


 Cite this: *Chem. Commun.*, 2024, 60, 10914

 Received 25th June 2024,  
Accepted 4th September 2024

DOI: 10.1039/d4cc03098b

rsc.li/chemcomm

## Secondary structure changes as the potential H<sub>2</sub> sensing mechanism of group D [FeFe]-hydrogenases†

 Ivan Voloshyn,<sup>a</sup> Conrad Schumann,<sup>b</sup> Princess R. Cabotaje,<sup>b</sup> Afridi Zamader,<sup>‡</sup> Henrik Land<sup>b</sup> and Moritz Senger<sup>\*,ab</sup>

**[FeFe]-hydrogenases function as both H<sub>2</sub> catalysts and sensors. While catalysis is well investigated, details regarding the H<sub>2</sub> sensing mechanism are limited. Here, we relate protein structure changes to H<sub>2</sub> sensing, similar to light-driven bio-sensors. Our results highlight how identical cofactors incorporated in alternative protein scaffolds serve different functions in nature.**

In nature, similar enzymes can serve multiple functions. A prime example of this versatility are the metalloenzyme [FeFe]-hydrogenases, which are mainly known for their high catalytic hydrogen (H<sub>2</sub>) turnover rates, making them an attractive target in the field of sustainable fuels research.<sup>1</sup> Besides catalysis, certain [FeFe]-hydrogenases are proposed to have a H<sub>2</sub> sensory function with the potential to regulate cellular metabolism.<sup>2–4</sup> The involvement of [FeFe]-hydrogenases in the H<sub>2</sub> metabolism of microbes shines new light on their role in medicine and health research.<sup>5,6</sup> [FeFe]-hydrogenases share the same cofactor, known as the H-cluster and are composed of a diiron site linked to a [4Fe–4S] cluster *via* the sulphur atom of a cysteine residue. The two irons of the diiron site are ligated by a carbon monoxide (CO) and cyanide (CN<sup>−</sup>) ligand each, and share a bridging CO molecule and an azadithiolate (ADT) bridge (Fig. 1A). [FeFe]-hydrogenases can be grouped into different phylogenetic groups (group A–G).<sup>7</sup> These groups exhibit variations in the second coordination sphere of the cofactor and overall protein architecture.<sup>1</sup> The most extensively studied

group A [FeFe]-hydrogenases are highly active, while groups C and D are proposed to serve a sensory function.<sup>1,8</sup> High turnover rates in group A have been associated with a sulphur-rich second coordination sphere of the diiron site and an alternative configuration of their unique proton transfer pathway (PTP) upon reduction.<sup>9,10</sup> In line with observations for the sensory [FeFe]-hydrogenases of group C, the second coordination sphere of the H-cluster in group D representative [FeFe]-hydrogenase from *Thermoanaerobacter mathranii*, *TamHydS*, lacks these sulphur-rich amino acids characteristic of group A [FeFe]-hydrogenases.<sup>3,4,9,11</sup> The low turnover rates of characterized representatives from groups C and D and the relative stability of the reduced potential signalling state, H<sub>red</sub>, upon H<sub>2</sub> exposure in *TamHydS* further implies a role in signalling rather than catalysis.<sup>3,12</sup> Moreover, we recently identified a novel PTP in group D of which the operational mechanism remains to be elucidated. (Fig. 1A).<sup>9</sup> Notably, the outer coordination sphere for *TamHydS* features a C-terminal extension which harbours an additional [4Fe–4S] cluster motif (Fig. 1B). The group C sensory [FeFe]-hydrogenase from *Thermotoga maritima*, *TamHydS*, features the same C-terminal FeS-cluster domain that is followed by a Per-Arnt-Sim (PAS) domain, which in combination with a Ser/Thr protein phosphatase might regulate downstream group A catalytic [FeFe]-hydrogenases.<sup>2,4</sup> However, in group D, the absence of this PAS domain necessitates an alternative sensing cascade, leading to their classification as putatively sensory [FeFe]-hydrogenases. The mechanism by which H<sub>2</sub> sensing is facilitated in group D enzymes remains an open question.

Here, we investigate the putative sensory function of group D [FeFe]-hydrogenase, *TamHydS*. Our genomic analysis supports a putative signalling function, particularly when compared to sensory group C enzymes. We demonstrate the enrichment of the potential signalling state, H<sub>red</sub>, in two ways, *via* photoreduction or exposure to H<sub>2</sub>. In contrast to the catalytic group A [FeFe]-hydrogenases, we do not observe a rearrangement of the H-bonding network of the PTP when H<sub>red</sub> is populated. Instead, we detect a secondary structural

<sup>a</sup> Department of Chemistry – BMC, Biochemistry, Uppsala University, 75120 Uppsala, Sweden. E-mail: moritz.senger@kemi.uu.se

<sup>b</sup> Department of Chemistry – Ångström Laboratory, Molecular Biomimetics, Uppsala University, 75120 Uppsala, Sweden

† Electronic supplementary information (ESI) available: Experimental procedures, infrared band patterns of different redox states, FTIR difference spectra of the H<sub>ox</sub> to H<sub>red</sub> transition and band kinetics, H/D exchange data, fit parameters, annotated genome region of *T. mathranii*. See DOI: <https://doi.org/10.1039/d4cc03098b>

‡ Current address: Laboratoire d'Electrochimie Moléculaire (LEM), Université Paris Cité, CNRS, F-75006, Paris, France.





**Fig. 1** H-cluster, PTPs and AlphaFold model of *TamHydS*. (A) The H-cluster and its connection to PTPs in group A and D [FeFe]-hydrogenases. Hydrogen bonding networks constitute the PTPs that are composed of amino acids and water molecules (red spheres) conserved within either group A (grey sticks) or group D<sup>9</sup> (green sticks). (B) The YASARA-generated homology model (green) of *TamHydS*<sup>3</sup> was generated on the basis of *Cpl*'s crystal structure (PDB ID 4XDC)<sup>13</sup> and the C-terminal domain was generated by AlphaFold (red). The [4Fe-4S] clusters and the H-cluster are shown as sticks (C: grey, Fe: orange, S: yellow, N: blue, O: red). RMSD values indicate closer alignment of the homology model (1.541 Å) than the AlphaFold model (3.692 Å) with *Cpl*. The fourth [4Fe-4S] cluster in the C-terminal domain is visually represented by manually integrating a [4Fe-4S] cluster from the *Cpl* structure into the *TamHydS* AlphaFold model, binding to cysteines C379, C382, C387, and C404 (red sticks), via PyMOL.

rearrangement using Attenuated total reflectance-Fourier transform infrared (ATR-FTIR) spectroscopy. We propose that this secondary structure change is involved in the signalling mechanism of *TamHydS*. In a broader sense, our results shine light on how nature evolved to facilitate different functions via alternative protein scaffolds harbouring the identical cofactors.

To substantiate the putative H<sub>2</sub> sensing role, we analysed the genome of *T. mathranii* surrounding *TamHydS* (Fig. 2). We identified two operons located in direct vicinity harbouring genes for *TamHydS* and a group A [FeFe]-hydrogenase respectively. In Operon 1 upstream to the *TamHydS* gene, a Ser/Thr protein kinase is encoded that is proposed to be involved in signal transduction.<sup>2</sup> Further downstream, we identified a sequence that could be assigned to either a phosphotransferase,



**Fig. 2** Genome region surrounding *TamHydS*. Both *TamHydS* and a group A [FeFe]-hydrogenase are present (light green) in neighbouring operons. directly upstream of *TamHydS* a Ser/Thr protein kinase is found (861, orange), downstream, a DRTGG protein (863, light blue), followed by PHP domain protein (864, yellow) are found. In Operon 2 a NADH dehydrogenase (ubiquinone) and NADH dehydrogenase (quinone) are marked as blue, 865 and 868 respectively. A His kinase (866, purple) can be found downstream of the group A [FeFe]-hydrogenase. Conserved hypothetical proteins present in the region are marked as red arrows. Promoters are labelled as black arrows.

a polymerase and histidinol phosphatase (PHP), or a phosphoesterase PHP domain protein. Any of these could assume the putative signal transfer role of the Ser/Thr protein phosphatase found in the genomic context of group C [FeFe]-hydrogenases. However, in group C the phosphatase is proposed to be regulated by the PAS domain, which is absent in group D. Instead, the *T. mathranii* genome encodes a DRTGG protein immediately following the *TamHydS* gene. This DRTGG protein has been implicated in the regulation of pyrophosphatase in *Clostridium perfringens* and *Desulfotobacterium hafniense*.<sup>14,15</sup> In Operon 2, we found a group A [FeFe]-hydrogenase with a NADH dehydrogenase (ubiquinone), a histidine kinase and a NADH dehydrogenase (quinone) found upstream. This resembles the group A [FeFe]-hydrogenases commonly encoded downstream in the genome of organisms containing group C sensory [FeFe]-hydrogenases.<sup>1,2,4,11</sup> Collectively, these findings strongly suggest an involvement of *TamHydS* in H<sub>2</sub> sensing and signal transduction.

We investigated the potential H<sub>2</sub> sensing mechanism of *TamHydS* at a molecular level via ATR-FTIR difference spectroscopy. A solution of purified *TamHydS* enzyme was deposited on the surface of the ATR crystal, dried and rehydrated under humidified N<sub>2</sub> gas as reported previously.<sup>16</sup> We populated the potential signalling state, H<sub>red</sub>, in two ways (i) via H<sub>2</sub> uptake and (ii) via photo-reduction (Fig. 3 inset). Upon exposure to H<sub>2</sub> gas, we observed the formation of the diiron site reduced state and the potential signalling state, H<sub>red</sub>, to the expense of the oxidised resting state, H<sub>ox</sub>. The corresponding difference spectrum (Fig. 3 red) showed positive bands corresponding to H<sub>red</sub> (2063, 2032, 1922, 1896, 1802 cm<sup>-1</sup>) and negative bands associated with H<sub>ox</sub> (2082, 2074, 1970, 1948, 1787 cm<sup>-1</sup>), respectively. Populating H<sub>red</sub> via our previously developed photo-reduction protocol<sup>10</sup> resulted in a difference spectrum nearly identical to that obtained via H<sub>2</sub> exposure (Fig. 3 black spectrum). The band positions of H<sub>red</sub> enriched via the photo-reduction approach are slightly shifted (ca. 1 cm<sup>-1</sup>) mainly to lower wavenumbers, compared to the H<sub>red</sub> signature enriched via H<sub>2</sub> exposure, most likely due to additional reduction of F-clusters.<sup>17,18</sup> Furthermore, we detected a  $\mu$ CO band (1801–1803 cm<sup>-1</sup>) associated with H<sub>red</sub> for both reduction methods. We were able to monitor the co-population of all bands assigned to a single redox state over the course of the photo-reduction experiment (Table S1 and Fig. S1, ESI<sup>†</sup>). Beyond the





**Fig. 3** FTIR difference spectra showing the population of the signalling state,  $H_{red}$ , via  $H_2$  uptake and via photo-reduction. We observed population of the reduced state  $H_{red}$  (positive peaks: 2063, 2032, 1922, 1896, 1802  $cm^{-1}$ ) and a de-population of  $H_{ox}$  (negative peaks: 2082, 2074, 1970, 1948, 1787  $cm^{-1}$ ) upon exposure of *TamHydS* to either  $H_2$  (red spectrum) or illumination of *TamHydS* in the presence of eosin Y and EDTA as sacrificial electron donor (black spectrum, photo-reduction). The respective vibrational regions are indicated above the graph. Band positions are indicated by coloured bars. The  $H_2$  uptake spectrum is scaled by 0.3 for better comparability. (inset) Scheme illustrating the two ways to populate  $H_{red}$  starting from  $H_{ox}$  either via  $H_2$  uptake or photo-reduction.

fingerprint region of the H-cluster ligand bands (2120–1780  $cm^{-1}$ ), we detected difference features in the region of C=O vibrations (1750–1600  $cm^{-1}$ ), which will be discussed in detail in the next section.

In Fig. 4, the difference spectra in the C=O region of the  $H_{ox}$  to  $H_{red}$  transition induced via photo-reduction for group D [FeFe]-hydrogenase *TamHydS* (black) and group A representative *CrHydA1* (blue, data from ref.<sup>10</sup>) are overlaid. These spectra are scaled based on the negative band for  $H_{ox}$  to facilitate comparison (Fig. S2, ESI<sup>†</sup>). The C=O region can be separated into the C=O vibrations of carboxylic acid residues found between 1750–1690  $cm^{-1}$  and of C=O vibrations from the peptide back bone found at 1690–1600  $cm^{-1}$ , which is denoted as amide I vibration (Fig. 4).<sup>19,20</sup> In *CrHydA1*, the difference features related to carboxylic acid residues between 1750–1690  $cm^{-1}$  with its most prominent peaks at 1715 and 1700  $cm^{-1}$  report a rearrangement of the H-bonding network in their unique PTP.<sup>10,21</sup> *TamHydS* is missing the carboxylic acids assigned to the 1700/1715  $cm^{-1}$  feature in group A. Furthermore, no similar difference features were detected from 1750–1690  $cm^{-1}$ , indicating no changes in the H-bonding network of the novel PTP upon reduction. Instead, we observed difference features in the amide I C=O region (1690–1600  $cm^{-1}$ ). The main difference features are negative peaks at 1687 and 1672  $cm^{-1}$ , as well as a positive peak at 1649  $cm^{-1}$ , which all dominate the  $H_2$  induced difference spectrum as well (Fig. 4 and Fig. S2, ESI<sup>†</sup>). The 1687/1672/1649  $cm^{-1}$  difference signals correlate with the CO and CN<sup>-</sup> bands associated



**Fig. 4** FTIR difference spectra of the C=O region indicate secondary structure changes associated with the  $H_{ox}$  to  $H_{red}$  transition in *TamHydS*. In the  $H_{red}$ – $H_{ox}$  difference spectrum of *TamHydS* induced via photo-reduction (black) we observed a large difference feature in the amide I C=O region with a positive peak at 1649  $cm^{-1}$  and negative peaks at 1687/1672  $cm^{-1}$  which we assign to a change in secondary structure. In the difference spectrum of the same transition in group A [FeFe]-hydrogenase *HydA1* (blue spectrum) no changes in the amide I region were detected. Instead, in *CrHydA1* a difference feature in the carboxylic acid C=O region (1715 and 1700  $cm^{-1}$ ) indicate a rearrangement of the hydrogen bonding network of the PTP. In the same region no difference features indicative of a similar rearrangement of the PTP can be detected in *TamHydS*. The respective vibrational regions are indicated above the graph. The *HydA1* spectrum is scaled by a factor of 0.12 to match the intensity of the negative  $H_{ox}$  band to allow for comparison (Fig. S2, ESI<sup>†</sup>). *CrHydA1* data modified from ref. 10.

with  $H_{ox}$  and  $H_{red}$ , respectively (Fig. S1, ESI<sup>†</sup>). Note that in the *HydA1* difference spectrum, no changes of comparable intensity could be detected in the amide I C=O region.

When considering the alternative PTP of *TamHydS*, it would be tempting to assign the difference bands at 1671  $cm^{-1}$  and 1648  $cm^{-1}$  to changes of the carboxylic acid residue(s) (E252 and E289) that are close to the H-cluster, in analogy to group A [FeFe]-hydrogenases (compare Fig. 1). However, these difference bands are two to three times more intense and are found at lower wavenumbers when compared to group A [FeFe]-hydrogenase. Importantly, the C=O vibrations of carboxylic acid residues (1750–1690  $cm^{-1}$ ) in proteins are normally not found at these low wavenumbers (1672 and 1649  $cm^{-1}$ ).<sup>19,20</sup>

In *HydA1*, we could assign the low wavenumber band at 1681  $cm^{-1}$  to the deprotonation of an arginine. However, no arginine is found in the putative PTP of *TamHydS*.<sup>9</sup> Upon H/D exchange, the negative band at 1672  $cm^{-1}$  shifts by 2  $cm^{-1}$  and the positive band at 1649  $cm^{-1}$  by 4  $cm^{-1}$  to lower wavenumbers (Fig. S3, ESI<sup>†</sup>). These minor shifts argue against arginine or localised water molecules as the origin of the difference feature. The exclusion of these possibilities supports our assignment of the observed bands to changes in amide I



vibrations induced by secondary structure changes. Similar amide I band positions were assigned to secondary structure changes upon ferredoxin binding in group A [FeFe]-hydrogenase recently.<sup>22</sup>

Several characteristics of *TamHydS* indicate its role as a H<sub>2</sub> sensor. Once populated, the potential signalling state, H<sub>red</sub>, exhibits high stability.<sup>3,12</sup> Furthermore, the low turnover rates disfavour a catalytic purpose, similar to what has been observed for group C sensory [FeFe]-hydrogenases.<sup>3,12</sup> The alternative function of *TamHydS* is additionally supported by its different PTP mechanism when compared to group A representative HydA1. In catalytically highly active group A [FeFe]-hydrogenases, the alternative configuration of the PTP upon reduction was proposed as one of the factors enabling fast catalysis.<sup>10,21</sup> In contrast in *TamHydS*, we detected no such alternative configuration of the PTP that would favour the faster depopulation of the potential signalling state.

Instead, we detected secondary structure changes that we propose to be involved in a potential sensing mechanism. In other biological sensors, large secondary structure changes are well established to facilitate signalling (*e.g.*, the light-sensitive phytochrome proteins).<sup>23</sup> In H<sub>2</sub> sensing group C [FeFe]-hydrogenases, a PAS domain protein is the most likely signal receptor that triggers a subsequent signal cascade.<sup>2</sup> In *TamHydS*, this PAS domain is absent. However, we identified key enzymes in the genome region that could constitute an alternative signalling cascade. The secondary structure change detected in this study provides the first insight on how H<sub>2</sub> sensing in [FeFe]-hydrogenases is facilitated at a molecular level. Understanding and being able to modulate the H<sub>2</sub> metabolism in microbes, in particular in antibiotic resistant pathogens, holds promising potential for future treatment of diseases. From a more fundamental research perspective our study gives insight on how nature tunes the function of enzymes harbouring identical cofactors.

M. S. designed experiments, performed the FTIR spectroscopy, analysed data and wrote the first draft of the manuscript. I. V. performed genomic analysis. A. Z. performed the synthesis work. C. S., P. R. C. and H. L. isolated the proteins. P. R. C. performed structural alignment. All authors were involved in the analysis and revision of the manuscript and have given approval to the final version of the manuscript.

The authors thank Leopold Fichet for contributions to the FTIR spectroscopy experiments. The work presented in this article is supported by the Novo Nordisk Foundation (NNF23 OC0085682 to M. S.) and by the Swedish Research Council (2023-04593 to M. S. and I. V.). The Olle Engkvist stiftelse (grant no 220-0226 to M. S.) is gratefully acknowledged for funding.

## Data availability

The methods and data supporting the findings of this study are available in the ESI.†

## Conflicts of interest

There are no conflicts to declare.

## Notes and references

- H. Land, M. Senger, G. Berggren and S. T. Stripp, *ACS Catal.*, 2020, **10**, 7069–7086.
- Y. Zheng, J. Kahnt, I. H. Kwon, R. I. Mackie and R. K. Thauer, *J. Bacteriol.*, 2014, **196**, 3840–3852.
- H. Land, A. Sekretareva, P. Huang, H. J. Redman, B. Nemeth, N. Polidori, L. S. Meszaros, M. Senger, S. T. Stripp and G. Berggren, *Chem. Sci.*, 2020, **11**, 12789–12801.
- N. Chongdar, J. A. Birrell, K. Pawlak, C. Sommer, E. J. Reijerse, O. Rudiger, W. Lubitz and H. Ogata, *J. Am. Chem. Soc.*, 2018, **140**, 1057–1068.
- S. L. Benoit, R. J. Maier, R. G. Sawers and C. Greening, *Microbiol. Mol. Biol. Rev.*, 2020, **84**, DOI: [10.1128/mmlbr.00092-00019](https://doi.org/10.1128/mmlbr.00092-00019).
- F. Carbonero, A. C. Benefiel and H. R. Gaskins, *Nat. Rev. Gastroenterol. Hepatol.*, 2012, **9**, 504–518.
- C. Greening, P. R. Cabotaje, L. E. Valentin Alvarado, P. M. Leung, H. Land, T. Rodrigues-Oliveira, R. I. Ponce-Toledo, M. Senger, M. A. Klamke, M. Milton, R. Lappan, S. Mullen, J. West-Roberts, J. Mao, J. Song, M. Schoelmerich, C. W. Stairs, C. Schleper, R. Grinter, A. Spang, J. F. Banfield and G. Berggren, *Cell*, 2024, **187**, 3357–3372.
- S. Morra, *Front. Microbiol.*, 2022, **13**, 853626.
- P. R. Cabotaje, K. Walter, A. Zamader, P. Huang, F. Ho, H. Land, M. Senger and G. Berggren, *ACS Catal.*, 2023, **13**, 10435–10446.
- M. Senger, V. Eichmann, K. Laun, J. Duan, F. Wittkamp, G. Knor, U. P. Apfel, T. Happe, M. Winkler, J. Heberle and S. T. Stripp, *J. Am. Chem. Soc.*, 2019, **141**, 17394–17403.
- N. Chongdar, P. Rodriguez-Macia, E. J. Reijerse, W. Lubitz, H. Ogata and J. A. Birrell, *Chem. Sci.*, 2023, **14**, 3682–3692.
- H. Land, P. Ceccaldi, L. S. Meszaros, M. Lorenzi, H. J. Redman, M. Senger, S. T. Stripp and G. Berggren, *Chem. Sci.*, 2019, **10**, 9941–9948.
- J. Esselborn, N. Muraki, K. Klein, V. Engelbrecht, N. Metzler-Nolte, U. P. Apfel, E. Hofmann, G. Kurisu and T. Happe, *Chem. Sci.*, 2016, **7**, 959–968.
- H. Tuominen, A. Salminen, E. Oksanen, J. Jamsen, O. Heikkila, L. Lehtio, N. N. Magretova, A. Goldman, A. A. Baykov and R. Lahti, *J. Mol. Biol.*, 2010, **398**, 400–413.
- V. A. Anashkin, A. Salminen, V. N. Orlov, R. Lahti and A. A. Baykov, *Arch. Biochem. Biophys.*, 2020, **692**, 108537.
- M. Senger, T. Kernmayr, M. Lorenzi, H. J. Redman and G. Berggren, *Chem Commun.*, 2022, **58**, 7184–7187.
- P. Rodríguez-Maciá, N. Breuer, S. DeBeer and J. A. Birrell, *ACS Catal.*, 2020, **10**, 13084–13095.
- P. Rodríguez-Macia, K. Pawlak, O. Rudiger, E. J. Reijerse, W. Lubitz and J. A. Birrell, *J. Am. Chem. Soc.*, 2017, **139**, 15122–15134.
- V. A. Lorenz-Fonfria, *Chem. Rev.*, 2020, **120**, 3466–3576.
- A. Barth, *Biochim. Biophys. Acta.*, 2007, **1767**, 1073–1101.
- J. Duan, A. Hemschemeier, D. J. Burr, S. T. Stripp, E. Hofmann and T. Happe, *Angew. Chem., Int. Ed.*, 2023, **62**, e202216903.
- S. Şahin, J. Brazard, T. B. M. Adachi, R. D. Milton and S. T. Stripp, *ChemRxiv*, 2024, preprint, DOI: [10.26434/chemrxiv-2024-pnm11](https://doi.org/10.26434/chemrxiv-2024-pnm11).
- H. Takala, A. Björöling, O. Berntsson, H. Lehtivuori, S. Niebling, M. Hoernke, I. Kosheleva, R. Henning, A. Menzel, J. A. Ihalainen and S. Westenhoff, *Nature*, 2014, **509**, 245–248.

