

Cite this: *Chem. Sci.*, 2020, 11, 12789

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# Characterization of a putative sensory [FeFe]-hydrogenase provides new insight into the role of the active site architecture†

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[FeFe]-hydrogenases are known for their high rates of hydrogen turnover, and are intensively studied in the context of biotechnological applications. Evolution has generated a plethora of different subclasses with widely different characteristics. The M2e subclass is phylogenetically distinct from previously characterized members of this enzyme family and its biological role is unknown. It features significant differences in domain- and active site architecture, and is most closely related to the putative sensory [FeFe]-hydrogenases. Here we report the first comprehensive biochemical and spectroscopical characterization of an M2e enzyme, derived from *Thermoanaerobacter mathranii*. As compared to other [FeFe]-hydrogenases characterized to-date, this enzyme displays an increased H<sub>2</sub> affinity, higher activation enthalpies for H<sup>+</sup>/H<sub>2</sub> interconversion, and unusual reactivity towards known hydrogenase inhibitors. These properties are related to differences in active site architecture between the M2e [FeFe]-hydrogenase and “prototypical” [FeFe]-hydrogenases. Thus, this study provides new insight into the role of this subclass in hydrogen metabolism and the influence of the active site pocket on the chemistry of the H-cluster.

Received 15th June 2020  
Accepted 19th September 2020

DOI: 10.1039/d0sc03319g

rsc.li/chemical-science

## Introduction

Hydrogenase enzymes play a central role in hydrogen metabolism, where they catalyze the interconversion between protons and molecular hydrogen (H<sub>2</sub>). The [FeFe]-hydrogenases are generally considered the most active, operating close to the thermodynamic limit with reported H<sub>2</sub> production rates exceeding 9000 s<sup>-1</sup>.<sup>1,2</sup> Consequently, they have been intensively studied, both for their biotechnological potential and as a model system for the design of synthetic catalysts.<sup>3,4</sup> Phylogenetically, [FeFe]-hydrogenases can be broadly divided into four main groups, denoted group A, B, C, and D, which in turn contain numerous subclasses.<sup>5–9</sup> Considering the well-conserved nature of the auxiliary proteins involved in cofactor

assembly (HydEFG),<sup>6</sup> they all arguably share a dependence on the same hexanuclear iron cofactor, the “H-cluster”. This biologically unique cofactor consists of a canonical [4Fe–4S] cluster ([4Fe–4S]<sub>H</sub>) connected to a low valent dinuclear iron complex ([2Fe]<sub>H</sub>).<sup>10–13</sup> The [2Fe]<sub>H</sub> subsite is coordinated by CO and CN<sup>-</sup> ligands, and bridged by an azadithiolate ligand (adt = <sup>-</sup>SCH<sub>2</sub>NHCH<sub>2</sub>S<sup>-</sup>). The overwhelming majority of biochemically characterized [FeFe]-hydrogenases belong to group A, with a primary focus on the “prototypical” [FeFe]-hydrogenases, e.g., *Cr* HydA1 from *Chlamydomonas reinhardtii*,<sup>14,15</sup> *Dd* HydAB from *Desulfovibrio desulfuricans*,<sup>11,16,17</sup> *CpI* from *Clostridium pasteurianum*,<sup>10,18</sup> as well as the multimeric electron bifurcating [FeFe]-hydrogenase from *Thermotoga maritima*.<sup>18–20</sup> Studies of these enzymes form the foundation for our understanding of [FeFe]-hydrogenase biochemistry. Spectroscopy has identified numerous redox and protonation states of the H-cluster, around which various mechanistic proposals have been put forth.<sup>21–24</sup> In short, the active-ready resting state (H<sub>ox</sub>) features a mixed valence Fe(II)Fe(I) form of the [2Fe]<sub>H</sub> subsite and an oxidized [4Fe–4S]<sub>H</sub> cluster (2+). One-electron reduction results in either the H<sub>red'</sub> or H<sub>red</sub> state, where H<sub>red'</sub> features a reduced [4Fe–4S]<sub>H</sub> cluster while H<sub>red</sub> features a reduced and protonated [2Fe]<sub>H</sub> subsite.<sup>25</sup> Further reduction results in the formation of the so-called H<sub>hyd</sub> state featuring a terminal hydride on the [2Fe]<sub>H</sub> subsite.<sup>26–28</sup> Protonation of H<sub>hyd</sub> results in H<sub>2</sub> release, potentially proceeding *via* a discrete

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0sc03319g

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intermediate ( $H_{hyd}H^+$ ),<sup>27,29</sup> and returns the H-cluster to the  $H_{ox}$  state. Additionally, CO can reversibly bind to the H-cluster, giving rise to the inhibited  $H_{ox}$ -CO and  $H_{red}$ -CO states.<sup>30</sup>

Considering the diverse nature of [FeFe]-hydrogenase, both with regards to structure as well as function, it is clear that characterization of representative examples from other subclasses is necessary to complete our understanding of this enzyme family and H-cluster chemistry. It has repeatedly been shown that [FeFe]-hydrogenases can operate at minimal overpotentials, albeit specific enzymes generally display a bias for either  $H^+$  reduction or  $H_2$  oxidation.<sup>31–34</sup> Indeed, even in the relatively narrow selection of enzymes studied to-date significant differences in catalytic rates, stability of different H-cluster states and reactivity towards inhibitors (e.g., CO and  $O_2$ ) have been observed.<sup>22,31,35–39</sup> On a fundamental level, further insight

into subclass-specific reactivities is critical for our understanding of hydrogen metabolism, and elucidating the interplay between the H-cluster and the protein. It will also serve to strengthen efforts related to biotechnological energy applications and potentially facilitate the development of selective antibiotics.<sup>8,9</sup>

We recently reported the whole-cell characterization of an [FeFe]-hydrogenase from the thermophilic firmicute *Thermoanaerobacter mathranii* in *E. coli*.<sup>40</sup> The enzyme belongs to the hitherto uncharacterised M2e subclass, which displays a number of well-conserved differences in amino acid sequence as compared to the prototypical group A hydrogenases; namely in the active site cavity and the proton transfer pathway (Fig. 1).<sup>41</sup> The M2e subclass has been proposed to form a distinct group of [FeFe]-hydrogenases, group D, and their physiological function is

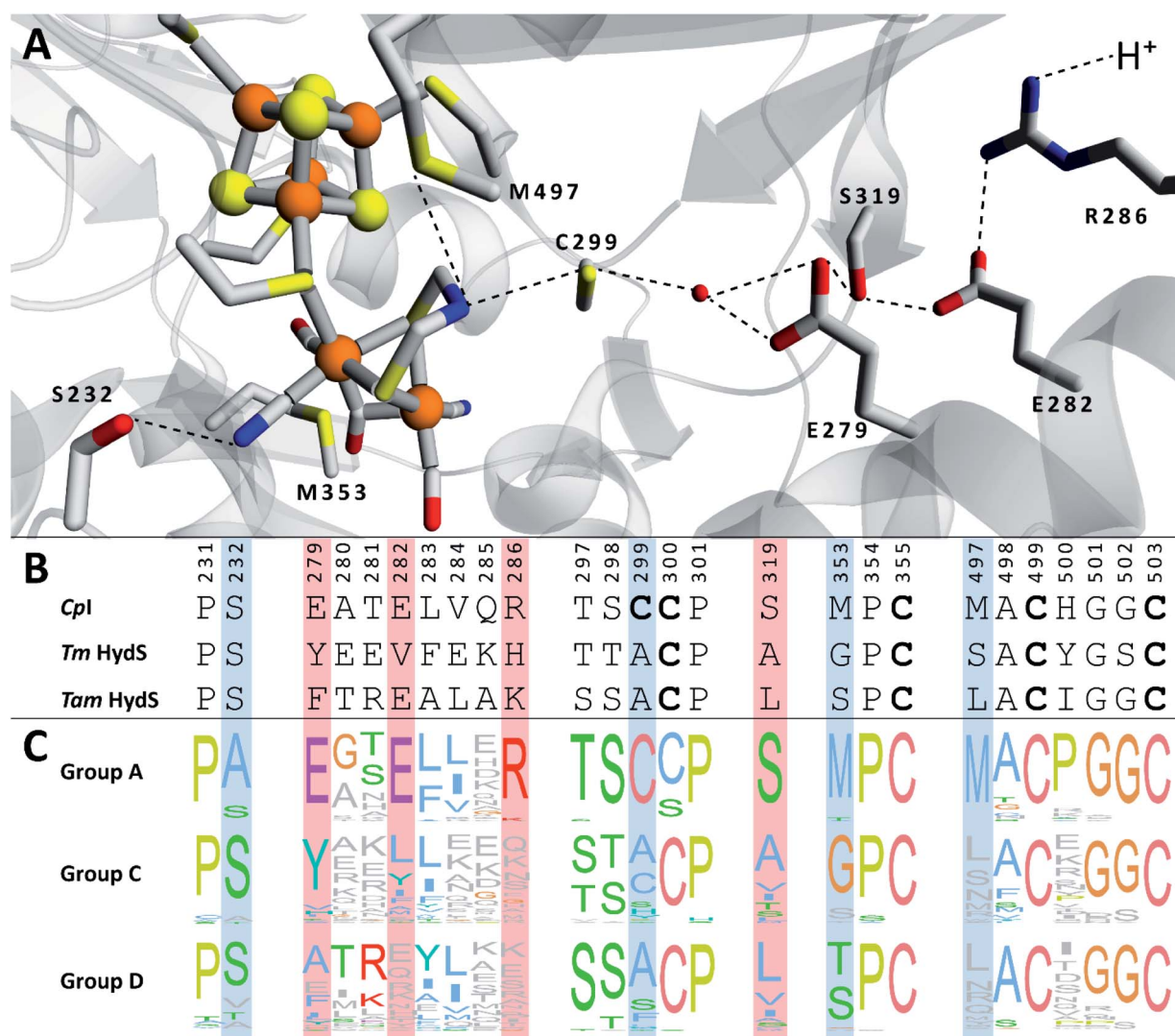


Fig. 1 (A) Structural view of the active site and proton transfer pathway of a prototypical [FeFe]-hydrogenase. Structure and numbering based on *Cpl* (PDB ID: 4XDC). Shown amino acid residues are either involved in interactions with the H-cluster or in the proton transfer pathway and show large variations between groups A, C and D. Potential interactions are shown with dashed lines. (B) Amino acid sequence comparison of *Cpl*, *Tm HydS* and *Tam HydS* (*Cpl* numbering) based on a Clustal $\Omega$  sequence alignment<sup>60</sup> of sequences retrieved from Greening *et al.* 2016 (ref. 7) and homology modeling. H-cluster interacting cysteine residues are highlighted in bold. (C) Normalized consensus logos of [FeFe]-hydrogenase groups A, C and D generated in Jalview using the sequence alignment in (B). Coloring is based on the Clustal X color scheme. Amino acid residues involved in H-cluster interaction and proton transfer that show variation between the groups are highlighted in blue and red, respectively.



unknown.<sup>5</sup> Still, it is phylogenetically most closely related to the M2f subclass of group C, which has been identified as “putative sensory” and includes the recently characterized *Tm* HydS enzyme from *Thermotoga maritima*.<sup>5,7,8,38</sup> In addition to the H-domain, which harbors the H-cluster, both subclasses feature an N-terminal domain with two [4Fe–4S] cluster-binding motifs as well as a C-terminal domain with a four-cysteine motif (Cx<sub>2</sub>-Cx<sub>4</sub>Cx<sub>16</sub>C) indicative of iron–sulfur (FeS) cluster binding. Additionally, the highly conserved cysteine residue initiating the proton transfer pathway from the adt amine in group A [FeFe]-hydrogenase is not conserved in the M2e and M2f subclasses (C299, *CpI* numbering is used throughout the text unless otherwise noted). Indeed, all residues that have previously been shown to be crucial for proton transfer in group A are missing in these subclasses (Fig. 1).<sup>42–44</sup> Moreover, the genes encoding M2e and M2f enzymes are often located on the same operon as group A hydrogenases.<sup>5</sup> In light of these similarities, we have now denoted the enzyme as *Tam* HydS (previously *Tam* HydA<sup>40</sup>). However, there are key differences between the two subclasses. In contrast to the M2f enzymes, the M2e subclass does not feature a Per-Arnt-Sim (PAS) domain, which was used to propose the sensing function of group C, as it is usually involved in the regulation of histidine kinases associated with signal transduction.<sup>5</sup> In addition, the active site architecture displays distinct differences between the two subclasses (Fig. 1). The two methionine residues framing the diiron site, M497 and M353, are exchanged against serine and glycine in *Tm* HydS; and leucine and serine, respectively, in *Tam* HydS. These methionines are considered critical for modulating the reactivity of prototypical [FeFe]-hydrogenases,<sup>31,41</sup> and it is noteworthy that the exchange of M353 to a hydroxyl donor (serine or threonine) appears to be a well-conserved property of M2e enzymes. On the other hand, M497 shows a low level of conservation, both for group C and D enzymes.

Herein, we report the first detailed characterization of an M2e enzyme, and by extension a group D [FeFe]-hydrogenase. The aforementioned *Tam* HydS enzyme has been isolated following heterologous expression, and characterized through electron paramagnetic resonance (EPR) spectroscopy, attenuated total reflection Fourier-transform infrared (ATR FTIR) spectroscopy, and protein film electrochemistry (PFE). Despite lacking several amino acid residues considered critical for the activity of prototypical [FeFe]-hydrogenase, *Tam* HydS shows reversible H<sup>+</sup>/H<sub>2</sub> interconversion close to the thermodynamic potential, with a slight bias for H<sup>+</sup> reduction. These findings show that the active site pocket can be significantly altered while still retaining the catalytic function of the H-cluster. However, in contrast to previously characterized [FeFe]-hydrogenases, an over-potential is observed at low temperatures; and the overall catalytic rates are low. Moreover, the M2e enzyme displays significant differences in its reactivity towards CO and O<sub>2</sub>, as compared to previously characterized [FeFe]-hydrogenases. We propose that the catalytic rates are influenced by intramolecular proton transfer. The study, furthermore, highlights the importance of the active site pocket in modulating the reactivity towards known inhibitors, and the stability of specific H-cluster states. Based on the aforementioned properties of *Tam* HydS which resemble those of known

regulatory [NiFe]-hydrogenases, in combination with analysis on a genome level, we hypothesize that group D [FeFe]-hydrogenases serve a sensory rather than catalytic function.

## Results and discussion

### Isolation and characterization of *apo-Tam* HydS

Aerobic heterologous expression of *Tam* HydS in *E. coli*, in the absence of the HydEFG maturation enzymes, has been shown to result in soluble *apo*-protein that could be activated *in vivo* using [2Fe]<sub>H</sub> subsite mimics.<sup>40</sup> In this study, anaerobic isolation of *apo-Tam* HydS resulted in a purified protein (Fig. S1†) containing 13.9 ± 0.85 Fe/protein. Additional reconstitution of the FeS clusters resulted in a final Fe content of 16.1 ± 0.29 Fe/protein, as expected from the incorporation of four [4Fe–4S] clusters. Isolation of *apo-Tam* HydS also under aerobic conditions provided a protein with an Fe content of 10.6 ± 0.06 Fe/protein.

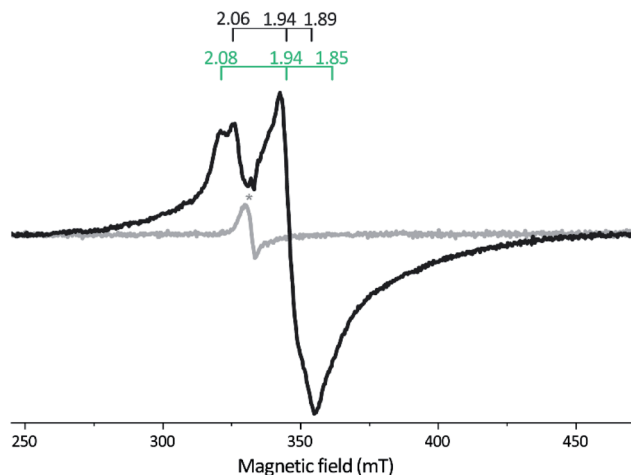
All three different purified forms of *apo-Tam* HydS displayed hydrogenase activity in *in vitro* sodium dithionite/methyl viologen (NaDT/MV<sup>2+</sup>) enzymatic assays, following anaerobic incubation with the synthetic [2Fe]<sub>H</sub> analogue [Fe<sub>2</sub>(adt)(CO)<sub>4</sub>(CN)<sub>2</sub>]<sup>2-</sup> ([2Fe]<sup>adt</sup>). Activation of aerobically and anaerobically purified *apo-Tam* HydS samples resulted in H<sub>2</sub> production activities of 0.068 ± 0.001 and 0.077 ± 0.009 μmol H<sub>2</sub> per min per mg, respectively. A further increase in the H<sub>2</sub> production activity, to 0.090 ± 0.005 μmol H<sub>2</sub> per min per mg was observed after activation of the reconstituted samples (Fig. 2).

X-Band EPR spectra recorded on samples of as-isolated and reconstituted *apo-Tam* HydS are essentially silent, indicating that all FeS-clusters are present at a [4Fe–4S]<sup>2+</sup> oxidation state with spin *s* = 0, thus diamagnetic. Reduction of the reconstituted enzyme by NaDT at pH 8 brought the sample to an EPR active state, displaying a broad rhombic EPR spectrum near the *g* ~ 2 region, typical of *s* = 1/2 [4Fe–4S]<sup>+</sup> species (Fig. 3). Spin



Fig. 2 Specific H<sub>2</sub> production activities of *Tam* HydS isolated under different conditions and activated *in vitro* with [2Fe]<sup>adt</sup>. The Fe content of the different preparations prior to insertion of [2Fe]<sup>adt</sup> were 10.9 ± 0.06 (aerobic purification), 13.9 ± 0.85 (anaerobic purification) and 16.1 ± 0.29 (reconstituted) Fe/protein. Reactions were performed in sodium phosphate buffer (100 mM, pH 6.8) with Triton X-100 (1% v/v), methyl viologen (10 mM) and sodium dithionite (100 mM).





**Fig. 3** X-band EPR spectra of reconstituted *apo-Tam* HydS (50  $\mu$ M). Gray spectrum: as prepared; black spectrum: NaDT reduced. The observed rhombic signal in the reduced sample is typical for  $[4\text{Fe}-4\text{S}]^+$  clusters. The  $g$ -tensors are indicated for the two contributing components, identified through comparison to NaDT reduced *holo-Tam* HydS (see Fig. S3†). The weak signal marked with \* appearing at  $g = 2.02$  is attributed to a trace amount of  $[3\text{Fe}-4\text{S}]^{2+}$ . EPR settings:  $T = 17$  K; modulation frequency 100 kHz, amplitude 15 G; microwave frequency 9.4 GHz, power 8 mW.

quantification resulted in two spin centers per protein, suggesting that two of the  $[4\text{Fe}-4\text{S}]$  clusters are susceptible to NaDT reduction.

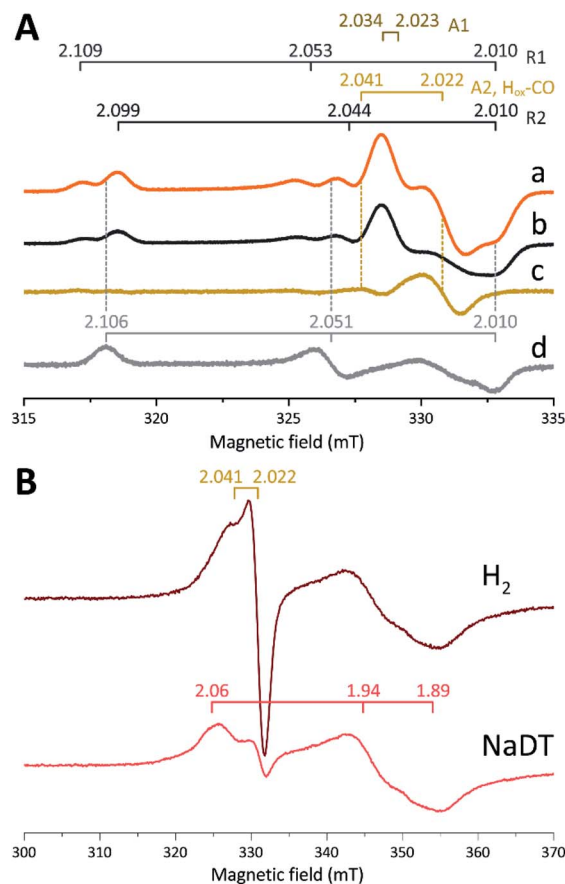
The iron quantification supports the presence of four  $[4\text{Fe}-4\text{S}]$  clusters in the reconstituted protein, including the active-site  $[4\text{Fe}-4\text{S}]_{\text{H}}$  cluster. This assignment is further supported by EPR spectroscopy. Albeit only two of the clusters were clearly discernable, no signal attributable to  $[2\text{Fe}-2\text{S}]$  clusters was observed upon reduction, and only traces of a  $[3\text{Fe}-4\text{S}]$  cluster signal were present in the non-reduced samples.<sup>45,46</sup> Similar results were recently reported for the putative sensory M2f enzyme *Tm* HydS, and it appears to be a shared trait between the two subclasses.<sup>38</sup> This FeS cluster composition is also supported by the linear increase in activity as iron content is increased up to 16 Fe/protein. Although the  $\text{H}_2$  production activity of *Tam* HydS, observed in *in vitro* assays, is low compared to most known prototypical  $[\text{FeFe}]$ -hydrogenases, it is similar to *Tm* HydS.<sup>38</sup> As the two N-terminal  $[4\text{Fe}-4\text{S}]$  clusters are well conserved across numerous  $[\text{FeFe}]$ -hydrogenases, including the aforementioned *Dd* HydAB as well as the M2 enzyme from *Megasphaera elsdenii* this arguably represents the catalytic electron transfer pathway also in the M2e enzymes.<sup>21,32,39</sup> In contrast, the biochemical and physiological role of the  $[4\text{Fe}-4\text{S}]$  cluster coordinated by the C-terminal  $\text{Cx}_2\text{Cx}_4\text{Cx}_{16}\text{C}$  motif is unknown. As it is located between the H-domain and the PAS domain in M2f  $[\text{FeFe}]$ -hydrogenases, it is likely to be involved in a signaling process.

### EPR characterization of *holo-Tam* HydS

The *in vitro* enzymatic assays revealed that treating *apo-Tam* HydS with  $[2\text{Fe}]^{\text{adt}}$  resulted in spontaneous formation of *holo-*

*Tam* HydS on a minute time-scale, similar to what was previously observed under whole-cell conditions.<sup>40</sup> In the absence of reductant (NaDT), this treatment is expected to yield an oxidized form of the H-cluster, either the  $\text{H}_{\text{ox}}$  state or the  $\text{H}_{\text{ox}}\text{-CO}$  state.<sup>13,47-49</sup> Both states are paramagnetic and best described as  $[4\text{Fe}-4\text{S}]_{\text{H}}^{2+}-[2\text{Fe}(\text{II},\text{I})]$  ( $s = 1/2$ ) species, thus EPR spectroscopy can be employed to monitor H-cluster assembly. It is well established that the EPR spectra of  $\text{H}_{\text{ox}}$  and  $\text{H}_{\text{ox}}\text{-CO}$  generally display rhombic and axial signals, respectively, with small anisotropy and spin transitions at  $g \approx 2$ .<sup>22</sup>

Expecting a mixture of  $\text{H}_{\text{ox}}$  and  $\text{H}_{\text{ox}}\text{-CO}$ , the X-band EPR spectra collected on solution samples of as-prepared *holo-Tam* HydS, generated under an inert argon atmosphere, displayed a surprisingly complex pattern. As seen in Fig. 4A (spectrum a),



**Fig. 4** (A) Signals observed for *holo-Tam* HydS and attributed to the H-cluster, (a) *holo-Tam* HydS activated with  $[2\text{Fe}]^{\text{adt}}$  as isolated; (b) *holo-Tam* HydS flushed by  $\text{N}_2$  to remove  $\text{H}_{\text{ox}}\text{-CO}$  contributions (contains components R1, R2, A1 and A2); (c) difference spectra obtained from subtraction of spectrum (b) from (a) revealing an axial signal attributed to  $\text{H}_{\text{ox}}\text{-CO}$  ("component A2,  $\text{H}_{\text{ox}}\text{-CO}$ "); (d) *Tam* HydS activated with  $[2\text{Fe}]^{\text{pdt}}$  (*pdt-Tam* HydS, as isolated). EPR settings:  $T$  21 K; modulation frequency 100 kHz, amplitude 10 G; microwave frequency 9.4 GHz, power 16  $\mu$ W. (B) Reduction of *holo-Tam* HydS activated with  $[2\text{Fe}]^{\text{adt}}$  by  $\text{H}_2$  and NaDT resulting in disappearance of the H-cluster signals and appearance of a rhombic signal attributed to a  $[4\text{Fe}-4\text{S}]^+$  cluster. EPR settings:  $T$  10 K; modulation frequency 100 kHz, amplitude 15 G; microwave frequency 9.4 GHz, power 80  $\mu$ W.  $g$ -values indicated in the figure on horizontal bars, for details see main text.

at least seven features were resolved. The spectrum displayed characteristics suggesting the presence of  $H_{ox}$  and  $H_{ox-CO}$  but the unusual complexity of the spectrum shows that more than two species contribute to the overall spectrum. Spin quantification of a representative spectrum resulted in 0.64 spin per protein, indicating that a fraction of the enzyme also resided in an EPR silent state, assigned by FTIR spectroscopy to the  $H_{red}$  state (see below). The relaxation behavior of the signal(s) for *holo-Tam* HydS was estimated by monitoring the dominant  $g \approx 2.04$ – $2.02$  feature (Fig. S4†). All components indicated in the spectrum followed similar saturation trends, and displayed low  $P_{1/2}$  values (73  $\mu$ W and 1.15 mW at 15 and 21 K, respectively). This suggests that spin relaxation is a slow process, most likely due to isolation of the H-cluster from the lattice.

Flushing of *holo-Tam* HydS solutions with  $N_2$  prior to freezing resulted in approx. 50% decrease of total spin density, reflected in a minor decrease in amplitude of the low-field features ( $g \approx 2.1$ ) while a pseudo-axial component represented by a feature at  $g = 2.022$  was almost completely lost (Fig. 4A, spectrum b). Subtraction of the spectrum obtained following  $N_2$  flushing (spectrum b) from the spectrum of the as-prepared sample (spectrum a) provided a “pure” pseudo-axial spectrum (Fig. 4A, spectrum c) with  $g_{||} = 2.041$  and  $g_{\perp} = 2.022$ . This signal is assigned to the  $H_{ox-CO}$  state (Fig. 4A, component “A2,  $H_{ox-CO}$ ”). Similarly, the  $H_{ox-CO}$  component was diminished in samples of *holo-Tam* HydS isolated following H-cluster assembly in the presence of NaDT (Fig. S5†). The residual spectrum (spectrum b) appears to feature two sets of rhombic signals (components R1 and R2), in combination with an additional narrow axial signal (component A1). The  $g_{||} = 2.034$  tensor of the latter signal is readily apparent while the  $g_{\perp}$  tensor is tentatively assigned to 2.023 from simulations (indicated in dark brown in Fig. 4A, see Fig. S5† for simulation details). With regards to the rhombic signals, the  $g$ -values are 2.109 and 2.099 for the two features at the lower field wing, 2.053 and 2.044 in the centre region and 2.010 in the high field wing. The overall spectral shape was highly similar between biological repeats prepared at pH 8. The relative signal amplitudes observed in spectrum (b) were also retained at pH 5 (Fig. S5†) but this acidification resulted in a 2–3 gauss downshift of one set of the rhombic EPR signals, which facilitated separation into two sets of separate  $g$ -tensors ( $g_{zyx} = 2.109, 2.053, 2.010$ , “component R1”; and  $g_{zyx} = 2.099, 2.044, 2.010$ , “component R2”; Fig. 4A). It should be noted that the assignment of  $g_x = 2.010$  is speculative due to its overlap with the adjacent axial signal.

To further clarify the EPR spectrum observed for *Tam* HydS, the *holo*-enzyme was generated using the modified cofactor  $[Fe_2(pdt)(CO)_4(CN)_2]^{2-}$  ( $[2Fe]^{pdt}$ ,  $pdt = ^-SCH_2CH_2CH_2S^-$ ; *pdt-Tam* HydS). It has been shown for both group A and C  $[FeFe]$ -hydrogenases that replacing the amine-bridgehead of the adt ligand with a methylene group destabilizes the  $H_{ox-CO}$  state.<sup>13,30,38,50,51</sup> Thus, analogous samples where  $[2Fe]^{pdt}$  replaced  $[2Fe]^{adt}$  were examined by EPR under the same recording conditions. The obtained spectrum showed a rhombic anisotropy with  $g$ -values of 2.106, 2.051 and 2.010 (Fig. 4A, spectrum d). This signal is attributed to the formation of a single pure  $H_{ox}$  state, in good agreement with FTIR spectroscopy (Fig. S6†), as

well as earlier studies of *Tam* HydS under whole-cell conditions.<sup>40</sup> A comparison between *holo-Tam* HydS generated with  $[2Fe]^{adt}$  (Fig. 4A, spectra a and b) and  $[2Fe]^{pdt}$  (Fig. 4A, spectrum d) reveals that the rhombic features observed in spectra (a) and (b) display a significant overlap with the signal observed for *pdt-Tam* HydS (indicated with dashed lines in Fig. 4A). Consequently, the rhombic components R1 and R2 are assigned to two distinct  $H_{ox}$ -like species. The non-overlapping features of the spectrum correspond to the pseudo-axial components (A1 and A2), in agreement with the assignment of A2 to the  $H_{ox-CO}$  state.

Reduction of *holo-Tam* HydS with NaDT resulted in disappearance of the aforementioned H-cluster signals with concomitant appearance of a broader rhombic EPR spectrum (Fig. 4B, spectrum NaDT). The loss of the  $H_{ox-CO}$  and  $H_{ox}$ -like signals is attributed to a one-electron reduction of the H-cluster to the diamagnetic  $H_{red}$  state (see below). The new signal partially resembles that observed for reduced *apo-Tam* HydS (Fig. 3), and spin quantification showed one spin per protein. Thus, one  $[4Fe-4S]$  cluster, with an EPR signature of  $g_{zyx} = 2.06, 1.94$  and  $1.89$ , is susceptible to NaDT reduction in *holo-Tam* HydS. Subtraction of the *holo-Tam* HydS signal from that of *apo-Tam* HydS revealed another broad rhombic EPR signal ( $g_{zyx} = 2.08, 1.94$  and  $1.85$ , see Fig. S3,† green spectrum). As this signal was present in the *apo*-protein but lost upon H-cluster formation, it is tentatively attributed to the  $[4Fe-4S]_H$  cluster of *apo-Tam* HydS. Reduction of *holo-Tam* HydS with  $H_2$  provided a similar result compared to reduction with NaDT, although a larger fraction of the  $H_{ox-CO}$  state remained (Fig. 4B, spectrum  $H_2$ ).

In summary, the combined EPR data from as-prepared and gas-flushed solution samples reveal an unusually complex mixture of oxidized states. Still, three of the contributing species can be assigned with relatively high certainty. Comparison of as-prepared and  $N_2$  flushed samples show that a standard  $H_{ox-CO}$  species can form also in *holo-Tam* HydS. Conversely, the “split”  $H_{ox}$ -like signal observed in *holo-Tam* HydS is suggestive of the formation of two distinguishable  $H_{ox}$ -like states. Based on FTIR spectroscopy (see below), R2 is attributed to a state highly similar to the well-known  $H_{ox}$  state of prototypical  $[FeFe]$ -hydrogenases ( $g_{zyx} = 2.099, 2.044, 2.010$ ), and the second rhombic EPR signal, R1, to a state similar to  $H_{ox}H$  ( $g_{zyx} = 2.109, 2.053, 2.010$ ). A similar downshift of the  $H_{ox}$ -signal upon formation of  $H_{ox}H$  has recently been reported for *Cr* HydA1.<sup>9</sup> As comparing spectra of samples prepared at mildly basic and acidic conditions did not reveal significant changes in the relative amplitudes of the rhombic signals their interconversion appears to be more complicated than an acid–base equilibrium. The structural details of this  $H_{ox}H$ -like state in *Tam* HydS remains to be fully elucidated. Still, both appear catalytically competent, as they were both lost upon exposure to  $H_2$ .

### FTIR characterization of *holo-Tam* HydS

The H-cluster of *holo-Tam* HydS was further investigated using ATR FTIR spectroscopy at different pH values and in the presence of  $H_2$ ,  $N_2$ , CO, or  $O_2$ . The enzyme adopted  $H_{red}$  as a semi-



stable resting state under our experimental conditions (20 °C, N<sub>2</sub> atmosphere with approx. 1% H<sub>2</sub>, hydrated protein films at pH 7). A quantitative enrichment of H<sub>ox</sub> was achieved only after 20–30 h of continuous purging with pure N<sub>2</sub>. In contrast to what is generally reported for prototypical [FeFe]-hydrogenases,<sup>52</sup> the H<sub>ox</sub>-state ( $\approx 65\%$ ) accumulated together with a small fraction ( $\approx 25\%$ ) of an alternative state displaying an H<sub>ox</sub>-like spectrum at higher frequencies (Fig. S7†). This hypsochromically-shifted signature is attributed to an H<sub>ox</sub>H-like state, albeit this state is generally not observed at neutral pH in prototypical [FeFe]-hydrogenases. This mixture of H<sub>ox</sub>-like states and their relative ratio is in agreement with the observation of two rhombic EPR signals under similar conditions (Fig. 4). The H<sub>red</sub> to H<sub>ox</sub> transition was further analyzed by ATR FTIR spectro-electrochemistry, revealing two unusual properties of the *Tam* HydS enzyme (Fig. S8†). The reaction displayed a significant over-potential requirement, and while the quasi-reversible nature of the process prevented an exact assignment of the H<sub>ox</sub>/H<sub>red</sub> reduction potential it was clearly shifted in an anodic direction as compared to previously studied prototypical [FeFe] hydrogenase, with  $E_{m,s}$  of  $\approx -350$  to  $-450$  mVs reported for *Cr* HydA1 and *Dd* HydAB.<sup>39,53,54</sup> During the reductive scan, H<sub>ox</sub>/H<sub>red</sub> interconversion in *Tam* HydS was observed at  $E_m \approx -300$  mV vs. SHE, while the re-oxidation did not occur until a potential of approx. +200 mV was applied (at pH 8). No intermediates were observed in the process. Thus, H<sub>red</sub> appears to be both kinetically and thermodynamically stabilized in *Tam* HydS. A relatively anodic H<sub>ox</sub>/H<sub>red</sub> midpoint potential has been reported also for the putative sensory Group C hydrogenase *Tm* HydS.<sup>38</sup>

Fig. 5A shows the IR signatures of H<sub>ox</sub>, H<sub>ox</sub>-CO, and H<sub>red</sub> observed for *Tam* HydS. The assignment of the spectra to specific H-cluster states was facilitated by their overall similarities to spectra previously reported for prototypical [FeFe]-hydrogenases. Still, the frequencies of the terminal CO/CN<sup>-</sup> ligands are upshifted in comparison to *Cr* HydA1 and *Dd* HydAB and closer to *CpI* and *CaI* from *C. acetobutylicum*.<sup>31,37,52,55</sup> The high-frequency CO band of H<sub>ox</sub>-CO (2026 cm<sup>-1</sup>) indicates a constrained geometry.<sup>51</sup> In contrast to these upshifts, the  $\mu$ CO band of H<sub>ox</sub> (1788 cm<sup>-1</sup>) and H<sub>ox</sub>-CO (1786 cm<sup>-1</sup>) was found at lower frequencies than typically observed. These latter differences, as compared to group A and C [FeFe]-hydrogenases, are likely attributable to the M393S variation in *Tam* HydS as this residue is in close contact with the bridging  $\mu$ CO ligand. A distinct feature in the  $\mu$ CO region for the H<sub>red</sub> state could not be discerned. At low pH and high concentrations of NaDT, accumulation of H<sub>ox</sub>H over H<sub>ox</sub> was achieved (Fig. 5B), although the protein film never fully converted into H<sub>ox</sub>H (Fig. S7†). Note that the signature of H<sub>ox</sub>H at low pH is in excellent agreement with the hypsochromically shifted H<sub>ox</sub>-like spectrum observed at pH 8. In contrast to what has been reported for prototypical [FeFe]-hydrogenases, no accumulation of H<sub>hyd</sub> was observed, e.g., when low pH samples were exposed to H<sub>2</sub>.<sup>56</sup> Moreover, H<sub>sred</sub> and H<sub>red</sub> were never detected. Table 1 summarizes the IR signature of all observed H-cluster states.

The absorbance spectra of the H-cluster states identified in Fig. 5 were fitted and used to describe the interconversion reactions as a function of gas composition and time. For

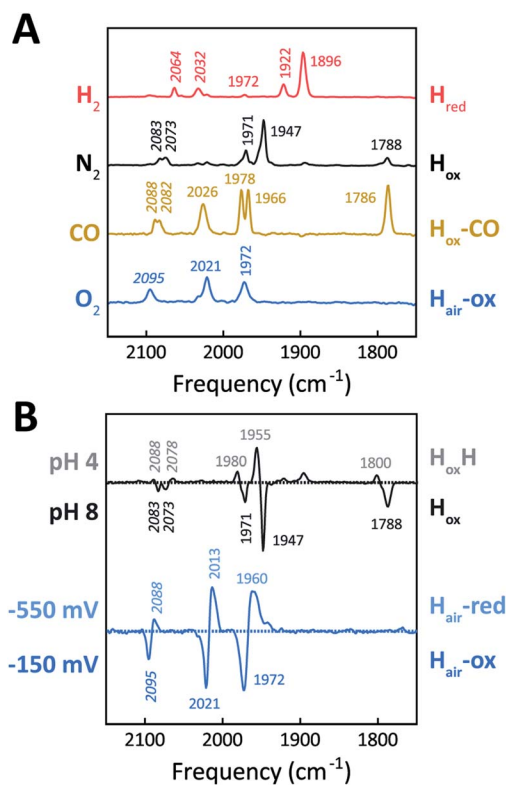


Fig. 5 ATR FTIR characterization of *holo-Tam* HydS. All data recorded at RT. (A) Infrared signature of the H-cluster in *Tam* HydS in the presence of H<sub>2</sub> (red, H<sub>red</sub>), N<sub>2</sub> (black, H<sub>ox</sub> and H<sub>ox</sub>H), CO (brown, H<sub>ox</sub>-CO), and after reaction with O<sub>2</sub> (blue, H<sub>air</sub>). (B) The upper difference spectrum shows accumulation of H<sub>ox</sub>H (positive bands, grey, pH 4) over H<sub>ox</sub> (negative bands, black, pH 8) under N<sub>2</sub>. Note: accumulation of H<sub>ox</sub>H also required NaDT addition. In the lower difference spectrum, FTIR spectro-electrochemistry was used to accumulate H<sub>air</sub>-red (positive bands, light blue,  $-550$  mV vs. SHE) over H<sub>air</sub>-ox (negative bands, blue,  $-150$  mV).

comparative purposes, analogous experiments were performed with the prototypical [FeFe]-hydrogenases *Dd* HydAB or *Cr* HydA1. Fig. 6A depicts the rapid conversion of H<sub>ox</sub> into H<sub>red</sub> for *Tam* HydS and *Dd* HydAB at 1%, 10%, and 100% H<sub>2</sub> over N<sub>2</sub>. *Dd* HydAB was chosen for comparison because it shows a similar, albeit not identical, composition of reduced H-cluster states under H<sub>2</sub> (Fig. S10†). The identity of H<sub>red</sub> as resting state in *Tam* HydS is illustrated by the pronounced persistence of H<sub>red</sub> when H<sub>2</sub> was removed from the gas phase ( $t > 16.5$  min) whereas *Dd* HydAB immediately converted into H<sub>ox</sub>. This accumulation of H<sub>ox</sub> is a consequence of auto-oxidation, i.e. due to H<sub>2</sub> release. In

Table 1 Vibrational frequencies observed for the CN<sup>-</sup> and CO ligands

	CN <sup>-</sup> (cm <sup>-1</sup> )		CO (cm <sup>-1</sup> )		
H <sub>red</sub>	2064	2032	1972	1922	1896
H <sub>ox</sub>	2083	2073	1971	1947	1788
H <sub>ox</sub> H	2088	2078	1980	1955	1800
H <sub>ox</sub> -CO	2088	2082	2026	1978	1966
H <sub>air</sub> -ox	2095		2021	1972	
H <sub>air</sub> -red	2088		2013	1960	

the next step, the influence of temperature on the  $H_{\text{red}}$  to  $H_{\text{ox}}$  transition of *Tam* HydS and *Dd* HydAB was investigated. We addressed the kinetics of auto-oxidation for five temperature points in the range between 20–40 °C. The enzymes were reduced in the presence of 1%  $H_2$  and subjected to pure  $N_2$  for 10 min, before they were re-reduced with 1%  $H_2$ . Fig. 6B depicts the changing population of  $H_{\text{ox}}$  in *Tam* HydS as a function of gas, time, and temperature. Higher temperature increased the rate of  $H_{\text{ox}}$  formation, upon removal of  $H_2$  from the atmosphere, and induced a higher percentage of  $H_{\text{ox}}$  accumulation (*i.e.*, after 10 min). The same set of experiments was performed for *Dd* HydAB. Albeit apparent instability of the H-cluster in *Dd* HydAB at  $T > 30$  °C prevented a complete study, the net-oxidation rate in *Dd* HydAB is significantly higher than in *Tam* HydS (Fig. S11†). Moreover, it should be noted that an increase in temperature also resulted in an increase in steady-state concentration of  $H_{\text{ox}}$  already when *Tam* HydS equilibrated under 1%  $H_2$ , highlighting a positive entropy contribution for the  $H_{\text{red}}$  to  $H_{\text{ox}}$  transition (Fig. 6B).

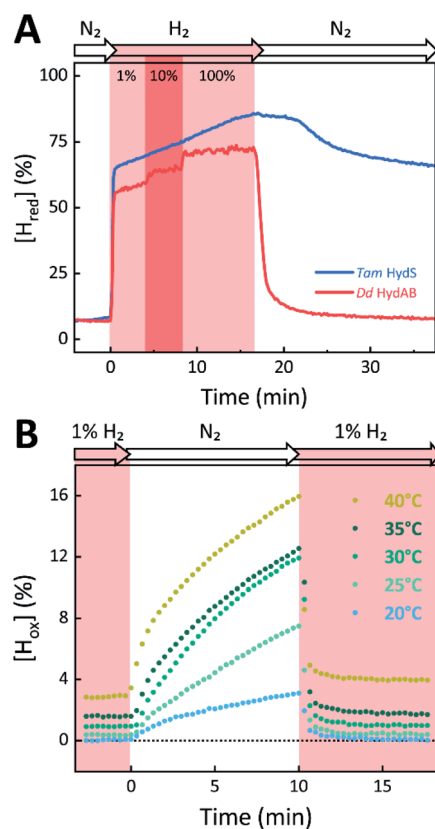


Fig. 6 The reactivity of *Tam* HydS and prototypical [FeFe]-hydrogenase (*Dd* HydAB) towards  $H_2$  (A) and temperature dependence of the auto-oxidation activity (B) monitored by time-resolved ATR FTIR spectroscopy. (A) Kinetic traces of  $H_{\text{red}}$  for *Tam* HydS (blue) and *Dd* HydAB (red) that show the reaction with different concentrations of  $H_2$  (RT). Note the persistence of  $H_{\text{red}}$  in the absence of  $H_2$  for *Tam* HydS. (B) Change of  $[H_{\text{ox}}]$  as a function of gas, time, and temperature. Representative data set for *Tam* HydS, recorded at pH 7. The equilibrium under 1%  $H_2$  is slightly shifted in favour of  $H_{\text{ox}}$  over  $H_{\text{red}}$  as temperature is increased, and so does rate of  $[H_{\text{ox}}]$  formation as the atmosphere is changed to pure  $N_2$ .

The reactivity towards known [FeFe]-hydrogenase inhibitors was probed by exposing protein films to CO or  $O_2$ . Fig. 7A depicts the conversion of  $H_{\text{ox}}$  into  $H_{\text{ox}}\text{-CO}$  for *Tam* HydS and *Dd* HydAB at 1%, 10%, and 100% CO over  $N_2$ . Here, *Tam* HydS displays a notable lack of CO inhibition. Even under 100% CO, only 60% of the H-cluster population converted into  $H_{\text{ox}}\text{-CO}$ . Similar trends were observed with 10%  $H_2$  in the  $N_2$  carrier gas (Fig. S10†). In the absence of CO gas,  $H_{\text{red}}$  recovered quickly. Adjusted for the CO-insensitive contamination of  $\sim 30\%$   $H_{\text{inact}}$  (Fig. S10†), *Dd* HydAB showed immediate, complete, and enduring CO inhibition. Fig. 7B depicts the reaction of oxidized *Tam* HydS and *Cr* HydA1 with 1 atm air. We chose *Cr* HydA1 for comparison because *Dd* HydAB partly converted into unready states like  $H_{\text{inact}}$  in the presence of  $O_2$  whereas  $O_2$  exposure rapidly destroyed the H-cluster in *Cr* HydA1 (Fig. S12†). In contrast, *Tam* HydS converted into an unprecedented species, denoted  $H_{\text{air}}$ , and the formation of this state was observed regardless of whether the H-cluster resided in the  $H_{\text{red}}$  or  $H_{\text{ox}}$  state upon  $O_2$  exposure (Fig. S9†). As seen in Fig. 5A (blue

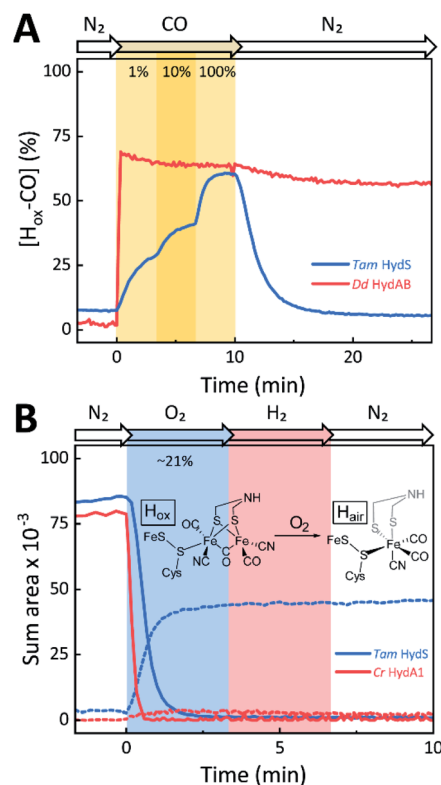


Fig. 7 The reactivity of *Tam* HydS and prototypical [FeFe]-hydrogenase (*Dd* HydAB or *Cr* HydA1) towards CO (A) and  $O_2$  (B) monitored by time-resolved ATR FTIR spectroscopy. (A) Kinetic traces of  $H_{\text{ox}}\text{-CO}$  for *Tam* HydS (blue) and *Dd* HydAB (red) that show the reaction with different concentrations of CO. About 30% of the *Dd* HydAB sample were arrested in the CO-insensitive  $H_{\text{inact}}$  state. Thus, the observed accumulation of  $\sim 70\%$   $H_{\text{ox}}\text{-CO}$  can be considered complete. (B) Kinetic traces for *Tam* HydS (blue) and *Cr* HydA1 (red) that show the reaction with  $\sim 21\%$   $O_2$  (air). Solid traces depict  $H_{\text{ox}}$ . While nearly 100% of the H-cluster is lost in *Cr* HydA1, a stable  $\text{Fe}(\text{CO})_2\text{CN}$  species prevails in *Tam* HydS (dashed traces). The proposed reaction between the oxidized H-cluster ( $H_{\text{ox}}$ ) and  $O_2$  is depicted.  $\text{FeS} = [4\text{Fe}-4\text{S}]_{\text{H}}$ .



spectrum) this new state featured two bands in the CO region and one band in the  $\text{CN}^-$  region of the spectrum, suggestive of partial degradation of the  $[\text{2Fe}]_{\text{H}}$  subsite. Moreover, it was found to be unreactive towards  $\text{N}_2$ ,  $\text{H}_2$ , and CO. EPR samples collected of *Tam* HydS exposed to air did not reveal any discernable EPR signal, apart from minor features at  $g \approx 4.3$  and 2.02, attributable to small amounts of  $\text{Fe}^{3+}$  ions (“junk iron”) and  $[\text{3Fe-4S}]$  cluster species, respectively. Similarly, a NaDT reduced anaerobic sample of  $\text{H}_{\text{air}}$  was also essentially EPR silent, albeit trace amounts of a  $[\text{4Fe-4S}]^+$  species became discernable (Fig. S13†). A mononuclear version of the  $[\text{2Fe}]_{\text{H}}$  subsite has previously been observed by X-ray crystallography in the prototypical *Cpl*  $[\text{FeFe}]$ -hydrogenase, following extended  $\text{O}_2$  exposure of the enzyme *in crystallo*.<sup>57</sup> The overall FTIR spectral features in combination with  $^{13}\text{CO}$  isotope editing clearly supports the assignment of a mononuclear  $\text{Fe}(\text{CO})_2\text{CN}$  species (Fig. S14†). Spectroelectrochemistry also suggests that this mononuclear complex is bound to the  $[\text{4Fe-4S}]_{\text{H}}$  cluster and that the modified H-cluster displays at least one redox transition, enabling accumulation of “ $\text{H}_{\text{air-red}}$ ” and “ $\text{H}_{\text{air-ox}}$ ” (Fig. 5B, S14 and S15†).

### The catalytic properties of *Tam* HydS

The catalytic properties of *holo-Tam* HydS were investigated using protein film electrochemistry (PFE), revealing pH and temperature dependent catalytic currents for both  $\text{H}^+$  reduction and  $\text{H}_2$  oxidation (Fig. 8A). Four different procedures were tested for electrode immobilization of the enzyme with the best method being absorption on a pyrolytic graphite electrode in the presence of the polycationic polymyxin B sulfate (Fig. S16†). The observation that the enzyme favors interaction with a positively charged surface suggests that it has a negative net surface charge at the pH of immobilization (pH 7, theoretical  $\text{pI} = 5.87$ , ExPASy ProtParam tool).

It is important to note that in all experiments even at large over-potentials for both  $\text{H}^+$  reduction and  $\text{H}_2$  oxidation the catalytic current does not reach a steady-state value, but increases almost linearly with over-potential. Such behavior has been rationalized by disorder among the adsorbed enzyme molecules, resulting in a dispersion of interfacial electron

transfer rate constants.<sup>58–60</sup> In this case, the steady-state limiting current ( $i_{\text{lim}}$ ) can be estimated from a linear fit of the high driving force part of the cyclic voltammograms (CVs), where the slope ( $\partial i/\partial E$ ) is:<sup>60,61</sup>

$$\partial i/\partial E = \frac{i_{\text{lim}}}{\beta d_0} \frac{F}{2RT} \quad (1)$$

Eqn (1) predicts that the product of the slope and temperature is proportional to the limiting current and therefore to the activity.

We first evaluated the enzyme affinity towards  $\text{H}_2$ . It has been noted earlier that  $K_{\text{M}}^{\text{H}_2}$  values determined from PFE experiments can be potential-dependent.<sup>62,63</sup> Therefore, we recorded CVs at various concentrations of  $\text{H}_2$  in a broad potential window (Fig. S17†). For  $K_{\text{M}}^{\text{H}_2}$  estimation it is important to measure the current response under conditions where it is limited by the catalytic rate of the enzyme, *i.e.* proportional to the catalytic rate rather than mass transport or interfacial electron transfer.<sup>64</sup> Thus,  $K_{\text{M}}^{\text{H}_2}$  values at various over-potentials were calculated, and the measurements were performed at 30 and 60 °C. Moreover, to ensure that the catalytic rate is not limited by mass transport, CVs were recorded at rotation rates of 2000 and 3000 rpm at each concentration of  $\text{H}_2$ . At over-potentials starting from 200 mV (30 °C) and 100 mV (60 °C), calculated  $K_{\text{M}}^{\text{H}_2}$  values were identical within error (Table 2), indicating that the observed current is dominated by the catalytic reaction. Similar  $K_{\text{M}}^{\text{H}_2}$  values were obtained from linear fits of the high driving force part of the cyclic voltammograms at various  $\text{H}_2$  concentrations, further confirming prevalence of the catalytic reaction over interfacial electron transfer at high driving forces (Table S3†).

We further scrutinized the effect of temperature and pH on the catalytic activity of *holo-Tam* HydS under conditions when the catalytic current for  $\text{H}_2$  oxidation is not limited by mass transport (3000 rpm rotation speed and 1 atm  $\text{H}_2$ ). Fig. 8A and C displays CVs recorded at various temperatures (10–70 °C) at pH 7, and different pH values at 30 °C, respectively. Fig. 8B shows the temperature dependence of the CVs at high driving force (eqn (1)) for  $\text{H}^+$  reduction and  $\text{H}_2$  oxidation. Fig. 8D shows the

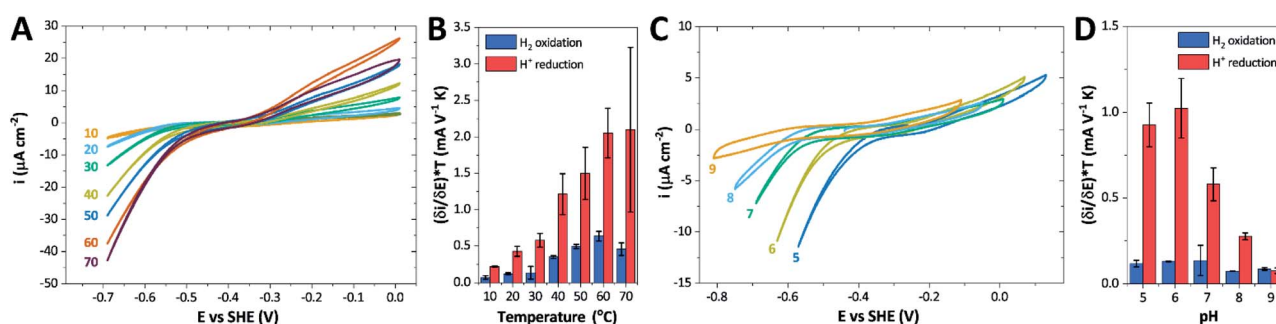


Fig. 8 CVs obtained at a rotating disc PGE modified with *Tam* HydS under 1 atm  $\text{H}_2$  at (A) various temperatures in the 10–70 °C range at pH 7 and (C) various pH values from 5 to 9 at 30 °C. The scan rate is  $2 \text{ mV s}^{-1}$ , the rotation rate is 3000 rpm. The data shown in (A) and (C) are obtained from single films cycled up and down in temperature and pH, respectively. Film stability was verified at the end of each experiment by returning the solution to its starting state (pH 7 solution at 30 °C). Dependence of the high driving force slopes of the voltammograms (eqn (1)) times the temperature for  $\text{H}_2$  oxidation and  $\text{H}^+$  reduction on (B) temperature and (D) pH. Error bars show standard deviation between three films.





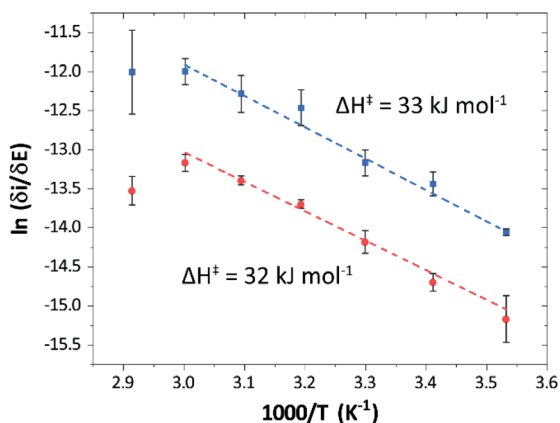
**Table 2**  $K_M^{H_2}$ -values for *Tam* HydS determined at various over-potentials and temperatures at pH 7

Over-potential (mV)	$K_M^{H_2}$ <sup>a</sup> (mM)	
	30 °C	60 °C
100	0.05 ± 0.01	0.15 ± 0.05
200	0.08 ± 0.02	0.12 ± 0.02
300	0.10 ± 0.02	0.13 ± 0.03
400	0.10 ± 0.03	0.15 ± 0.05
<b>Average</b>	<b>0.09 ± 0.03</b>	<b>0.14 ± 0.05</b>

<sup>a</sup> The errors for over-potentials 100–400 mV show standard deviation between three films.

corresponding data as a function of pH. When pH is decreased, the catalytic activity towards  $H^+$  reduction increases and the catalytic wave is shifted to more anodic potentials, consistent with higher  $H^+$  concentration (Fig. 8C). Conversely, the magnitude of the catalytic current for  $H_2$  oxidation does not vary smoothly with pH. The oxidation process is pH independent between pHs 8–9, increases at pH 6–7 and remains stable at this higher current down to pH 5. Moreover, over-potential for both  $H_2$  oxidation and  $H^+$  reduction is lowest at pH 5 (Fig. S18<sup>†</sup>). The origin of this pH switching behavior for  $H_2$  oxidation is currently not identified.

Temperature was found to have a strong influence on  $H_2$  oxidation and  $H^+$  reduction between 10–60 °C at pH 7 (Fig. 8A). Increasing the temperature not only resulted in higher overall currents, but also a significant decrease in over-potential. At room temperature, an over-potential of 50–60 mVs was observed in both catalytic directions, in contrast to previously characterized group A [FeFe]-hydrogenases. With the exception of specific mutants the latter enzymes generally display rapid increase in currents around the thermodynamic midpoint potential.<sup>44,65,66</sup> The over-potential requirement of *Tam* HydS is in line with the quasi-reversible nature of the  $H_{ox}/H_{red}$  transition observed by FTIR spectro-electrochemistry (Fig. S8<sup>†</sup>). At



**Fig. 9** Eyring plots for  $H^+$  reduction (red) and  $H_2$  oxidation (blue) with linear fits (dashed lines). Plots prepared using estimated  $i_{lim}$  based on data from Fig. 8B (data observed at 70 °C excluded from the linear fit).

**Table 3** Activation enthalpies ( $\Delta H^\ddagger$ ) for  $H^+$  reduction and  $H_2$  oxidation observed for different [FeFe]- and [NiFe]-hydrogenases

Enzyme	$\Delta H^\ddagger$ (kJ mol <sup>-1</sup> )	
	$H^+$ reduction	$H_2$ oxidation
<i>Tam</i> HydS (this work)	32 ± 3	33 ± 2
<i>CaI</i> <sup>a</sup>	29	19
<i>Cr</i> HydA1 <sup>a</sup>	20 <sup>b</sup>	26 <sup>b</sup>
<i>Ec</i> Hyd1 <sup>a,c</sup>	—	48
<i>Ec</i> Hyd2 <sup>a,d</sup>	65	37

<sup>a</sup> Experimental data obtained from ref. 62, determined at pH 6, 30 °C. <sup>b</sup> Only reached at high over-potential. <sup>c</sup> [NiFe]-hydrogenase *E. coli* Hyd1. <sup>d</sup> [NiFe]-hydrogenase *E. coli* Hyd2.

temperatures of 50–60 °C the over-potential decreased to approx. 10 mV. The decrease of catalytic currents at 70 °C is attributed to the deactivation of the protein, since we did not observe any significant protein loss during the experiment at temperatures up to 60 °C (Fig. S19<sup>†</sup>). It should also be noted that no oxidative inactivation<sup>67,68</sup> was observed when cycling up to ±0 mV vs. SHE. Stability and increased catalytic activity of the protein at elevated temperatures is not surprising considering the thermophilic nature of *T. mathranii*.<sup>69</sup> The activation enthalpies ( $\Delta H^\ddagger$ ) of the  $H_2$  oxidation and  $H^+$  reduction reactions were estimated through Eyring plots based on the change of the high potential slope as a function of temperature (Fig. 9), and found to be similar in both catalytic directions (Table 3). In the case of the prototypical [FeFe]-hydrogenases *Cr* HydA1 and *CaI* (Table 3),<sup>70</sup> distinctly lower activation enthalpies ( $\Delta H^\ddagger$ ) for either  $H^+$  reduction or  $H_2$  oxidation, respectively, have been reported, suggesting that *Tam* HydS is exceptionally well balanced for bidirectional catalysis. Moreover, albeit the activation enthalpies are higher for *Tam* HydS than *Cr* HydA1 and *CaI*, they are still significantly lower than what has been reported from the *E. coli* [NiFe]-hydrogenases *Ec* Hyd1 and *Ec* Hyd2 (Table 3). Thus, the low specific activity observed for *Tam* HydS (Fig. 2) cannot be explained by differences in activation enthalpies alone. Rather, the low catalytic rate of *Tam* HydS is governed by mass transfer, e.g. proton- or  $H_2$  transfer within the protein. Impaired proton transfer could also, at least partially, explain the over-potential observed at low temperature.<sup>44</sup> Finally, it is noteworthy that films prepared of the enzyme exposed to air, to induce the formation of the  $H_{air}$  state, displayed limited capacity for  $H^+$  reduction but a complete loss of  $H_2$  oxidation function (Fig. S20<sup>†</sup>). The catalytic properties of  $H_{air}$  was further supported by *in vitro* assays, showing a  $H^+$  reduction activity approximately hundred-fold lower than the native enzyme.

## Conclusions

This report represents the first biochemical and biophysical characterization of a group D [FeFe]-hydrogenase. As described herein, *Tam* HydS features a number of properties similar to regulatory [NiFe]-hydrogenases,<sup>71</sup> including a relatively low  $K_M^{H_2}$  and an increased tolerance against CO inhibition. This is in line with a potential sensory function, as expected from its close



relationship to the putatively sensory group C [FeFe]-hydrogenases. Still, a  $K_M^{H_2}$  of around 0.1 mM is 5–10 times higher than what has been reported for regulatory [NiFe]-hydrogenases,<sup>71,72</sup> but is arguably in agreement with the overall higher  $K_M^{H_2}$  and  $K_i^{H_2}$  reported for prototypical [FeFe]-hydrogenase.<sup>73</sup> Moreover, analysis on a genomic level shows that *Tam* HydS is encoded upstream of a heterotrimeric bifurcating group A [FeFe]-hydrogenase. Thus, we hypothesize that M2e represents an alternative type of sensory [FeFe]-hydrogenase. This raises the question of how signal transduction is achieved. In the case of group C putative sensory [FeFe]-hydrogenase, a PAS-domain is fused to the H-domain. Similarly, to the best of our knowledge, all regulatory [NiFe]-hydrogenases identified to-date feature the so-called HoxJ subunit, encoding a PAS domain-like sequence. A PAS-domain cannot be identified in the operon encoding for *Tam* HydS but the gene is flanked by a histidine kinase-like ATPase and an AraC family transcriptional regulator. These genes are commonly involved in signal transduction and regulation of transcription, which further suggests a regulatory role for the M2e hydrogenases.<sup>74,75</sup>

In the context of H-cluster function, *Tam* HydS displays a number of diverging properties. As reflected in the tolerance towards CO, the unusual mixture of  $H_{ox}$ -like states, reaction enthalpies ( $\Delta H^\ddagger$ ) that were practically identical for both  $H^+$  reduction and  $H_2$  oxidation, and a stabilization of the  $H_{red}$ -state. Moreover, it is noteworthy that despite its high tolerance towards CO *Tam* HydS reacted rapidly with  $O_2$  to form a stable mononuclear version of the  $[2Fe]_H$  subsite ( $H_{air}$ ). This argues against gas diffusion (mass transfer) to the active site as the main defense mechanism towards CO inhibition, which we instead attribute to the unusual hydrogen-bonding network around the H-cluster afforded by variations of *e.g.* Cys299, Met353 and Met497 (Fig. 1). Indeed, an increased resistance towards CO has also been reported for the putative-sensory group C [FeFe]-hydrogenase *Tm* HydS, featuring different alterations of the same residues.<sup>38</sup> The influence of the hydrogen-bonding network on CO affinity has also been suggested by studies of cofactor variants and mutants of *Cr* HydA1, in which the nearby cysteine was replaced by an alanine residue.<sup>51</sup>

The effect of  $O_2$  on prototypical group A [FeFe]-hydrogenases has been intensively studied, nevertheless formation of  $H_{air}$  has not been reported for any other [FeFe]-hydrogenase. Instead, the reaction generally causes complete H-cluster degradation or formation of reversibly inhibited states like  $H_{inact}$  by coordination of thiol ligands to the  $[2Fe]$  subsite.<sup>37,76</sup> Whether  $H_{air}$  has a physiological role remains uncertain. However, as this state appears unreactive towards  $H_2$  while retaining a limited  $H_2$  production capacity, it is unlikely to be relevant in  $H_2$  sensing. Still, its formation provides a striking example of how the active site environment modulates the reactivity of the H-cluster. It should be noted that formation of  $H_{air}$  is dependent on  $O_2$  as the equivalent state does not seem to be formed under oxidizing electrochemical conditions. As  $O_2$  acts as a chemical oxidant it can lead to an alternative pathway that remains to be fully elucidated. Arguably, access to electrons and protons has a large influence on the reaction between the H-cluster and  $O_2$  and

several factors need to be considered to explain the formation of  $H_{air}$  in *Tam* HydS.<sup>77,78</sup> It has been shown that in the case of [NiFe]-hydrogenase,  $O_2$  can be efficiently reduced to  $H_2O$  through rapid electron injection to the [NiFe] cofactor involving an unusual  $[4Fe-4S]$  cluster.<sup>79</sup> The presence of a C-terminal  $[4Fe-4S]$  cluster and a disrupted proton transfer pathway differentiates M2e enzymes from prototypical [FeFe]-hydrogenases. Still, formation of  $H_{air}$  was not reported in the case of *Tm* HydS which features the same differences.<sup>38</sup> This suggests that the reaction between the H-cluster and  $O_2$  is significantly influenced by other variations in the active site architecture.

We presume that the lack of a cysteine in position 299 also influences the catalytic activity of *Tam* HydS, as hindered proton release could stabilize  $H_{red}$  kinetically, following reduction of the H-cluster with  $H_2$ . A similar stabilization of  $H_{red}$  has been noted for *Tm* HydS.<sup>38</sup> However, while the latter enzyme is suggested to form an unusual, unprotonated, CO-bridged form of  $H_{red}$  ( $H_{red}^*$ ), *Tam* HydS appears to form an  $H_{red}$  state highly similar to that of prototypical [FeFe]-hydrogenase. Also, the complete lack of the well-conserved proton transfer pathway found in group A [FeFe]-hydrogenases raises the question of where an alternative pathway could reside. In the homology model of *Tam* HydS, a conserved glutamate residue close to the H-cluster (Glu252) is located in a preferable position for initiating proton transfer (Fig. S21†). However, there is no clear continuation of this potential pathway. In addition to changes in the proton transfer pathway, variation of Met353 and Met497 certainly also modulates the electronic properties of the H-cluster. It has been previously proposed that Met353 is important for tuning  $K_M^{H_2}$  as well as the catalytic bias of prototypical [FeFe]-hydrogenase, in favor of either  $H^+$  reduction or  $H_2$  oxidation.<sup>31,41</sup> Still, the exact contributions of the Met353Ser variation on the activity of *Tam* HydS remain to be fully elucidated.

In closing, this study shows that the *Tam* HydS M2e enzyme displays significant differences in reactivity as compared to previously studied group A as well as group C [FeFe]-hydrogenases. Thus, in addition to proposing a biological function for the group D [FeFe]-hydrogenases, it underscores how mutations of known [FeFe]-hydrogenases need to be complemented with further studies of the biodiversity to fully realize the chemical space of this fascinating family of enzymes. Mapping out the reactivity of these diverse enzymes is certainly critical for our understanding of hydrogen metabolism and envisioned biotechnological applications. It is also of high relevance in the context of bioinspired catalyst design, as it will provide new model systems for elucidating the influence of the protein environment and the outer coordination sphere on the reactivity of the H-cluster.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This project was supported by the Novo Nordisk Foundation (HL contract no. NNF19OC0055613), the European Research Council



under the European Union's Seventh Framework Programme (grant agreement No. 714102 to GB) and Horizon 2020 research and innovation programme (Marie Skłodowska-Curie grant agreement No. 897555 to MS), and the Deutsche Forschungsgemeinschaft through the priority program 1927 (grant agreement No. 1554/5-1 to STS). A. S. acknowledges support from the Knut and Alice Wallenberg Foundation (KAW 2015.0418).

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