Caught in the H_{inact}: Crystal Structure and Spectroscopy Reveal a Sulfur Bound to the Active Site of an O_{2}-stable State of [FeFe] Hydrogenase

Patricia Rodríguez-Maciá, Lisa M. Galle, Ragnar Björnsson, Christian Lorent, Ingo Zebger, Yoshitaka Yoda, Stephen P. Cramer, Serena DeBeer, Ingrid Span, and James A. Birrell

Abstract: [FeFe] hydrogenases are the most active H_{2} converting catalysts in nature, but their extreme oxygen sensitivity limits their use in technological applications. The [FeFe] hydrogenases from sulfate reducing bacteria can be purified in an O_{2}-stable state called H_{inact}. To date, the structure and mechanism of formation of H_{inact} remain unknown. Our 1.65 Å crystal structure of this state reveals a sulfur ligand bound to the open coordination site. Furthermore, in-depth spectroscopic characterization by X-ray absorption spectroscopy (XAS), nuclear resonance vibrational spectroscopy (NRVS), resonance Raman (RR) spectroscopy and infrared (IR) spectroscopy, together with hybrid quantum mechanical and molecular mechanical (QM/MM) calculations, provide detailed chemical insight into the H_{inact} state and its mechanism of formation. This may facilitate the design of O_{2}-stable hydrogenases and molecular catalysts.

Introduction

Hydrogen is a promising green energy carrier for the future because it can easily be produced by water electrolysis using renewable energy and, later used in a fuel cell to generate energy producing only water as a byproduct.[1] Currently, H_{2} is produced mostly from fossil fuels or to a small extent by water electrolysis using expensive noble metal catalysts. In nature, efficient and reversible H_{2} conversion is performed by a group of metalloenzymes called hydrogenases.[2] These biocatalysts use earth abundant metals such as nickel and/or iron in their active site.[3] Of the three groups of hydrogenases ([NiFe] hydrogenases, [FeFe] hydrogenases and [Fe] hydrogenases), the [FeFe] hydrogenases are the most active (10 000 s^{-1} in H_{2} oxidation and up to 10 000 s^{-1} in H^{+} reduction).[4] However, these enzymes are extremely oxygen sensitive.[5] Vigorous efforts have been made in order to protect [FeFe] hydrogenases, and hydrogenases in general, from oxygen. [6] Although various oxygen inactivation mechanisms have been proposed,[7] there is still a lack of understanding on how exactly O_{2} attacks their active site. Such insights may help in designing strategies to protect hydrogenases and molecular catalysts from O_{2} damage.

The active site of the [FeFe] hydrogenases, the H-cluster, consists of a binuclear [2Fe] sub-cluster ([2Fe]_{H}) covalently attached by a cysteine sulfur to a [4Fe–4S] cluster ([4Fe–4S]_{H}).[8] [2Fe]_{H} contains two irons bridged by the thiol groups of an aza-propane 1,3-dithiolate (ADT) ligand,[9] a bridging CO ligand, with an additional CN– and CO ligated to each iron. The (proximal) iron (Fe_{p}) directly bound to the [4Fe–4S]_{H} sub-cluster is always coordinatively saturated, while the distal iron (Fe_{d}) possesses an open coordination site in most catalytic states, where substrates (H_{2} and H^{+}) and inhibitors (including CO and O_{2}) can bind. The nitrogen atom in the ADT bridge serves as a base and Fe_{d} acts as a Lewis acid, together forming a frustrated Lewis pair, which is essential to heterolytically split H_{2} at Fe_{d}.[10] The catalytic cycle of these enzymes has been extensively studied through different spectroscopic techniques.[10]

When purified aerobiically from the native organism, the [FeFe] hydrogenase from Desulfovibrio desulfuricans remains...
in an inactive oxygen-stable state called $H_{\text{inact}}$ (or $H_{\text{ox}}^{\text{air}}$), which can be reactivated upon reduction.[4,11] This state is thought to be “overoxidized” with an Fe$^{III}$Fe$^{II}$ configuration at the binuclear site and an additional ligand bound to Fe$_{d}$.[11] The reduction of $H_{\text{inact}}$ to an intermediate state $H_{\text{trans}}$ is reversible while the further conversion of $H_{\text{trans}}$ to $H_{\text{ox}}$ is thought to be irreversible, involving the release of the putative ligand from Fe$_{d}$.[11] The nature of this putative ligand in the $H_{\text{inact}}$ state has remained a mystery for more than two decades. Despite considerable spectroscopic analysis,[12] new approaches are clearly needed to define the electronic and geometric configuration of the $H$-cluster, and identify the nature of the exogenous ligand. Theoretical calculations have suggested that the extra ligand could be H$_2$O or OH$^-$. [13]

Interestingly, an [FeFe] hydrogenase from Clostridium beijerinckii has been shown to convert into the $H_{\text{inact}}$ state in a highly reversible fashion, but the presence of an extra ligand bound in this state is so far unknown and its formation mechanism remains elusive.[14]

Recently, we showed that the $H_{\text{inact}}$ state is formed upon oxidation of $Dd$HydAB in the presence of sulfide (Na$_2$S). Based on this result, we suggested that the extra ligand bound to the open coordination site might be a sulfur species, possibly SH$^-$. [15] However, we were unable to identify whether sulfide was directly bound to the $H$-cluster, in what configuration, and whether there were any additional changes to the enzyme during $H_{\text{inact}}$ formation. In this work, we identify the nature of the additional ligand as SH$^-$ through combined crystallographic and spectroscopic investigations. These results together with hybrid QM/MM calculations provide deeper understanding on the formation mechanism of this state and how it is protected against O$_2$. This new insight may allow the general protection of metalloenzymes against oxygen, enabling their implementation in fuel cells and ultimately, it may provide design principles for developing O$_2$-stable bio-inspired molecular catalysts.

**Results and Discussion**

**Crystal Structure of $Dd$HydAB in the $H_{\text{inact}}$ State**

$Dd$HydAB in the $H_{\text{inact}}$ state was crystallized under aerobic conditions at 12°C. Brown crystals (indicating the presence of iron–sulfur clusters) were observed within three days and retained their dark color for at least two weeks. IR spectra of crystals taken from the same drop confirmed that the $Dd$HydAB was in the $H_{\text{inact}}$ state (Figure S1 in the Supporting Information). $Dd$HydAB in the $H_{\text{inact}}$ state crystallized in an orthorhombic space group $P2_12_12_1$, and the asymmetric unit contains one biological assembly. In contrast, the previously reported structure was obtained from crystals with the space group $P2_12_12_1$ and the asymmetric unit contained two biological assemblies. The structure of $H_{\text{inact}}$ was solved using molecular replacement with the structure published by Nicolet et al.[8b] (PDB ID 1HFE) as a starting model, and was refined to a resolution of 1.65 Å (crystal parameters and refinement statistics in Table S1). The structure by Nicolet et al. is the only available structure of $Dd$HydAB and the redox state of the enzyme in these crystals was not defined but assumed to be a mixture of the $H_{\text{ox}}$ and $H_{\text{red}}$ states.

The overall architecture of $Dd$HydAB in the $H_{\text{inact}}$ state is essentially identical to the starting model with a root mean square deviation (RMSD) of 0.631 Å (calculated for all Ca atoms of residues 2–397 without outlier rejection, Figure 1A). The electron density for the $H$-cluster in the active site is well-defined (Figure 2A); however, the occupancy of the [2Fe] sub-cluster had to be reduced to 0.6 to fit the experimental data. The low [2Fe] content indicates the presence of some apo protein in the preparation, partly a limitation of the artificial maturation procedure (see methods in the Supporting Information),[16] and partly from some decomposition of the $H$-cluster.[15] A more detailed analysis of the atomic
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No significant oxidative damage during crystallization or radiation damage to the accessory iron–sulfur clusters during the measurement were observed (Figure S3).

The electron density map reveals that the H-cluster is intact and contains two CN⁻ ligands, two terminal CO ligands, the bridging ADT ligand, as well as the bridging CO ligand (Figure 2 A). In addition, the electron density in the active site clearly shows the presence of an additional ligand in the apical position on Fe₄ at a distance of 2.4 Å (see Figure S4). The ligand consists of only one non-hydrogen atom, in agreement with the presence of a (hydro)sulfide, hydroxide or oxo ligand. Modelling of the sulfur ligand with the same occupancy as that of the [2Fe] subcluster (0.6) resulted in a good fit, but with a high B-factor (45 Å²), indicating some intrinsic disorder of the exogenous ligand. Modelling with an oxygen ligand gave a similar occupancy, but a slightly lower B-factor (38 Å²). Thus, anomalous scattering on exactly the same crystal was utilized to provide further information about the nature of the additional ligand. By measuring diffraction data at 6 keV, the anomalous signal from the iron atoms is suppressed, while that from other heavy atoms (such as S or Cl) is enhanced. The resulting anomalous electron density map shows clear evidence for anomalous density at the apical position on Fe₄ (Figure 2 B and Figure S5 A and B). To further support this observation, anomalous diffraction measurements were also performed before native data collection on a second crystal obtained under identical conditions (see Supporting Information, Table S1). The anomalous density map of the second crystal also showed distinct anomalous density at the apical position on Fe₄. While this strongly supports that the additional ligand is actually a sulfur species, we cannot exclude the possibility of a Cl⁻ ligand. Interestingly, Cl⁻ has been suggested to bind to the H-cluster under certain circumstances.[19] Hence, we investigated the spectroscopic properties of the H₅₆ₑｔ state to provide further insight.

Characterization of the H₅₆ₑᵗ State by X-Ray Absorption Spectroscopy

X-ray absorption spectroscopy (XAS) on the H₅₆ₑᵗ state (containing the additional ligand) and the well-characterized H₅₆ₑᵗ state (lacking the additional ligand) were measured for comparison. Figure 3 shows the Fourier-transformed (FT) spectrum of the extended X-ray absorption fine structure (EXAFS) region for H₅₆ₑᵗ (Figure 3 A) and H₅₆ₑᵗ (Figure 3 B) after subtraction of the [4Fe–4S] cluster contribution (see discussion in the Supporting Information). The presented data thus correspond to the average environment around the two iron atoms of the H-cluster. Comparison of the FTs clearly shows that the H-cluster of H₅₆ₑᵗ has greater amplitude than that of H₅₆ₑᵗ consistent with the presence of an additional heavy scatterer in the first coordination sphere of H₅₆ₑᵗ. H₅₆ₑᵗ is best fit with 3 Fe–C scatterers at 1.80 Å (from the terminal CN, terminal CO and the bridging CO), 2.5 Fe–S scatterers at 2.26 Å (from ADT ligand sulfurs and the Fe₄-bound cysteine sulfur), and an Fe–Fe scattering path at 2.60 Å (Table S4). In addition, Fe–C–O/N multiple scattering paths have to be included in the fit. The Fe–C–O/N multiple scattering paths
and the Fe–Fe scattering path are highly correlated, resulting in a somewhat larger error in the fit of the Fe–Fe distance.

For H inact, the first shell consists of the same scattering paths (Fe–C, Fe–S, Fe–Fe and Fe–C–O multiple scattering) with the same degeneracy for every path as for H ox except for the Fe–S path, for which the degeneracy needed to be increased to N = 3 (see EXAFS Discussion and Table S5 in the Supporting Information). This is consistent with the presence of an additional S ligand in H inact coordinated to one of the H–cluster Fe atoms. Attempts to separate the Fe–S contributions into shorter and longer Fe–S distances (as observed in the crystal structure) resulted in the fit paths coalescing to the same distance. This suggests that the separation of the two sulfur contributions is beyond the resolution of our data (≈0.16 Å). The ability to fit unique Fe–S contributions is further complicated by the strong correlation of the various scattering paths in our system.

Further, we note that similar to the protein crystallography, the EXAFS cannot distinguish between Cl or S as the additional ligand. Nevertheless, the first shell distances extracted from the EXAFS fits are in reasonable agreement, within the associated errors (≈0.1 Å), with the crystal structure (Table S5). For various crystal structures obtained with similar resolution diffraction data errors in the positions of the atoms, and hence bond lengths also, of up to 0.1 Å have been determined.[20] Figure 4A presents the Fe K-edge XAS spectra of DdHydAB in the H inact states (with the [4Fe–4S] cluster contribution subtracted, see Supporting Information for details). The shift of the rising edge toward higher energy for H inact is consistent with a more oxidized binuclear site (homovalent FeIIFeII in H inact vs. mixed-valent FeIIFeIII in H ox). The differences in the pre-edge region (7110–7115 eV) suggest a different coordination environment of the [2Fe] sub-cluster in the two states.

To understand the features in the experimental XAS spectra and to gain insight into the origins of the observed...
changes, time dependent density functional theory (TDDFT) calculations were performed with \( H_{\text{o}} \) and \( H_{\text{inact-SH}} \) quantum mechanics/molecular mechanics (QM/MM) models (see Supporting Information Figure S7). The TDDFT calculated pre-edge spectra (Figure 4B) reproduce the general experimental trends in terms of the pre-edge energies, intensity distributions and the onset of the rising edge. This indicates that the QM/MM models are consistent with the XAS data. Overall, these results support an oxidized \( \text{Fe}^{II}\text{Fe}^{II} \) sub-cluster for \( H_{\text{inact}} \) with a bound ligand at \( \text{Fe}_p \). We note, however, that the XAS edges are not sensitive to the exact nature of the apical ligand (see Supporting Information).

**Characterization of the \( H_{\text{inact}} \) State by Vibrational Spectroscopy**

Figure 5 shows experimental and calculated IR spectra of the \( H_{\text{ox}} \) and \( H_{\text{inact}} \) states. In the experimental spectra, all the bands of \( H_{\text{inact}} \) are shifted toward higher energy with respect to \( H_{\text{ox}} \). This is consistent with a more oxidized \( [2\text{Fe} ] \) sub-cluster in \( H_{\text{inact}} \), which leads to reduced backbonding into the \( \pi^* \) orbitals of the ligands resulting in shorter CO and CN bonds.[21] The calculated IR spectra for the \( H_{\text{ox}} \) and \( H_{\text{inact-SH}} \) QM/MM models show that the calculated frequencies are in reasonable agreement (see Supporting Information for more details). The magnitudes of the shifts are underestimated, especially for the terminal CO groups, suggesting that the experimental change in backbonding upon oxidation is not quite reproduced by the calculations (even though the \( H_{\text{inact-SH}} \) model is oxidized). Importantly, the calculated shift of the bridging CO, which should be sensitive to the addition of a new ligand is consistent with the experimental shift, albeit slightly underestimated as well.

Nuclear resonance vibrational spectroscopy (NRVS) measures vibrational sidebands coupled to nuclear transitions for Mössbauer-active nuclei, including \( ^{57}\text{Fe} \).[22] NRVS has already been used to study states of \( [\text{FeFe}] \) hydrogenases, including the catalytic intermediate \( H_{\text{hyd}} \).[23] Artificial maturation with \( ^{57}\text{Fe} \)-labelled \( [2\text{Fe} ] \) precursors results in selectively labeled \( [2\text{Fe} ]^{57}\text{Fe} \) sub-clusters.[24] As such, predominantly vibrations associated with the \( [2\text{Fe} ] \) sub-cluster are observed. Figure 6A presents the experimental NRVS spectra of \( H_{\text{ox}} \) and \( H_{\text{inact}} \), where clear differences can be observed. Low energy features in the 150–400 cm\(^{-1}\) region emerge primarily from Fe–S vibrations (bending and stretching motions). Bands around 450 cm\(^{-1}\) are mostly due to Fe–CN motion, while the strong bands between 500–600 cm\(^{-1}\) arise predominantly from Fe–CO bending and stretching modes. By correlating the experimental spectra to QM/MM NRVS calculations, the most important differences in the spectra can be interpreted. The calculated NRVS spectra (B) using the \( H_{\text{ox}} \) and \( H_{\text{inact-SH}} \) models correlate well with the experimental. Plots C and D in Figure 6 highlight the Fe–S region of the NRVS spectra. Compelling evidence for an extra sulfur bound to \( H_{\text{inact}} \) arise from the peak at \( \approx 350 \) cm\(^{-1}\) (356.52 cm\(^{-1}\) in the calculated spectrum) in the Fe–S region (marked with an asterisk). We note that the absolute prediction of complete NRVS spectra from theoretical calculations is a challenge due to the densely populated spectra and the nature of the low-energy modes involved, which are sensitive to the computational model. It is, therefore, advantageous to focus on the difference between \( H_{\text{ox}} \) and \( H_{\text{inact}} \) and on the Fe–S region.

Calculations reveal that the increased intensity in this region of the \( H_{\text{inact-SH}} \) model (compared to \( H_{\text{ox}} \)) arises from the Fe–S stretching vibration associated with an SH ligand. This feature is reproduced by the \( H_{\text{inact-SH}} \) model but cannot be reproduced in models with lighter ligands such as OH (see Supporting Information, Figure S23). The calculations cannot
exclude a Cl⁻ ligand bound to Fe₂ due to its similar mass and covalency, which gives a comparable spectrum in this region (see Supporting Information, Figure S23). However, the H_inact state can be formed in the strict absence of chloride (Figure S20), suggesting that a Cl⁻ bound to the open coordination site is unlikely. Furthermore, the experimental peak at ≈322 cm⁻¹ (324 cm⁻¹ in the calculated spectra) in H_inact (marked with a hashtag) is assigned to Fe₅S modes from the ADT ligand and cysteine, which are shifted to higher energy (compared to H_ox) due to a more oxidized [2Fe] subcluster. The calculations demonstrate the sensitivity of NRVS spectra with respect to cluster oxidation state, but also with respect to the light vs. heavy atom nature of the ligand.

In order to directly identify the exogenous sulfur ligand, we compared NRVS spectra of H_inact samples prepared using natural abundance (95% ³²S) and ³⁴S-labelled sodium sulfide. We observed small differences between the ³²S and ³⁴S spectra, particularly in the 340–360 cm⁻¹ region (Figure S24). Similar results were obtained with resonance Raman (RR) spectroscopy (Figure 7A and Figure S25), where small changes can be observed in the 340–360 cm⁻¹ region when going from the ³²S spectrum to the ³⁴S spectrum, coinciding with the expected lower frequencies of vibrations related to a heavier atom. Calculations support the assignment of two peaks in this region to Fe–S stretching modes from exogenously bound SH⁻ (Figure 7C and Table S12).

### Mechanism of H_inact Formation and Implications

Using our QM/MM model, we calculated the binding of H₂S to the H_ox state, (see Scheme in Figure S26). Sulfur likely reaches the H-cluster via diffusion through the same hydrophobic gas channels used by H₂, CO and O₂. Thus, protonation to H₂S(pKₐ 7) in solution facilitates this process. Interestingly, partial formation of a very similar H_inact state with Na₂Se could be achieved (Figure S27), but only at pH 4, where the enzyme is not very stable. This supports the idea that the neutral species (H₂S or H₂Se) are involved, as H₂Se has a much lower pKₐ (3.89) than H₂S. Direct H₂S binding to H_ox is calculated to be thermoneutral (ΔG = 0.2 kcal mol⁻¹), while deprotonation of bound H₂S by the NH group of the ADT is quite favorable (ΔG = -4.9 kcal mol⁻¹). This leads to an Fe–SH bound intermediate that is favorable with respect to free H₂S (ΔG = -4.7 kcal mol⁻¹). The H-cluster is subsequently oxidized to give an Fe³⁻FeⅡ binuclear sub-cluster, a process driven by application of oxidizing potentials. This likely proceeds via proton-coupled...
electronic rearrangement, whereby the electron is first transferred from the binuclear sub-cluster to the \([4\text{Fe}–4\text{S}]\) sub-cluster, followed by its oxidation. The oxidation steps take place by one-electron transfers from the \([4\text{Fe}–4\text{S}]\) sub-cluster, via the F-clusters, to the available high potential oxidant (including oxygen). Calculations performed with our Hinact-OH QM/MM model suggest that H₂O binds effectively at Hox but will not deprotonate via the ADT ligand, due to the larger pKₐ difference than for H₂S (see Supporting Information).

A number of other metalloenzymes bind sulfide under similar conditions. Both [NiFe] hydrogenase and Ni-dependent CO dehydrogenase are inhibited by sulfide at high applied potentials, presumably by sulfide binding in a bridging position between Ni and Fe. Recent studies suggest that, in nitrogenase, reductive displacement of an active site “belt” sulfide could be important for binding of substrates/inhibitors. Thus, binding of additional sulfides to compensate for increased positive charge on oxidized metal ions could be a common theme among enzymes, highlighting the importance of understanding how sulfide interacts with metals in nature. Diiron model complexes including thioether groups are also involved in oxidation state dependent sulfur coordination from the S group. However, in none of these cases has additional oxygen protection due to sulfur binding been demonstrated, as is observed in [FeFe] hydrogenase.

Handling air-sensitive enzymes such as [FeFe] hydrogenases under air has definite advantages, particularly, for crystallization and manipulation of crystals. Our DdHydAB Hinact structure is the first [FeFe] hydrogenase structure for which the redox state has been defined using single crystal spectroscopy, as previously demonstrated for [NiFe] hydrogenases. This provides the opportunity to directly correlate structural and spectroscopic properties of the H-cluster. Interestingly, there are very few differences in the structure of the H-cluster compared with previously published structures, suggesting an H-cluster environment that minimizes redox state dependent structural changes, lowering reorganization energy and enhancing catalysis. Air-stable [FeFe] hydrogenases may also be industrially useful for example, in fuel cells. Fuel cells containing [FeFe] hydrogenase embedded in a redox polymer have been prepared under strict anaerobic conditions, but could be prepared under air with the H inact state, simplifying the process and increasing the scalability.

The Relevance of O₂ Protection via H₅₇ In Vivo

The bacterium *Desulfovibrio desulfuricans* has evolved in anaerobic environments and, therefore, its hydrogenase is extremely oxygen sensitive, becoming inactivated irreversibly even by traces of O₂. Although the mechanism of oxygen inactivation is not yet completely understood, O₂ is believed to attack the active site by binding to the open coordination...
site on Fe$_{\text{a}}$.[16] If the bound-O$_2$ cannot be reduced to water, it may form reactive oxygen species, which could destroy the active site. As the open coordination site is blocked by sulfide in the H$_{\text{inact}}$ state, this prevents O$_2$ binding and destruction of the active site.

A plausible scenario is that in vivo, DdHydAB is constantly exposed to H$_2$S (since *Desulfovibrio desulfuricans* reduces sulfate to sulfide). Under reductive conditions H$_2$S is displaced by H$_2$, binding to the H-cluster. In the presence of oxygen, however, H$_2$S becomes locked to the H-cluster forming the H$_{\text{inact}}$ state to protect the enzyme from oxygen inactivation. It is interesting that the [FeFe] hydrogenase from *Clostridium beijerinckii* (ChH5A) can form the H$_{\text{inact}}$ state without exogenous sulfide.[14] How, the H$_{\text{inact}}$ state in this enzyme differs structurally from that in DdHydAB is not known, but it seems likely that, in the absence of available sulfide in this organism, an endogenous sulfur ligand, such as a cysteine nearby the active site, has evolved to play a role. Another important difference between these two enzymes is that DdHydAB functions as a periplasmic H$_2$ uptake enzyme with extremely high activity,[32] while ChH5A shows a strong bias for H$_2$ production.[14] This lack of activity in H$_2$ uptake may be due to spontaneous formation of H$_{\text{inact}}$ at high potentials. As DdHydAB requires exogenous sulfide, it will only be inactive when sulfide levels are high, which may serve to regulate metabolism.

**Conclusion**

Here, we provide direct structural and spectroscopic evidence for an exogenously bound sulfur in the apical coordination site of the [FeFe] hydrogenase from *Desulfovibrio desulfuricans* in the O$_2$-stable H$_{\text{inact}}$ state. In our previous work, we showed that exogenous sulfide was required for H$_{\text{inact}}$ formation, but we were unable to demonstrate if and how sulfide binds to the H-cluster. The 1.65Å crystal structure shows electron density at the apical position on the distal Fe and anomalous diffraction suggests this is consistent with sulfur. EXAFS shows an additional sulfur in the Fe-coordination environment of H$_{\text{inact}}$, compared with H$_{\text{ox}}$. Fe K-edge XAS data reveal a more oxidized [2Fe] sub-cluster in H$_{\text{inact}}$ compared to H$_{\text{ox}}$ and a different coordination environment of the Fe ions in the [2Fe] subcluster. Comparison of H$_{\text{inact}}$ and H$_{\text{ox}}$ NRVS spectra, as well as $^{32}$S/$^{34}$S isotope-labelling in both NRVS and resonance Raman spectroscopy, provide additional compelling evidence for an exogenous sulfur ligand. Calculations with an H$_{\text{inact}}$-SH model provide close agreement to all the experimental data and shed light on the mechanism of forming H$_{\text{inact}}$. In particular, the most likely pathway involved H$_2$S binding at the open coordination site, followed by proton transfer via the ADT ligand to the proton transfer channel. H$_{\text{inact}}$ formation is then completed upon proton coupled electronic reconfiguration of the H-cluster and oxidation of $[4\text{Fe}-4\text{S}^{2+}]$. Since we previously demonstrated that this in vitro H$_{\text{inact}}$ approach works for other [FeFe] hydrogenases such as *Chlamydomonas reinhardtii* (ChHydA1), it demonstrates the wider applicability of this method. Thus, it would be interesting to perform similar structural and spectroscopic studies of H$_{\text{inact}}$ in other enzymes, including ChH5A and CrHydA1. Our highly complementary structural, spectroscopic and theoretical approach represents a major advance for the understanding the function of this O$_2$-stable state and its mechanism of formation, as well as a possible implementation of these enzymes in biotechnological applications.

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Conflict of interest

The authors declare no conflict of interest.
Caught in the $H_{\text{inact}}$: [FeFe] hydrogenases are highly efficient catalysts for the interconversion of $H_2$ and $H^+$. However, they are highly $O_2$ sensitive. Here, the X-ray crystal structure of an $O_2$-stable state of [FeFe] hydrogenase, along with comprehensive spectroscopic analysis and molecular calculations reveals the presence of a sulfur ligand bound to the active site. These results provide crucial insight into $O_2$ stability mechanisms in proteins.