. In the [NiFe] enzymes, the  $H^+$  relay role is, instead, played by a proteinaceous Arg residue<sup>74</sup> (although the terminally bound Cys thiolates have also been suggested). The nitrogenase assembly shares the crucial need for rSAM enzymes with the [FeFe] hydrogenase H-cluster assembly, but, in this case, the rSAM enzyme NifB carries out distinctive chemistry, using SAM both as a methyl donor and as a radical coenzyme to cement two [4Fe–4S] clusters together with an SAM-methyl-derived bridging carbide. Thus, the organometallic



intermediate formed by NifB is very different to the Fe(CO)<sub>2</sub>(CN)-containing intermediate formed by HydG, although both enzymes harness the special power of rSAM chemistry.

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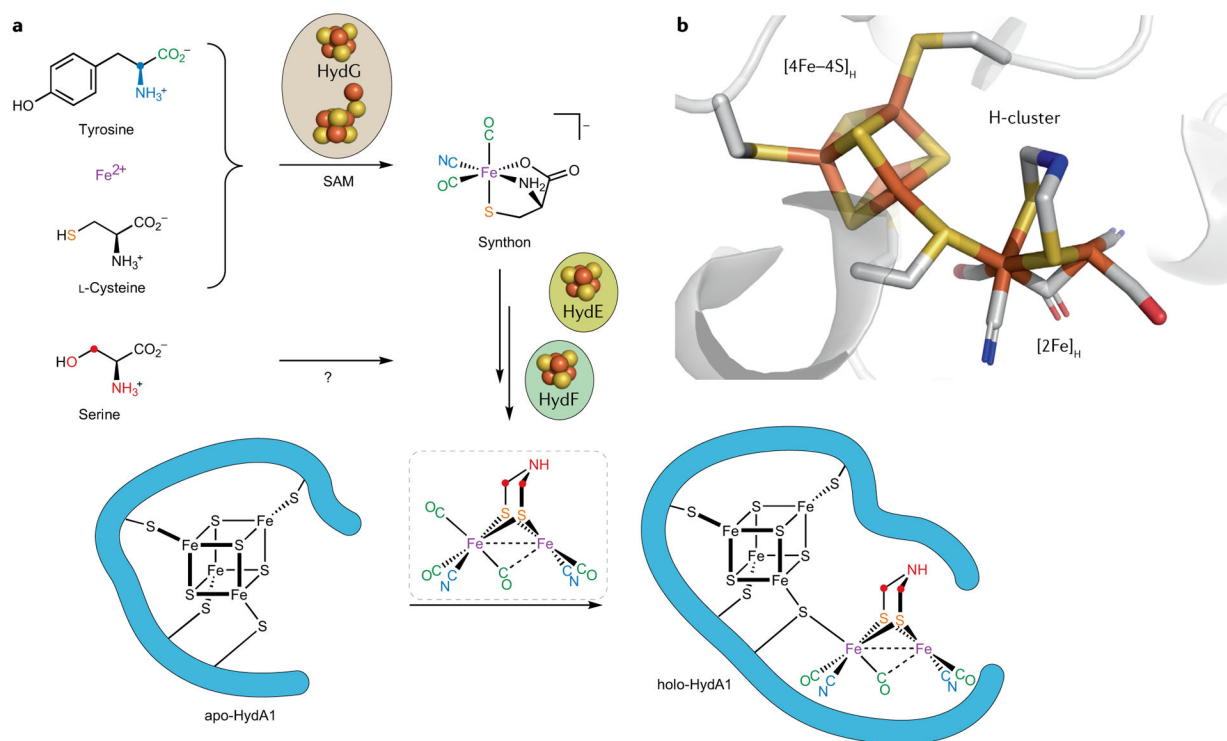
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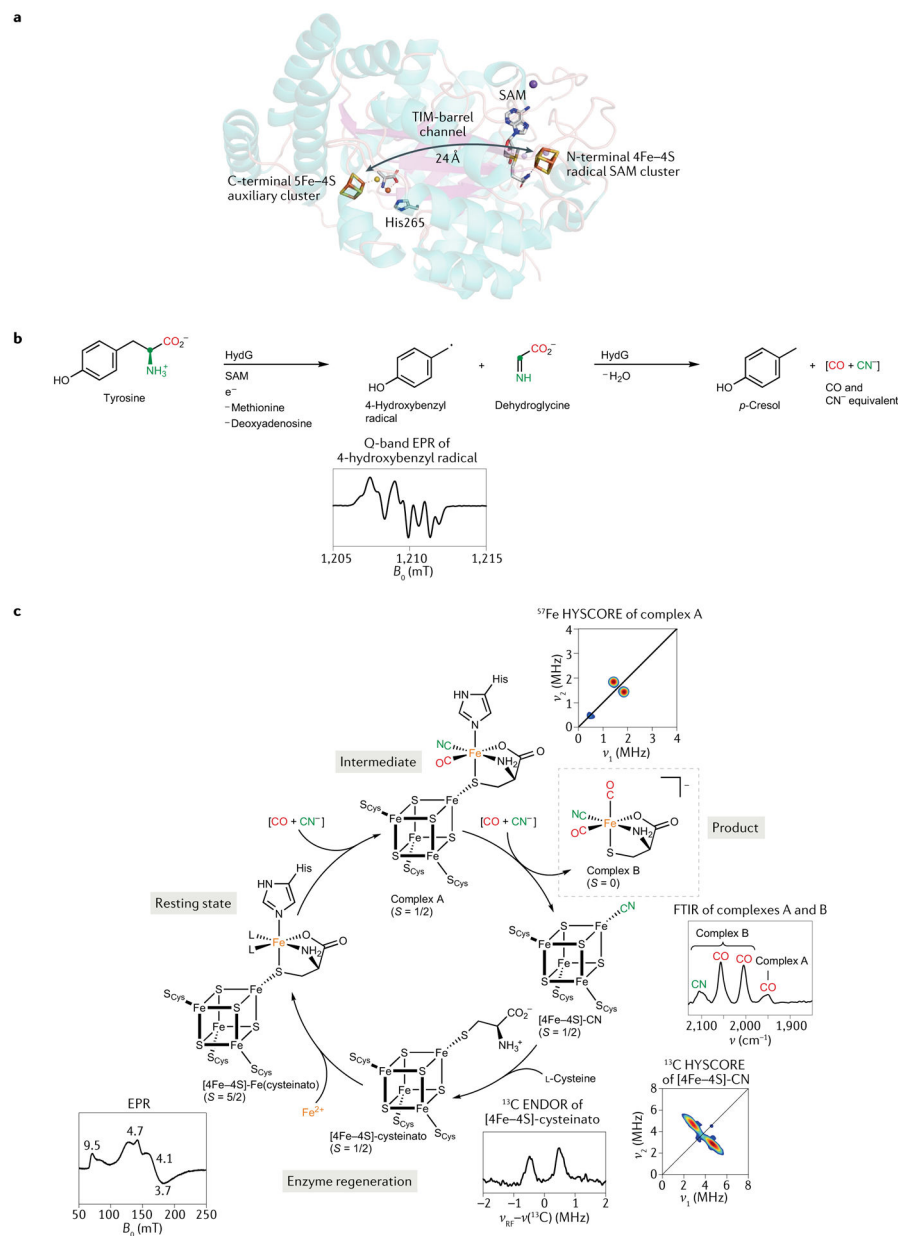
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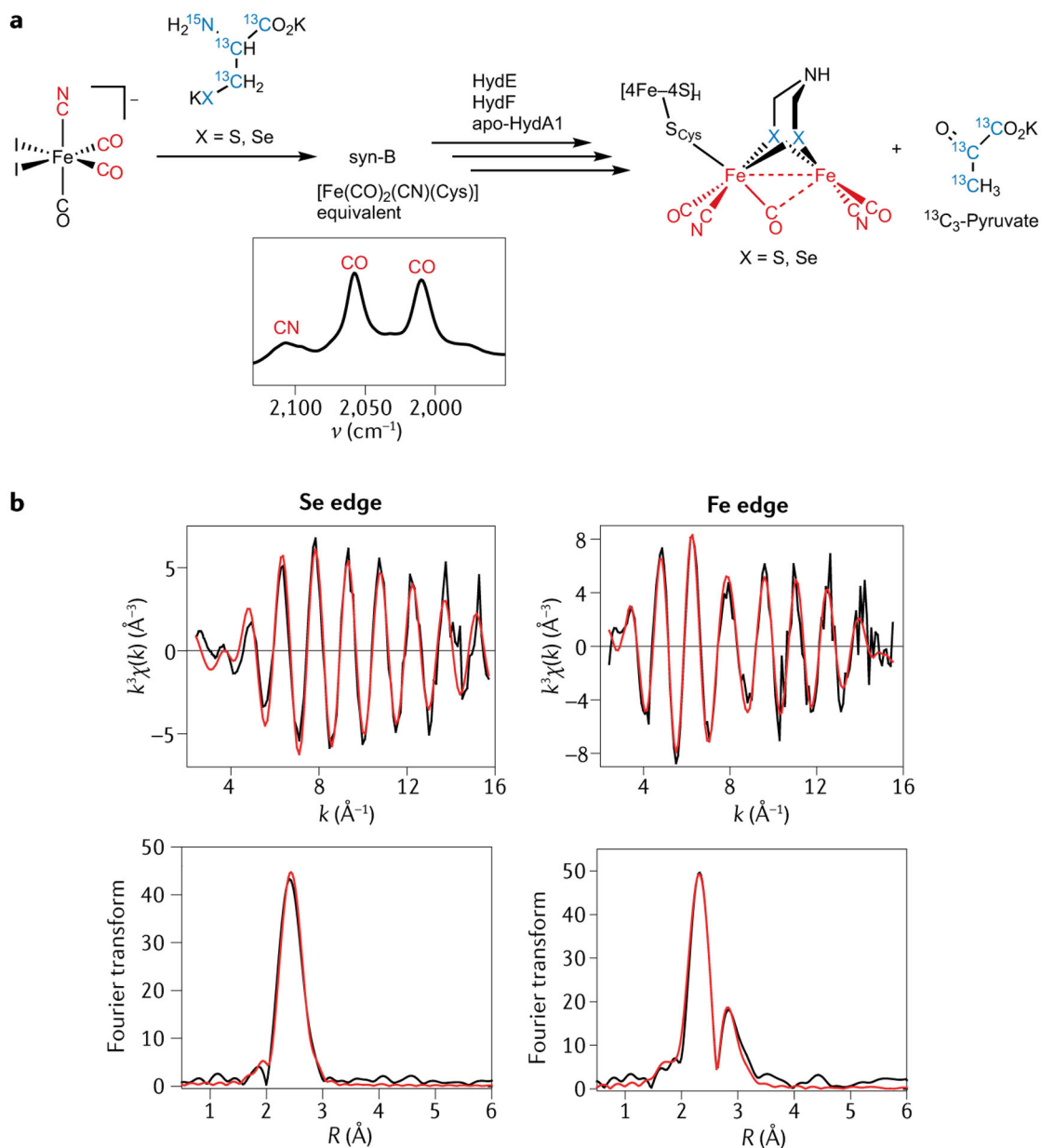
**Fig. 1 | The [FeFe] hydrogenase H-cluster and its biosynthetic origins.**

**a** | The present model of the [FeFe] hydrogenase H-cluster bioassembly accounts for the molecular origins of all atoms in the  $[2\text{Fe}]_{\text{H}}$  subcluster. **b** | The X-ray structure of the H-cluster in *Clostridium pasteurianum* [FeFe] hydrogenase (CpI) (Protein Data Bank identifier: 4XDC)<sup>9</sup>. SAM, *S*-adenosylmethionine.



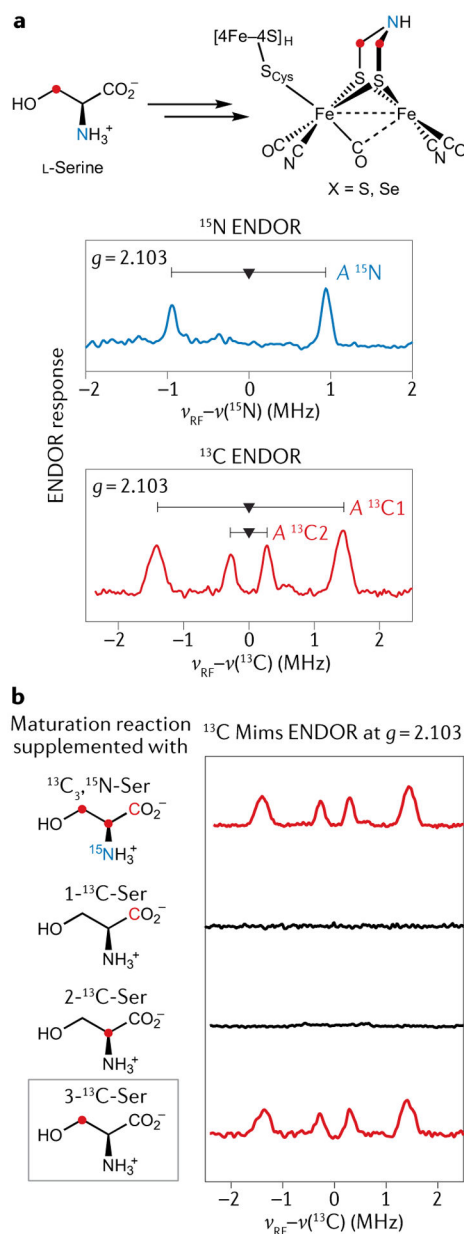
**Fig. 2 | Reactions carried out by the radical SAM maturase HydG.**

**a** | The X-ray structure of HydG, which features two Fe–S clusters (Protein Data Bank identifier: 4WCX)<sup>38</sup>. **b** | Tyrosine is subjected to radical reactions that see it fragment into the electron paramagnetic resonance (EPR)-active 4-hydroxybenzyl radical (4-OB•) and dehydroglycine (DHG). These further convert to *p*-cresol, CO and CN<sup>−</sup>. **c** | The organometallic synthon forms at the auxiliary 5Fe–4S cluster, as evidenced by FTIR spectroscopy. ENDOR, electron–nuclear double resonance; FTIR, Fourier transform infrared; HYSCORE, hyperfine sublevel correlation; SAM, *S*-adenosylmethionine. Part **b** adapted with permission from REF.<sup>30</sup>, AAAS. Part **c** adapted with permission from REF.<sup>35</sup>, AAAS.



**Fig. 3 |. Transfer of S/Se from the HydG synthon to make the  $adt^{2-}$  bridge.**

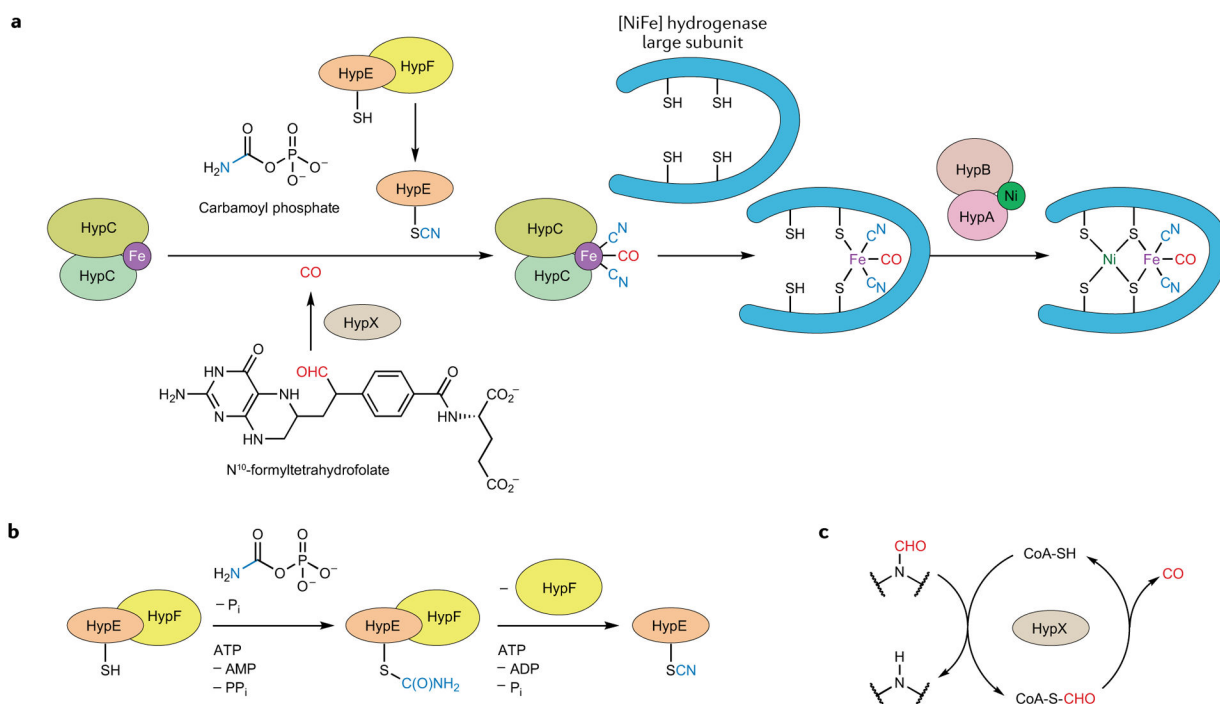
**a** | The semi-synthetic maturation of the H-cluster with the bridging S/Se sourced from cysteine/selenocysteine introduced by a synthetic HydG synthon carrier. **b** | Se and Fe edge extended X-ray absorption fine structure data show the incorporation of Se into the  $[2Fe]_H$  subcluster. Adapted with permission from REF.<sup>45</sup>, PNAS.



**Fig. 4 |. The CNC component of the azadithiolate is sourced from serine.**

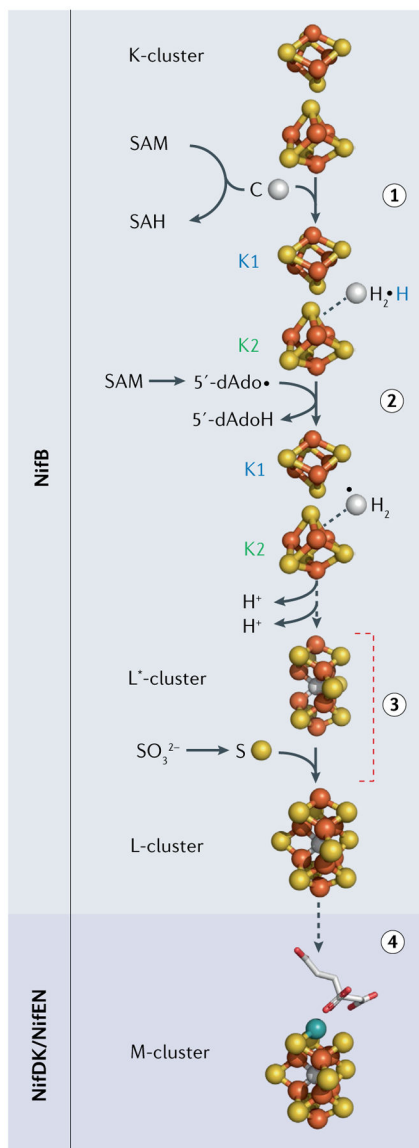
**a** | <sup>15</sup>N and <sup>13</sup>C electron–nuclear double resonance (ENDOR) spectra arising from the H<sub>ox</sub> H-cluster electron paramagnetic resonance signal of a HydA1 [FeFe] hydrogenase sample produced by in vitro maturation with added <sup>13</sup>C<sub>3</sub>, <sup>15</sup>N-labelled serine. **b** | <sup>13</sup>C ENDOR spectra of the H<sub>ox</sub> state of HydA1 [FeFe] hydrogenase samples produced via in vitro maturation with added <sup>13</sup>C<sub>3</sub>, <sup>15</sup>N-labelled, 1-<sup>13</sup>C-labelled, 2-<sup>13</sup>C-labelled and 3-<sup>13</sup>C-labelled serine, respectively, showing that the C atoms of the azadithiolate (adt<sup>2-</sup>) bridge originate from the 3-C of serine. Part **b** adapted with permission from REF.<sup>49</sup>, Royal Society of Chemistry.





**Fig. 5 | Aerobic bioassembly of the [NiFe] hydrogenase  $(\text{Cys})_2\text{Ni}(\mu\text{-Cys})_2\text{Fe}(\text{CO})(\text{CN})_2$  active site.**

**a** | Fe acquires its CO and  $\text{CN}^-$  ligands from organic sources, before the  $\{\text{Fe}(\text{CO})(\text{CN})_2\}$  fragment is delivered to the hydrogenase large subunit. The four Cys ligands then also bind Ni (REF.<sup>55</sup>). **b** | Biosynthesis of the  $\text{CN}^-$  ligand by the HypEF protein complex<sup>56</sup>. **c** | The HypX protein helps convert a formamide into the CO ligand via a thioformate intermediate<sup>58</sup>.



**Fig. 6 | Bioassembly of the nitrogenase M-cluster.**

The key initial step occurs at the radical *S*-adenosylmethionine (SAM) enzyme NifB, where two 4Fe–4S clusters are stitched together via an SAM methyl insertion, followed by a radical SAM H-atom abstraction to make the 8Fe–8S L\*-cluster (steps 1 and 2). Addition of a ninth S atom makes the final L-cluster (step 3), the target of the final specific metal substitution (V or Mo) capped by homocitrate (step 4). SAH, *S*-adenosylhomocysteine. Adapted from REF.<sup>75</sup>, CC BY 4.0.