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Accumulating the hydride state in the catalytic cycle of [FeFe]-hydrogenases

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H₂ turnover at the [FeFe]-hydrogenase cofactor (H-cluster) is assumed to follow a reversible heterolytic mechanism, first yielding a proton and a hydrido-species which again is double-oxidized to release another proton. Three of the four presumed catalytic intermediates (H_{ox}, H_{red}/H_{red} and H_{sred}) were characterized, using various spectroscopic techniques. However, in catalytically active enzyme, the state containing the hydrido-species, which is eponymous for the proposed heterolytic mechanism, has yet only been speculated about. We use different strategies to trap and spectroscopically characterize this transient hydride state (H_{hyd}) for three wild-type [FeFe]-hydrogenases. Applying a novel set-up for real-time attenuated total-reflection Fourier-transform infrared spectroscopy, we monitor compositional changes in the state-specific infrared signatures of [FeFe]-hydrogenases, varying buffer pH and gas composition. We selectively enrich the equilibrium concentration of H_{hyd}, applying Le Chatelier's principle by simultaneously increasing substrate and product concentrations (H₂/H⁺). Site-directed manipulation, targeting either the proton-transfer pathway or the adt ligand, significantly enhances H_{hyd} accumulation independent of pH.

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Hydrogen (H_2) turnover in [FeFe]-hydrogenases is proposed to occur at the distal iron site (Fe_d) of the $[2Fe_2S]$ moiety ($2Fe_H$), which is part of the H-cluster¹. The distal iron ion exchanges protons with a conserved proton-transfer pathway (PTP)^{2,3} via an azadithiolate ligand (adt) that bridges both iron sites (Fig. 1)^{4,5}. Different redox states of the H-cluster can be distinguished by the unique infrared vibrational signatures of the two CN^- and three CO ligands at the $2Fe_H$ moiety. To tailor [FeFe]-hydrogenases for specific applications or translate their catalytic principle into productive and durable synthetic mimics, detailed knowledge of the succession of turnover steps occurring at the H-cluster is required. The binuclear metal cofactors of [FeFe]- and [NiFe]-hydrogenases have been suggested to oxidize H_2 by enhancing the acidity of H_2 in the presence of a base, thus facilitating its heterolytic cleavage into a proton (H^+) and hydride (H^-) as a first intermediate step (Fig. 1)^{6–9}.

For [NiFe]-hydrogenases, electron paramagnetic resonance spectroscopy showed that a bridging metal hydride was present in the paramagnetic Ni-C state^{6,7}, while an ultra-high 0.89 Å resolution X-ray crystallography structure enabled the assignment of both, the proton and bridging hydride, in the first state following H_2 -binding (Ni-R)⁸. For algal-type [FeFe]-hydrogenases, a comparative density functional theory (DFT)/X-ray absorption spectroscopy analysis indicated a bridging hydride ($\mu-H^-$) in the double-reduced H_{sred} state¹⁰, albeit the importance of a $\mu-H^-$ state for fast catalytic turnover has been questioned. In fact, the terminal hydride ($t-H^-$) is thermodynamically less stable and more reactive compared to $\mu-H^-$ and would match the high turnover kinetics of [FeFe]-hydrogenases far better^{11–14}. According to the current working model for the catalytic mechanism of [FeFe]-hydrogenases¹, binding of H_2 to the oxidized active ready state (H_{ox}) results in the heterolytic cleavage of H_2 , with H_{hyd} as the first intermediate. On deprotonation, H_{hyd} is converted into the rather stable H_{sred} state before two successive oxidation steps recycle H_{ox} . A direct conversion of H_{hyd} into H_{red} has been proposed as well¹⁵. However, a terminal hydride has not been assigned to any of the known redox states¹⁰, nor has another catalytic wild-type state with the postulated $t-H^-$ been reported. This implies that the kinetically relevant hydride state is a transient one and difficult to trap under steady-state conditions. Owing to the rules governing steady-state kinetics, which favour thermodynamically stable intermediates in catalysis, the direct characterization of transient states relies on the utilization of time-resolved approaches with stopped-flow or single-turnover set-ups¹⁶.

We take a different approach, exploiting the properties of steady-state kinetics and using Le Chatelier's principle to

selectively enrich the transient $t-H^-$ state in three different [FeFe]-hydrogenases: HydA1; DdH; and CpI, which cover the complete range of increasingly complex structured monomeric hydrogenase subtypes (M1–M3)^{17–20}. In our experimental set-up, attenuated total reflection (ATR) Fourier-transform infrared (FTIR) spectroscopy is used to follow the population of different catalytic states. Compositional changes are monitored in the state-specific infrared signatures as a function of buffer/gas composition.

Results

Accumulation of H_{hyd} in wild-type [FeFe]-hydrogenases.

Figure 2a depicts FTIR difference spectra of HydA1 film at pH 8 recorded while changing the atmosphere from N_2 to H_2 (for details about the experimental setup see Supplementary Fig. 1). Negative bands mainly comprised the frequency pattern of H_{ox} ($1,964/1,940/1,802\text{ cm}^{-1}$). Positive bands included marker frequencies $1,915/1,891\text{ cm}^{-1}$ (H_{red}) and $1,953/1,918/1,882\text{ cm}^{-1}$ (H_{sred}), corresponding to earlier measurements with H_2 -purged HydA1 carried out in transmission FTIR systems²¹. On steady-state supply of substrate (that is, H_2), the enzyme accumulates the apparently rather stable reduced intermediates H_{red} and H_{sred} . However, bands at $1,978/1,960/1,860\text{ cm}^{-1}$ could be identified as minor contributions, indicating the presence of another state, which resembles the band pattern of H_{trans} in DdH (H_{trans} -like state)²² and was previously observed as a weak fraction of a complex band pattern in a sodium dithionite (NaDT)-treated HydA1 sample²³. According to our working model, deprotonation should lead from the presupposed H_{hyd} to the reduced states (Fig. 2b), and a plausible way to selectively enrich H_{hyd} would be a simultaneous increase in substrate (H_2) and product (H^+) pressure (Fig. 2d). We therefore titrated HydA1 to pH 4 before H_2 exposure, enhancing the proton pressure by four orders of magnitude (Fig. 2c). The presence of a band at $1,891\text{ cm}^{-1}$ in the resulting difference spectrum hinted at a minor contribution of H_{red} , while no H_{sred} could be detected. Instead, the H_{trans} -like state represented the main fraction of our protein sample. No loss of cofactor was observed in the time frame of hours (Supplementary Fig. 2a,b), and the back-titration of HydA1 sample buffer from pH 4 to pH 8 clearly demonstrates a pH-dependent reversibility of the indicated state transitions (Supplementary Fig. 2b,c). The same applies for H_2 -dependency at pH 4 (Supplementary Fig. 2d). This bi-directionality is not compatible with the supposed H_{trans} -like nature of the novel pattern²². Therefore, we will refer to this redox state in the following as H_{hyd} . Similar results were obtained when repeating this experiment with samples of native DdH and CpI as representatives of the other two major [FeFe]-hydrogenase subtypes, implying this state and behaviour to be general features of the [FeFe]-hydrogenase family (Supplementary Fig. 3).

The influence of proton-transfer efficiency. As the H_2/H^+ balance appears to affect the electronic configuration of the H-cluster, alternative approaches were designed to enrich H_{hyd} . HydA1 sample was once more purged with H_2 aerosol to first enrich the reduced states. When humidity was removed from the H_2 gas stream, sample films reacted by rapid dehydration. As predicted, this as well leads to a loss of H_{red}/H_{sred} in favour of the $1,978/1,960/1,860\text{ cm}^{-1}$ pattern (Supplementary Fig. 4a) indicating that the loss of bulk water has a similar effect to lowering the buffer pH, likely due to subsiding proton-transfer efficiency²⁴ (Supplementary Fig. 4).

To verify that an interference with the PTP enhances the population of the H_{hyd} state under H_2 exposure, we blocked the PTP in HydA1 and CpI via site-directed mutagenesis at two strictly conserved positions for each protein. Position 1 (C169 of

Figure 1 | Substrate/product transfer and heterolytic H_2 splitting in [FeFe]-hydrogenases. The electron-transfer path is shown in yellow, the PTP is depicted in blue. H_2 is released from, or reaches the $2Fe_H$ cluster through hydrophobic gas channels (green). At the catalytically pivotal $2Fe_H$ site, the presumptive first step in the catalysis of H_2 oxidation (H_2 heterolysis) is depicted, resulting in the unequal intermediates H^+ , binding at the adt-bridge (Lewis base) and H^- , binding at Fe_d (Lewis acid).

Figure 2 | Population of H_{hyd} in HydA1 by simultaneously increasing H_2 and H^+ pressure. (a) ATR FTIR difference spectrum ($N_2/pH8 \rightarrow H_2/pH8$), depicting the accumulation of the reduced states (H_{red}/H_{red} and H_{sred}) in the catalytic cycle of HydA1 under H_2 at pH 8. (c) ATR FTIR difference spectrum ($N_2/pH8 \rightarrow H_2/pH4$) depicting the accumulation of H_{hyd} in the catalytic cycle of HydA1 under H_2 at pH 4. Catalytic cycles (b,d) beside the corresponding spectra illustrate the shift in steady-state equilibrium depending on applied substrate and product pressures in the working model of the catalytic cycle, which comprises the already-characterized states H_{ox} , H_{red}/H_{red} and H_{sred} , and the missing H_2 -activated state (H_{hyd}). Peak labels for H_{ox} are presented in black, for H_{hyd} in blue, for H_{red} in green and for H_{sred} in magenta (red asterisks: H_{red}). For the complete state-specific CN^- -/ CO -vibrational spectra of HydA1 observed during sample analysis, see Supplementary Table 1. At pH 4, the vibrational signals of H_{ox} were slightly shifted to higher frequencies (for more information see Supplementary Table 1, Hox-blue).

HydA1/C299 of CpI; Fig. 1), which was shown in earlier experimental data to be involved in catalytic proton transfer^{3,5}, marks the most proximal position of the PTP to the $2Fe_H$ sub-cluster, while position 2 (E141 of HydA1/E279 of CpI; Fig. 1) is situated more distant from the H-cluster. An exchange to alanine almost completely abolished H_2 -release activity for all variants (Supplementary Fig. 6). To probe the structural integrity and unequivocally connect the monitored effects to an interrupted PTP, we solved the crystal structure of CpI E279A (Fig. 3). The X-ray structure illustrates a fully intact H-cluster and overall identical structural features compared to wild-type CpI (refs 19,25), with the exception of the clearly diminished electron density at position 279, which is in full accordance with an exchange from glutamate to alanine. As a consequence of this non-conservative exchange, the path of polar side chains is interrupted between S319 and the harnessed H_2O molecule. Figure 4a,b shows the reaction of enzyme variants C169A of HydA1 and E279A of CpI towards H_2 (black graph) at pH 8. Similar to the effect of acidification or sample drying, the oxidized state is lost in favour of bands at $1,978/1,962/1,863\text{ cm}^{-1}$ or $1,985/1,970/1,858\text{ cm}^{-1}$, respectively.

We further exploited the effect of H_{hyd} accumulation as a consequence of PTP obstruction to elucidate the proposed role of the azadithiolate ligand in the heterolytic cleavage of H_2 . The earlier-described procedure of *in vitro* maturation allows for the replacement of the native $2Fe_H$ site with derivatives such as odt- $2Fe_H$, which contains an oxodithiolate group ($((SCH_2)_2O, odt)$ instead of the native azadithiolate ($((SCH_2)_2NH, adt)$ ^{4,25,26}. After *in vitro* maturation with odt- $2Fe_H$ in presence of H_2 , HydA1^{odt} exhibits an infrared band pattern that strongly resembles H_{hyd} (refs 23,26). The oxidized state of HydA1^{odt} was obtained via extensive purging with humidified N_2 . Subsequent exposure to H_2 induced a decrease in the H_{ox} pattern in favour of bands $1,980, 1,962$ and $1,868\text{ cm}^{-1}$ (Fig. 4c, black graph), however at a significantly lower rate compared to the corresponding experiment with wild-type HydA1.

Figure 3 | Crystal structure of the PTP in wild-type CpI and PTP-defective CpI variant E279A. Stick models and simulated annealing omit $[Fe-Fc]$ electron density maps (blue) of the $2Fe_H$ cluster, and the amino-acid residues and the H_2O molecules involved in proton transfer embedded in the $2Fe-Fc$ electron density maps (grey) of the X-ray structures of CpI wild-type²⁵ (a) and variant E279A (b). The presumed H^+ -transfer pathway is indicated in yellow. Crystal structure data for wild-type CpI correspond to pdb database entry 4XDC²⁵. Details of the crystallographic data of E279A are summarized in Supplementary Table 2. For stereo view presentations see Supplementary Fig. 9.

Figure 4 | Manipulating H^+ transfer traps the H_{hyd} state under H_2 or D_2 regardless of buffer pH. Left: ATR FTIR difference spectra ($N_2/pH8 \rightarrow H_2$ or $D_2/pH8$) depicting the accumulation of H_{hyd} for variants of the PTP E279A (Cpl) (a) and C169A (HydA1) (b), as well as HydA1^{odt} (c) under H_2 or D_2 at pH 8. The grey spectrum in a depicts the absolute absorption bands of H_{hyd} . Negative bands are assigned to H_{ox} (Supplementary Table 1). Right: location of the corresponding PTP manipulation illustrated in a crystal structure model of Cpl (pdb ID: 4XDC). Accumulation of H_{hyd} under D_2 (red spectra) does not affect the tCO-frequencies but induces a selective redshift for the μCO signal of $6\text{--}8\text{ cm}^{-1}$ in all analysed variants. For the complete state-specific CN[−]/CO-vibrational spectra observed during sample analysis, see Supplementary Table 1.

A terminal hydride in wild-type [FeFe]-hydrogenases. To confirm the presence of a terminal hydride ($t\text{-H}^-$) in H_{hyd} for wild-type hydrogenase and all examined variants, we probed the effect of an H/D isotope exchange on the FTIR band position of the bridging CO ligand, μCO (see Fig. 4, red graph, and Supplementary Figs 7 and 8 including Supplementary Note 1). Terminal hydride species on metal carbonyl complexes are vibrationally coupled with the ligand positioned in *trans* due to resonant frequencies²⁷. Previously published DFT calculations on the H-cluster model of HydA1 variant C169S predict a selective ‘redshift’ of 6 cm^{-1} for the μCO frequency as a consequence of substituting a hydride for a deuteride species located *trans* to μCO (ref. 23). When switching the purging gas from H_2 to D_2 , Cpl E279A (Fig. 4a) and HydA1 C169A (Fig. 4b) showed a selective shift to lower energies by $5\text{--}6\text{ cm}^{-1}$ of the band at $1,860\text{ cm}^{-1}$, which corresponds to the stretching vibration of μCO . A similar ‘redshift’ was observed for HydA1^{odt} when the oxidized enzyme was extensively treated with D_2 (Fig. 4c and Supplementary Fig. 7b). In Supplementary Fig. 8 we show spectra that document the H/D effect on wild-type HydA1 in presence of pH 4 and H_2/D_2 . This confirms the corresponding behaviour of the H_{hyd} state for all protein variants examined here.

Discussion

Le Chatelier’s principle was applied to enrich the highly transient H-cluster intermediate H_{hyd} , which according to the current

working model of catalytic H_2 turnover is predicted to carry a terminal hydride species. By simultaneously enhancing substrate and product pressure, we accumulated H_{hyd} for three different wild-type [FeFe]-hydrogenases. Interestingly, a nearly identical FTIR band pattern was observed in our earlier spectroscopic analysis of the largely inactive HydA1 PTP variant C169S (ref. 28). Using DFT calculations, Mulder *et al.*^{15,23} assigned this band pattern and redox species to a model of C169S that carried a terminal hydride at Fe₄ (Supplementary Note 1 and Supplementary Fig. 5).

Accumulation of H_{hyd} was also observed on decreasing the humidity level of the protein sample. Recently, we demonstrated that freeze-drying renders [FeFe]-hydrogenase samples insensitive to O_2 exposure²⁴. Although X-ray absorption spectroscopy data showed the formation of a stable O_2 adduct at the $2Fe_H$ site, H-cluster degradation was not initiated, as opposed to hydrated enzyme in presence of O_2 (ref. 29). DFT calculations suggested a protonation step mandatory for generating destructive reactive oxygen species³⁰. Therefore, a lack of H^+ -transfer activity in the freeze-dried state appeared to be a plausible explanation for this effect and dehydration should have a similar effect as lowering the pH. It can be speculated that the loss of bulk water, being the acceptor pool for exported protons, leads to an accumulation of protons in the PTP from the inside. Accordingly, low pH and dehydration both provide a counter pressure that prevents the H_2 -activated state from being deprotonated and proceeding to the reduced states (Supplementary Fig. 4b,c).

The N_2/H_2 difference spectra of site-directed exchange variants of HydA1 and Cpl targeting positions C169/C299 and E141/E279 clearly demonstrate how the disrupted proton transport stabilizes the H_{hyd} state regardless of buffer pH. This highlights that the transient character of H_{hyd} essentially depends on a functional catalytic proton transfer, just as predicted from the working model (Figs 1 and 2). In return, the analogous behaviour of cysteine and glutamate variants provides experimental proof for the involvement of the median glutamic acid residue (E141^{HydA1}/E279^{Cpl}) in the proton-transfer mechanism of [FeFe]-hydrogenases. In future mutagenesis studies on the PTP of [FeFe]-hydrogenases, accumulation of H_{hyd} can be regarded as an experimental indicator for the involvement of peptide positions in the catalytic H^+ transfer.

The relevance of the bridgehead position of the $2Fe_H$ subcluster for catalytic proton transfer was clearly demonstrated by the fact that cofactor variant HydA1^{odt} could be trapped in H_{hyd} under H_2 independent of the pH, thus confirming the results of earlier studies, which pointed out the essential role of the native adt ligand as a proton relay to the substrate-binding site at Fe_d . The fact that H_2 exposure of HydA1^{odt} results in H_{hyd} accumulation suggests that the ether bridgehead of odt is capable to serve as a Lewis base and supports heterolytic H_2 cleavage. This reflects earlier investigations of Barton *et al.*,³¹ which demonstrated in a comparative study, including different diiron complexes that the weakly basic ether group of odt can assist in proton relay during H_2 evolution³¹. However, in contrast to C169A and E279A, the H_{hyd} spectrum of HydA1^{odt} could only be regained very slowly in presence of H_2 . While all examined enzyme variants exhibited an impaired proton transfer, for C169A and E279A the defect was restricted to the H^+ transfer across the protein shell (Fig. 4). In case of HydA1^{odt}, the impairment already affected proton abstraction during H_2 heterolysis. This finding experimentally verifies the hypothesis that the pending amine group in the native adt ligand significantly enhances the rate of heterolytic H_2 cleavage³¹. The selective shift of the μCO band observed in H_{hyd} on H/D exchange suggests the presence of a terminal hydride ligand for all PTP variants and catalytically competent wild-type enzymes examined here.

In addition, two recently published studies independently demonstrated a terminal hydride for inactive HydA1 variants assigned to the presumptive H_{hyd} state. Reijerse *et al.*³² reported the direct detection of Fe_d-H bending vibrations via nuclear resonance vibrational spectroscopy of a terminal hydride species in HydA1^{odt} after isolation under H_2 . This is in good agreement with our real-time analysis of HydA1^{odt}. Furthermore, Mulder *et al.*¹⁵ applied electron paramagnetic resonance and Mössbauer spectroscopy to provide experimental support for the presence of a metal hydride species in HydA1 variant C169S, which largely adopted H_{hyd} on supplementation with NaDT. Well in agreement with electrochemistry studies on HydA1 under turnover conditions¹⁴, the authors determined a H_{hyd} transition potential close to the H_2/H^+ redox couple. This argues in favour of the relevance of H_{hyd} in the catalytic cycle.

Both HydA1 C169S and HydA1^{odt} have been shown to be largely inactive^{4,15,23,25,26,28,32}, and a definite proof for the catalytic relevance of H_{hyd} can only be given by monitoring the succession of states during catalysis at sub-turnover time resolution as previously done for [NiFe]-hydrogenase¹⁶. However, this work strongly suggests that H_{hyd} , as identified by Mössbauer and nuclear resonance vibrational spectroscopy, corresponds to the first transient state in the catalytic mechanism of H_2 oxidation and is shared by all major subtypes of [FeFe]-hydrogenases. It is therefore plausible to assume that we successfully accumulated a state carrying a terminal hydride ligand in native [FeFe]-hydrogenases. In response to modulating

substrate and product concentrations (H^+/H_2), accumulation of H_{hyd} was found to be fully reversible showing that H_{hyd} is no artificial dead-end state but part of the dynamic redox equilibrium typically occurring in the catalytically competent enzyme. All available data strongly suggest that H_{hyd} is indeed the missing link in the catalytic mechanism of [FeFe]-hydrogenases.

Methods

Site-directed mutagenesis of HydA1 and Cpl. Plasmids pET21b-HydA1Cr and pET21b-Cpl were used as templates for site-directed mutagenesis, which was performed according to the QuikChange-PCR protocol published by Zheng *et al.*³³ For generating the constructs encoding HydA1 variants C169A, C169S and E141A, the 5'-overlapping mismatch primer pairs 5'-CAGCGCGTGTCCGGGCTGG ATTGC-3'/5'-CCGGACACGCGCTGGTAAACATCGG-3', 5'-CAGTTCATG TCCGGGCTGGATTGC-3'/5'-CCGGACATGAGCTGGTAAACATC-GG-3' and 5'-CATCATGGAAGCGGGCAGCGAACTGTCATCGTC-3'/5'-GCAGTTC-GCTGCCCGCTCCATGATGGTCAGATCC-3' were used, respectively.

For substitutions C299A and E279A within the polypeptide of Cpl, primer pairs 5'-CCTCTGCGTGCCAGGTTGGGTACGTC-3'/5'-CCTGGGCACGCAGAGG TAAACATT-GGGAAAGGG-3' and 5'-CCAGTTCGGTAGCCGCTCCATAA TGGTCATATC-3'/5'-TG-GAAGCGGCTACCGAACTGGTCAACG were utilized.

Heterologous expression of HydA1 and Cpl proteins. The original plasmids pET21b-HydA1Cr and pET21b-Cpl and their mutagenesis constructs were used for the heterologous expression of wild-type enzymes and variants of HydA1 and Cpl (refs 4,25,34,35). Apo-forms of Cpl and HydA1 (lacking $2Fe_H$) were expressed anaerobically in *Escherichia coli* strain BL21 ΔiscR (ref. 36). Cells first grew aerobically in lysogeny broth (LB) medium pH 7.4 (0.1 M morpholinepropanesulfonic acid (Mops)-NaOH) supplemented with 5 g l⁻¹ glucose and 2 mM ammonium iron citrate until an OD₆₀₀ of 0.35–0.6 was reached. Before switching to anaerobic cultivation, the culture was flush with N_2 for 30 min to remove residual O_2 and supplemented with 25 mM sodium fumarate. Induction of gene expression was initiated by adding 5 mM cysteine and 0.5 mM β-D-1-thiogalactopyranoside. Cells were collected after 16–24 h expression via centrifugation.

Purification of [FeFe]-hydrogenase apo-proteins. The apo-proteins of wild-type and mutant forms of HydA1 and Cpl, containing the $4Fe_H$ subcluster but lacking the $2Fe_H$ moiety, were expressed under anaerobic conditions in *E. coli* strain BL21 ΔiscR as described earlier^{4,34,36}. After cell disruption via ultrasonication and the separation of the soluble proteins from the cell debris by ultracentrifugation and subsequent filtering (pore size 0.2 μm), we exploited the C-terminally fused Strep-tag-II sequence (WSHPQFEK) for protein purification, performing affinity chromatography with Strep-Tactin Superflow resin (IBA GmbH) in 100 mM Tris-HCl buffer (pH 8) supplemented with 2 mM NaDT. Protein purity was verified by SDS-PAGE and protein concentration determined via Bradford assay (BioRad).

Synthesis of $2Fe_H$ cofactor complexes. The native-like $2Fe_H$ -cofactor-mimic complex $[Fe_2\{\mu-(SCH_2)_2NH\}(CN)_2(CO)_4][Et_4N]_2$ (adt- $2Fe_H$) and its odt derivative $[Fe_2\{\mu-(SCH_2)_2O\}(CN)_2(CO)_4][Et_4N]_2$ (odt- $2Fe_H$) were synthesized according to literature procedures²⁵.

In vitro maturation. To obtain the fully equipped holoprotein, heterologously expressed wild-type and mutant apo-proteins of HydA1 and Cpl were matured *in vitro* by adding adt- $2Fe_H$ (odt- $2Fe_H$) to a 10-fold excess in 0.1 M K_2HPO_4 /KH₂PO₄ buffer (pH 6.8), supplemented with 2 mM NaDT. To ensure complete sample maturation, the reaction mix was incubated for 1 h at room temperature. The resulting holoproteins were cleaned from surplus $2Fe_H$ complex by size-exclusion chromatography using NAP 5 columns of GE Healthcare and concentrated in Amicon Ultra centrifugal filters 30K (Millipore). Afterwards, methyl viologen-specific H_2 production activity was determined to assess catalytic competence.

In vitro assay to determine H_2 production activity. H_2 production activities have been determined *in vitro* for wild-type and mutant proteins using 0.2 μg ml⁻¹ of *in vitro* matured [FeFe]-hydrogenase in 2 ml of 100 mM postassium phosphate buffer, pH 6.8, supplemented with 100 mM NaDT and 10 mM methyl viologen. Suba-seal vessels were sealed with stoppers, degassed for 5 min with 100% Argon and incubated for 20 min in a shaking water bath adjusted to 37 °C. For product quantification, 400 μl of sample head-space was analysed via gas chromatography (Shimadzu GC 2010). When enzyme activity was below the detection limit, the assay was repeated with a 10-fold increased enzyme concentration (2 μg ml⁻¹).

Crystallization and structure analysis of Cpl variant E279A. Using a 1:1 mix of E279A holoprotein (10 mg ml⁻¹) and reservoir solution (0.1 M MES, pH 6,

supplemented with 0.4 M MgCl₂, 24% polyethylene glycol and 16% glycerol) in a hanging drop vapour diffusion set-up, single cuboid-shaped crystals of brownish colour were identified after 2–4 days at 277 K under anaerobic conditions. The selected protein crystal was mounted into a CryoLoop (Hampton Research) and flash-frozen in liquid N₂. Diffraction data were collected at 100 K at beamline ID23-2 of the ESRF in Grenoble, France and processed using the software package XDS³⁷. Molecular replacement and structure optimization were achieved with the software packages PHENIX³⁸ and Coot³⁹. For details on crystallographic data of Cpl E279A, see Supplementary Table 2.

Infrared spectroscopy. All FTIR spectroscopy was conducted on a rapid-scan Tensor 27 spectrometer (Bruker Optik, Germany) equipped with a three-reflection ZnSe/silicon crystal ATR cell (Smith Detection, USA). The spectrometer was situated in an anaerobic gas chamber (Coy Laboratories, USA) in a water-free atmosphere of typically 99% N₂ and 1% H₂. Supplementary Fig. 1 depicts the process flow diagram of the experimental set-up. Approximately 5 bar nitrogen carrier gas of ultra-high purity (5.0) was provided by a PN1450 nitrogen generator (Inmatec, Germany). The exact amount of N₂ was adjusted with a digital Smart-Trak mass flow controller (MFC, Sierra, USA). Molecular H₂ (H₂ 5.0, Linde, Germany) was injected separately via a second flow controller to create a well-defined mixture. The gas passed a 200 mbar check valve to protect the flow controllers from humidity. Afterwards, two sequential wash bottles (H₂O and miscellaneous content, for example, buffer) could be switched into the gas stream. A bypass loop allowed for an adjustable proportion of the gas to bypass the wash bottles entirely or to run through the liquid volume to create a mix of carrier gas, water vapour and microscopic drops of liquid water (aerosol). The aerosol stream was fed to a customized polychlorotrifluoroethylene (PCTFE) gas cell, screwed gas-tight onto the ATR crystal plate. An exit line eventually guided the aerosol to a gas dump (fume). The PCTFE gas cell was equipped with three inlets and a manometer (not shown in Supplementary Fig. 1). Directly over the silicon crystal of the ATR unit, a cylindrical cavity joined all inlets. A gas-tight lid was screwed on top of the gas cell and allowed irradiation via an acrylic glass window. At this position, an infrared- or ultraviolet/visible-transparent window with protein sample could be adjusted into the aerosol stream. Thus, transmission samples for, for example, ultraviolet/visible or Raman spectroscopy could be prepared under the same conditions as monitored via the ATR silicon crystal. All spectra were recorded with 80 kHz scanning velocity, at a spectral resolution of 1 cm⁻¹, and varying extent of co-additions. All pH titrations were performed with citrate buffer (SSC).

Data availability. The atomic coordinates and factors for the reported crystal structure for Cpl variant E279A have been deposited with the Protein Data Bank (PDB) under accession code 5LA3. Further data supporting the findings of this study are available within the article and its Supplementary Information file and from the corresponding authors on reasonable request.

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Author contributions

M.W., S.T.S. and T.H. designed research; M.W., S.T.S. and T.H. wrote the manuscript; U.-P.A. and F.W. synthesized the 2Fe_H -adt/-odt complexes for *in vitro* maturation; J.D. and M.W. produced, purified and matured the wild-type and mutant proteins of HydA1 and Cpl; S.T.S. and M.S. performed and designed ATR FTIR experiments; J.D., E.H. and J.E. performed or contributed to protein crystallization and X-ray structure analysis.

Additional information

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