

Cite this: *Chem. Sci.*, 2021, 12, 3239

All publication charges for this article have been paid for by the Royal Society of Chemistry

A [3Cu:2S] cluster provides insight into the assembly and function of the Cu_Z site of nitrous oxide reductase†

Lin Zhang,^a Eckhard Bill,^b Peter M. H. Kroneck^c and Oliver Einsle^{b,*a}

Nitrous oxide reductase (N₂OR) is the only known enzyme reducing environmentally critical nitrous oxide (N₂O) to dinitrogen (N₂) as the final step of bacterial denitrification. The assembly process of its unique catalytic [4Cu:2S] cluster Cu_Z remains scarcely understood. Here we report on a mutagenesis study of all seven histidine ligands coordinating this copper center, followed by spectroscopic and structural characterization and based on an established, functional expression system for *Pseudomonas stutzeri* N₂OR in *Escherichia coli*. While no copper ion was found in the Cu_Z binding site of variants H129A, H130A, H178A, H326A, H433A and H494A, the H382A variant carried a catalytically inactive [3Cu:2S] center, in which one sulfur ligand, S_{Z2}, had relocated to form a weak hydrogen bond to the sidechain of the nearby lysine residue K454. This link provides sufficient stability to avoid the loss of the sulfide anion. The UV-vis spectra of this cluster are strikingly similar to those of the active enzyme, implying that the flexibility of S_{Z2} may have been observed before, but not recognized. The sulfide shift changes the metal coordination in Cu_Z and is thus of high mechanistic interest.

Received 20th September 2020

Accepted 3rd January 2021

DOI: 10.1039/d0sc05204c

rsc.li/chemical-science

Introduction

Nitrous oxide (N₂O) is an inert, odorless and non-toxic gas that nevertheless acts as a greenhouse agent with a global warming potential exceeding that of carbon dioxide (CO₂) by a factor of 300.¹ Its atmospheric concentration is on a steady rise, at a rate of 0.2–0.3% per year,^{1,2} which led to its designation as the most significant ozone-depleting substance of the 21st century.³ The biological reduction of N₂O is catalyzed exclusively by nitrous oxide reductase (N₂OR) as the ultimate step of the bacterial metabolic pathway of denitrification.² N₂OR is a periplasmic metalloprotein of 130 kDa that contains two copper centers, Cu_A and Cu_Z, in each monomer. Midpoint reduction potentials and spectroscopic properties of the two centers are distinct and provide detailed information about the respective electronic state of the sites.^{4–6} The three-dimensional structures of N₂OR from *Marinobacter hydrocarbonoclasticus* (PDB 1QNI),⁷ *Paracoccus denitrificans* (PDB 1FWX),⁸ *Achromobacter cycloclastes* (PDB 2IWF),⁹ *Pseudomonas stutzeri* (PDB 3SBQ),¹⁰ and *Shewanella denitrificans* (PDB 5I5M)¹¹ were reported, providing structural details for various states of the copper centers. The

dimeric N₂OR forms a tight head-to-tail arrangement that juxtaposes the Cu_A site of one monomer with the Cu_Z site of the other, placing the two sites in sufficient proximity (10 Å) to form the composite active site of the enzyme at each dimer interface.^{7,8} Cu_A is a binuclear mixed-valent [Cu^{1.5+}:Cu^{1.5+}] site^{12,13} liganded by one methionine, one tryptophan, two cysteines and two histidines that is found in a very similar form in many respiratory heme:copper oxidases,¹⁴ and it mediates one-electron transfer at a midpoint potential of approximately +260 mV vs. SHE.^{6,15,16} When isolated under anoxic conditions – a state referred to as ‘form I’¹⁷ – the Cu_Z site that binds N₂O is a tetranuclear [4Cu:μ₄-S:μ₂-S] cluster coordinated by seven histidines, as has been demonstrated for *M. hydrocarbonoclasticus* N₂OR¹⁸ and *P. stutzeri* N₂OR.¹⁰ The Cu_Z site seems to be prone to decomposition, losing sulfide ligand S_{Z2} to yield a [4Cu:μ₄-S] site designated Cu_Z* in an enzyme referred to as form II.^{19,20} Notably, a conclusion of the present discussion will be that these terms require re-definition. In other structural analyses, Cu_Z sites were modelled to contain either a μ₂-bridging water molecule, two terminal waters or an iodide anion that acted as an inhibitor, adding to a lack of clarity regarding the relevant active forms of this site.^{7–9,20}

To date, understanding of the biogenesis of Cu_Z remains fragmentary, but a series of maturation factors involved in the process have been identified.⁵ Interestingly, the apoenzyme is exported as a folded dimer *via* the Tat pathway, although the entire maturation of the metal sites occurs in the periplasm.²¹ Here the dedicated metallochaperone NosL, a lipoprotein,

^aInstitut für Biochemie, Albert-Ludwigs-Universität Freiburg, Albertstrasse 21, 79104 Freiburg im Breisgau, Germany. E-mail: einsle@biochemie.uni-freiburg.de

^bMax-Planck-Institut für Chemische Energiekonversion, Stiftstr. 34-36, D-45470 Mülheim an der Ruhr, Germany

^cFachbereich Biologie, Universität Konstanz, 78457 Konstanz, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0sc05204c



binds Cu^+ and delivers it to apo- N_2OR ,²² and a recent study has suggested its role to be in particular for Cu_Z site assembly.²³

In addition, the ABC transporter NosFY in conjunction with the periplasmic NosD protein is required to provide a sulfur species to the periplasm in order to complete Cu_Z maturation.¹⁷ We recently established a recombinant system that included all essential genes *nosRZDFYLX* for N_2OR production, able to generate functional enzyme containing both Cu_A and Cu_Z sites in *E. coli*,²⁴ facilitating mutagenesis studies of the key residues coordinating the copper sites.

Results and discussion

Properties of the seven histidine variants of Cu_Z

In the present study we focus on understanding the role of the seven ligands of the Cu_Z site, namely H129, H130, H178, H326, H382, H433 and H494 of *P. stutzeri* N_2OR (PsN_2OR). All histidine residues were individually mutated to alanine. The corresponding variants were purified (Fig. S1†), followed by spectroscopic characterization and the determination of three-dimensional structures by X-ray crystallography (Table S1†).

Not unexpectedly, six of the seven variants of the coordinating histidine residues, namely H129A, H130A, H178A, H326A, H433A and H494A only showed spectral properties confirming the presence of the Cu_A site (Fig. S2†), with absorption maxima at 485 nm, 525 nm and 795 nm, although the relative occupancy of Cu_A was found to differ (Table S2†). However, the H382A variant showed UV-vis spectra that were nearly identical to those of wild-type N_2OR (Fig. 1). The oxically isolated H382A was of blue colour, with partially oxidized copper sites as indicated by comparing the spectra of sample as isolated (Fig. 1D, blue) and ascorbate-reduced (Fig. 1E, blue).

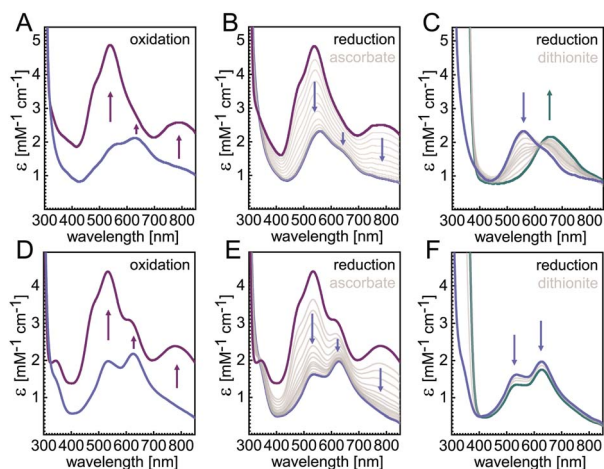


Fig. 1 UV-vis spectra of WT PsN_2OR (A–C) and variant H382A (D–F). As isolated, H382A (D) showed two absorption peaks at 535 nm and 625 nm (blue); upon oxidation with ferricyanide the spectra (purple) were similar to the wild type form I N_2OR (A). (B and E) Selective reduction of Cu_A with ascorbate yielded a two-peak spectrum indicative of a $[\text{4Cu:2S}]$ Cu_Z site (blue). Upon extended reduction with dithionite, the typical loss of the 535 nm band seen for WT PsN_2OR (C) was not observed in the H382A (F) variant (cyan).

After oxidation with ferricyanide the protein underwent a colour change to purple, with absorption maxima at 533 nm and 780 nm and two prominent shoulders at 485 nm and 625 nm (Fig. 1D, purple). These features do not merely reflect a mixed-valent Cu_A site, but correspond to the properties of a wild-type form I N_2OR (Fig. 1A),^{6,25,26} except for the more pronounced shoulder peak at 625 nm. The selective reduction of Cu_A by ascorbate resulted in two distinct bands at 535 nm and 625 nm (Fig. 1E, blue). We have earlier described this two-peak spectrum of Cu_Z as a signature of the $[\text{4Cu:2S}]$ form I, with the 625 nm maximum originating from S_{Z1} and the second peak, found at 562 nm in wild-type N_2OR , as originating from S_{Z2} ,¹⁰ although a difference assignment attributing this transition to also originate from S_{Z1} was made by other.¹⁸ The spectrum of H382A N_2OR thus indicated that – unlike in Cu_Z^* – sulfide S_{Z2} was still present, but the shifted peak indicated a change in its chemical environment. Also, the intensity ratio of the two Cu_Z bands indicated an incomplete occupancy for S_{Z2} . Intriguingly, further reduction with dithionite did not lead to a loss of either band (Fig. 1F, cyan), indicating the “ Cu_Z ” site in H382A was redox-inert. In wild-type N_2OR , dithionite reduction leads to a single charge transfer peak at 650 nm (Fig. 1C), which was assigned to the reduction of Cu_Z from a $[\text{2Cu}^+:\text{2Cu}^{2+}]$ state to a $[\text{3Cu}^+:\text{1Cu}^{2+}]$ form.^{4,18,27} We then determined the N_2O -reducing activity using reduced benzyl viologen as electron donor. H382A was not active in N_2O reduction (Fig. S3F†). We also determined the specific activities for the other six variants, and while H129A, H130A, H326A, H433A and H494A were completely inactive as expected (Fig. S3†), H178A showed low activity, with a decrease in v_{max} to $0.03 \pm 0.01 \mu\text{mol N}_2\text{O}$ per min per mg, approximately 50-fold lower than that of wild-type enzyme,²⁴ and K_M (N_2O) of $268 \pm 24 \mu\text{M}$ (Fig. S3D†). This residual activity (validated by 3 replicates) might be originated from a small portion of H178A containing a functional Cu_Z site, but the overall occupancy of possibly only 2% given the 50-fold lower activity was too low to be observed in the UV-vis spectra and crystal structure (*vide infra*).

We further proceeded to crystallize all variants and determined their three-dimensional structures to resolutions ranging from 1.67 Å to 1.49 Å (Table S3 and Fig. S4–S11†). As expected, the overall fold and dimeric structure of all variants remained unchanged,²⁴ with root-mean-squared deviations for all atoms from wild-type N_2OR at 0.34 Å (H129A), 0.28 Å (H130A), 0.14 Å (H178A), 0.48 Å (H326A), 0.11 Å (H382A), 0.32 Å (H433A), and 0.23 Å (H494A). Overall, the major structural difference was that the Ca^{2+} -binding loop (N257–D273) was disordered in some of variants. This disorder seemed to correlate with the occupancy of the Cu_A site (Tables 1 and S2†), in line with an early report that the presence of Ca^{2+} ions was required for a stable insertion of the center.¹¹

Although at different occupancies, the Cu_A site was present in all seven-variants (Table S2 and Fig. S4–S11†). The Cu_A ligands C618, W620, C622, H626, and M629 were in place to coordinate two copper ions, but the remaining ligand, H583, was in one of two possible conformations (*e.g.* Fig. S4†). In a ‘bound’ conformation, the N_δ atom of the imidazole moiety coordinated Cu_{A1} at a distance of 2.5 Å, and the N_ϵ atom formed



Table 1 Structural information on WT *PsN₂OR* the seven histidine variants

Variant	Side ^a	Ca ²⁺ -binding loop	Cu _Z	Cu _A -His583 ^b	
WT ^c	A	Ordered	[4Cu:2S]	Unbound	3.74 Å
	B	Ordered	[4Cu:2S]	Unbound	3.44 Å
H129A	A	Ordered	Empty	Bound	2.51 Å
	B	Disordered	1 Zn ²⁺	Unbound	3.46 Å
H130A	A	Ordered	Empty	Unbound	3.49 Å
	B	Disordered	Empty	Unbound	3.84 Å
H178A	A	Ordered	Empty	Bound	2.33 Å
	B	Ordered	Empty	Bound	2.46 Å
H326A	A	Disordered	1 Zn ²⁺	Unbound	3.93 Å
	B	Disordered	1 Zn ²⁺	Unbound	3.86 Å
H382A	A	Ordered	[3Cu:2S]	Unbound	3.69 Å
	B	Ordered	[3Cu:2S] ^d	Unbound	3.68 Å
H433A	A	Ordered	Empty	Unbound	3.47 Å
	B	Disordered	Empty	Unbound	3.85 Å
H494A	A	Ordered	1 Zn ²⁺	Bound	2.82 Å
	B	Disordered	2 Zn ²⁺	Unbound	3.29 Å

^a Side A shows Cu_A of chain A, Ca²⁺-binding loop and Cu_Z of chain B, which together form one active site; side B means the other way around. ^b Cu_A is present in all seven variants. 'Bound' indicates H583 is a ligand of Cu_{A1}, while 'unbound' means it is turned away. The given distances are between Cu₁ and H583; see Table S4 for more details. ^c PDB accession code 6RL0. ^d Chain A of H382A shows weak density for Cu₁ in the Cu_Z site. See Fig. S9B for more details.

a short hydrogen bond (2.7 Å) to residue D576 (Fig. S4A†). This is a state for Cu_A²⁸ that is most commonly observed and was also found in N₂OR of *M. hydrocarbonoclasticus*,⁷ *P. denitrificans*,⁸ and *A. cycloclastes*,⁹ as well as in cytochrome *c* oxidases (PDB ID: 2CUA).²⁹ In the second, 'unbound' state, however, the imidazole group of H583 was rotated away from Cu_A by approximately 135°, so that the N_δ atom now formed a hydrogen bond to residue S550, while the H-bond between N_ε atom and D576 remained unchanged (Fig. S4B†). This histidine flip at Cu_A was previously reported for *P. stutzeri* N₂OR,¹⁰ and we proposed a role in gating electron transfer from an external redox partner to Cu_A.²⁰ In our previous study, H583 showed only partial ligation of Cu_A in either form I and II of recombinant *PsN₂OR*.²⁴ However, the conformational switch of H583 was randomly distributed among the variants (Table 1). Geometry changes of Cu_A were also observed between the two conformations of H583. The major difference was that the bond lengths of Cu_{A2} to the sulfur atoms of C618 and C622 were about 0.1 Å longer when H583 was not a ligand (Table S4†).

The UV-vis spectra of the six variants H129A, H130A, H178A, H326A, H433A and H494A lacked the S → Cu charge transfer bands typically associated with the Cu_Z center (Fig. S2†). This is primarily indicative of an absence of sulfide, but the presence of histidine-coordinated Cu alone should also lead to similar N → Cu CT bands, albeit with lower intensity. Only the H326A variant might show such a peak at 630 nm (Fig. S2D†), while the other five gave no indication for the presence of Cu. Structural analysis revealed that in the variants H130A, H178A and H433A the Cu_Z site was only occupied by water molecules (Fig. S5, S6 and S10†). In the H129A, H326A and H494A variants, one or two zinc ions were instead found at the binding site for Cu_Z (Fig. S4,

S7 and S11†), as confirmed by anomalous scattering data collected at the X-ray absorption K-edge of Zn (9700 eV). The presence of Zn²⁺ in the Cu_Z site can be rationalized by non-specific incorporation *via* a periplasmic zinc chaperone such as ZinT³⁰ or ZraP,^{31,32} and the presence of multiple histidines, which are suitable ligands for zinc.³³ The metal might occupy this site if the regular Cu_Z maturation pathway is dysfunctional. Note that in no case either a single or two Cu ions were observed. Since the maturation factors NosDFYL were present for the production of all N₂OR variants in *E. coli*, copper (and sulfide) delivery should have been possible. The complete lack of Cu thus either means that each of the six histidines mutated here is essential for assembly, or that any site assembled without the full complement of ligands is unstable and the metal is quickly lost.

A [3Cu:2S] cluster in the H382A variant

The unexpected UV-vis spectra (Fig. 1D) were reflected in an unprecedented, only partially assembled [3Cu:μ₃S:μ-S] cluster at the Cu_Z site (Fig. 2B). In this cluster, Cu₁ was absent, leaving S_{Z1} as a μ₃-bridging sulfide ligating the remaining three copper ions. The histidine coordination of these was identical to native Cu_Z, in that Cu₂ was coordinated by H129 and H178, Cu₃ by H130 and H433, and Cu₄ by H494. Furthermore, the second sulfide, S_{Z2}, was still present as a ligand to Cu₄, in spite of the absence of Cu₁. It shifted position towards the nearby lysine K454, forming a hydrogen bond that stabilized the [3Cu:2S] cluster (Fig. 3B). As a consequence, the S_{Z1}-Cu₄-S_{Z2} bond angle in the [3Cu:2S] site increased by approximately 26° with respect to the one in the native, [4Cu:2S] Cu_Z. In contrast, the changes to the individual bond lengths in the cluster were insignificant (Fig. 3C).

The X-band EPR spectra for both WT N₂OR (Fig. 4A) and H382A variant (Fig. 4D) showed a similar 7-line hyperfine splitting pattern in the *g*_{||} region originating from the mixed-valent [Cu^{1.5+}:Cu^{1.5+}] state of oxidized Cu_A,^{12,14} although the

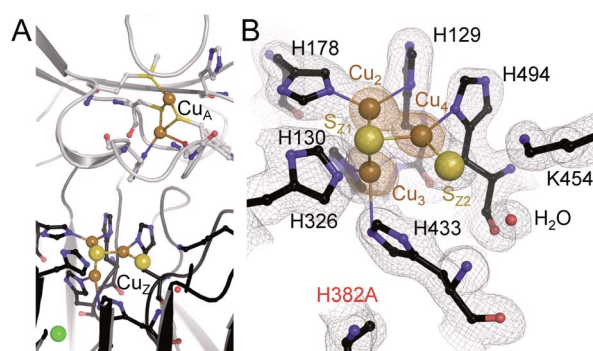


Fig. 2 Characterization of the *PsN₂OR* variant H382A. (A) The three-dimensional structure of H382A shows the presence of binuclear Cu_A site, while Cu_Z is an incomplete [3Cu:2S] cluster due to the loss of copper atom Cu_{Z1}. (B) Electron density map around the [3Cu:2S] site. The grey map is a 2F_o - F_c electron density map contoured at the 1σ level, and anomalous difference Fourier maps (data collected at the X-ray absorption edge of Cu, 9050 eV) are shown in orange at the 6σ level, confirming the presence of three copper ions.



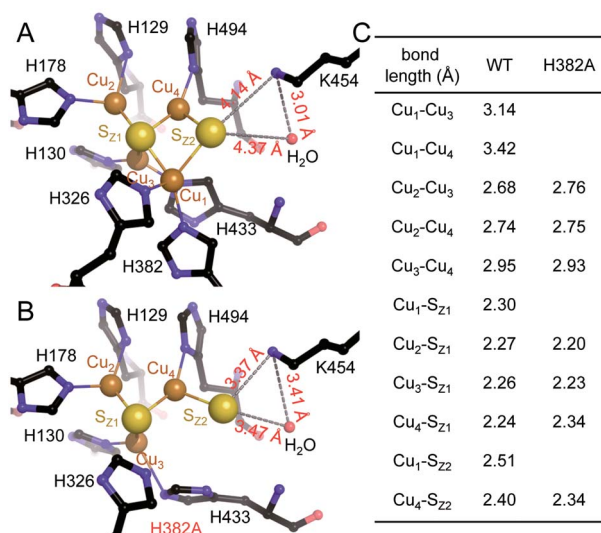


Fig. 3 Structural comparison of Cu₂ site and [3Cu:2S] site. (A) Structure of *holo*-N₂OR isolated from *P. stutzeri* (PDB code: 3SBQ) shows Cu₂ site in the [4Cu:2S] state coordinated by seven histidine residues. K454 is too far from S₂₂ (4.14 Å) to form a bond. (B) In [3Cu:2S] site of variant H382A, residues H129, H130, H178, H326, H494 are in the same conformation as observed in the wild-type. However, H433 rotates by approximately 55° due to the space released by H382A. The distance between K454 and S₂₂ is close enough to form a weak hydrogen bond (3.37 Å). (C) Comparison of bond lengths for Cu₂ and [3Cu:2S] sites.

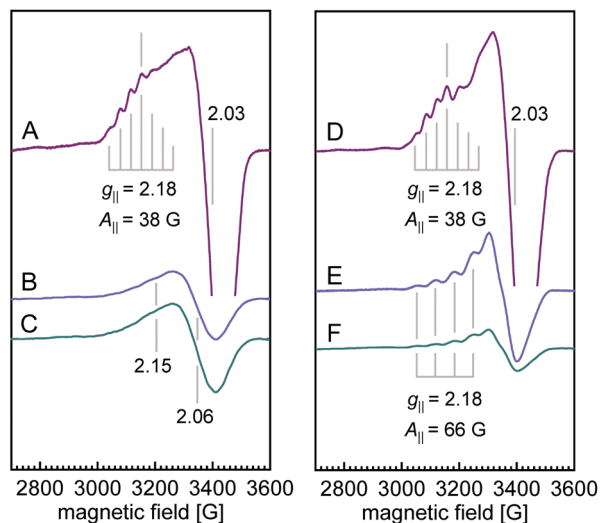


Fig. 4 X-band CW EPR spectra of WT *PsN₂OR* (A–C) and variant H382A (D–F). Spectra for ferricyanide-oxidized (purple), ascorbate-reduced (blue), and dithionite-reduced (cyan) samples are shown, respectively. The intensities are normalized to spins per protein. Temperature, 10 K; power, 0.2 mW; microwave frequency, 9.635 GHz; modulation amplitude, 7.46 G.

peak in the g_{\parallel} region of H382A was more pronounced. In the spectrum of WT *PsN₂OR*, reduction with ascorbate selectively removed the contribution of Cu_A, leaving Cu_Z in a di-cupric [2Cu²⁺:2Cu²⁺] state (Fig. 4B).⁴ The further addition of dithionite reduced Cu_Z to a [3Cu⁺:Cu²⁺] state (Fig. 4C).³⁴ In [2Cu⁺:2Cu²⁺]

Cu_Z, the two oxidized coppers couple antiferromagnetically,^{4,35} with a total spin of $S = 0$.²¹ The EPR signal shown in Fig. 4B was derived from residual [3Cu⁺:Cu²⁺] Cu_Z* that was described to have lost sulfide S₂₂.^{4,6,34} Interestingly, EPR spectra of both ascorbate- and dithionite-reduced H382A showed a 4-line hyperfine splitting pattern in the g_{\parallel} region ($g_{\parallel} = 2.18$, $A_{\parallel} = 66$ G) (Fig. 4E and F), indicating the presence of a Cu(II) ion ($S = 1/2$, ^{63/65}Cu $I = 3/2$) with its ground state in a 3d_{x²-y²}-derived molecular orbital,^{36,37} and consistent with that of typical mononuclear type 1 (T1) copper as found in plastocyanin,^{37,38} azurin,^{39–41} cucumber basic protein,⁴² as well as copper-containing nitrite reductase (NiR),^{43–45} especially from fungal laccase^{46–48} and Fet3p,^{49,50} where Cu(II) is trigonal-planar coordinated by one cysteine and two histidine residues.

Reduction with dithionite (Fig. 4F) reduced the signal intensity to about 1/3 of that for the ascorbate-reduced sample (Fig. 4E), and also destabilized the cluster, as shown by the broad peak in the $g = 2.3$ – 2.4 region (Fig. S12†). Therefore, the [3Cu:2S] cluster is very likely in a [2Cu⁺:Cu²⁺:2S²⁻]⁰ state, with Cu₄ (Fig. 3B) as Cu(II), which is trigonally coordinated in the S(μ₃-S)–N(His494)–S(μ₂-S) plane.

Residue H326, the second ligand to Cu₁ in the Cu_Z site (Fig. 3A), did not coordinate a metal in the [3Cu:2S] cluster (Fig. 3B). Thus, we expected the [3Cu:2S] cluster to be present in variant H326A as well. However, the H326A only contained a single Zn²⁺ (Fig. S7†), indicating the stabilizing effect of H326 and H382 to the Cu_Z site differs. We hypothesize that H326 is required already in the early stages of Cu_Z maturation, so that the site will not assemble if this histidine residue is mutated. H382, in contrast, only seems come into play once the entire cluster is assembled. Our data do not reveal whether Cu_Z is initially complete as a [4Cu:2S] cluster that is then prone to lose Cu₁ in the absence of the support by H382. Residue K454 (K397 in the case of N₂OR from *M. hydrocarbonoclasticus*) was proposed to play a role in the catalytic cycle of N₂OR by providing protons.^{34,51,52} Our data show that it is also involved in the stabilization – and possibly assembly – of Cu_Z through hydrogen-bonding interactions (Fig. 3B).

Concluding remarks

Beyond these questions regarding the assembly of Cu_Z *in vivo*, the [3Cu:2S] cluster in *PsN₂OR* H382A also has interesting functional implications. In the variant, the cluster retains sulfide S₂₂, but has it shifted towards residue K454, leading to UV/vis properties that fall between the forms I and II described earlier. Form II N₂OR was proposed to contain a [4Cu:S] Cu_Z* center, requiring the loss of S₂₂ as a prerequisite for reductive activation.⁴ Nevertheless the [4Cu:2S] Cu_Z state has been consistently isolated from cells grown under denitrifying conditions (*i.e.* after having turned over *in vivo*).^{4,6,10,18,52,53} The H382A variant may now help to reconcile these seemingly incompatible results, suggesting that S₂₂ can indeed change its position from ligating Cu₁ of Cu_Z to the nearby K454 (similar sulfur-shift mechanism was envisaged by Moura and Pauleta).⁵³ This would leave both Cu₁ and Cu₄ with three remaining ligands, and thus with the opportunity to bind an additional



exogenous ligand, the substrate N₂O, in a 1,3-bridging fashion. This binding mode is similar to the one proposed by Moura and Solomon,⁵² but does not require dissociation of S_{Z2} in accordance with our structural data.²⁴ It would also imply a binding mode of N₂O that is very much compatible with the N₂O binding site we observed at Cu_z after pressurizing crystals of the enzyme with the substrate gas.¹⁰

In particular, the UV-vis properties of the H382A variant highlight the fact that preparations of the enzyme that were typically assigned to a 'form II', or Cu_z* species that implies a [4Cu:S] site may well be of a different nature. The loss of the charge transfer band at 550 nm that characterizes this form II may be rooted in the shift of S_{Z2} towards K454, without being fully lost from the cluster. This finding is also in line with the frequent observation of a less well-defined, elongate electron density feature at the Cu₁-Cu₄ edge of the Cu_z cluster in different published and unpublished structures. Such features were frequently interpreted as two H₂O ligands,⁹ or inspired an originally suggested binding mode for N₂O⁵⁴ that was, however, never observed experimentally. The present data now offers an alternative rationalization for the reported spectroscopic features that may eventually lead to a unified picture of the structural and functional features of the unique and enigmatic Cu_z site.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the staff at beam lines X06SA and X06DA, Swiss Light Source, Villigen, CH, for excellent assistance with data collection, and Anne-Catherine Abel for help with the experiments. This work was supported by the European Research Council (grant 310656 to O. E.) and Deutsche Forschungsgemeinschaft (CRC 992, project no. 192904750).

Notes and references

- 1 K. Butterbach-Bahl, E. M. Baggs, M. Dannenmann, R. Kiese and S. Zechmeister-Boltenstern, *Philos. Trans. R. Soc., B*, 2013, **368**, 20130122.
- 2 A. J. Thomson, G. Giannopoulos, J. Pretty, E. M. Baggs and D. J. Richardson, *Philos. Trans. R. Soc., B*, 2012, **367**, 1157–1168.
- 3 A. R. Ravishankara, J. S. Daniel and R. W. Portmann, *Science*, 2009, **326**, 123–125.
- 4 E. M. Johnston, S. Dell'Acqua, S. Ramos, S. R. Pauleta, I. Moura and E. I. Solomon, *J. Am. Chem. Soc.*, 2014, **136**, 614–617.
- 5 S. R. Pauleta, S. Dell'Acqua and I. Moura, *Coord. Chem. Rev.*, 2013, **257**, 332–349.
- 6 T. Rasmussen, B. C. Berks, J. N. Butt and A. J. Thomson, *Biochem. J.*, 2002, **364**, 807–815.
- 7 K. Brown, M. Tegoni, M. Prudencio, A. S. Pereira, S. Besson, J. J. Moura, I. Moura and C. Cambillau, *Nat. Struct. Biol.*, 2000, **7**, 191–195.
- 8 K. Brown, K. Djinovic-Carugo, T. Haltia, I. Cabrito, M. Saraste, J. J. G. Moura, I. Moura, M. Tegoni and C. Cambillau, *J. Biol. Chem.*, 2000, **275**, 41133–41136.
- 9 K. Paraskevopoulos, S. V. Antonyuk, R. G. Sawers, R. R. Eady and S. S. Hasnain, *J. Mol. Biol.*, 2006, **362**, 55–65.
- 10 A. Pomowski, W. G. Zumft, P. M. Kroneck and O. Einsle, *Nature*, 2011, **477**, 234–237.
- 11 L. K. Schneider and O. Einsle, *Biochemistry*, 2016, **55**, 1433–1440.
- 12 J. A. Farrar, F. Neese, P. Lappalainen, P. M. H. Kroneck, M. Saraste, W. G. Zumft and A. J. Thomson, *J. Am. Chem. Soc.*, 1996, **118**, 11501–11514.
- 13 D. R. Gamelin, D. W. Randall, M. T. Hay, R. P. Houser, T. C. Mulder, G. W. Canters, S. de Vries, W. B. Tolman, Y. Lu and E. I. Solomon, *J. Am. Chem. Soc.*, 1998, **120**, 5246–5263.
- 14 M. E. Llases, M. N. Lisa, M. N. Morgada, E. Giannini, P. M. Alzari and A. J. Vila, *FEBS J.*, 2020, **287**, 749–762.
- 15 C. L. Coyle, W. G. Zumft, P. M. H. Kroneck, H. Körner and W. Jakob, *Eur. J. Biochem.*, 1985, **153**, 459–467.
- 16 C. Carreira, M. M. C. Dos Santos, S. R. Pauleta and I. Moura, *Bioelectrochemistry*, 2020, **133**, 107483.
- 17 W. G. Zumft and P. M. H. Kroneck, *Adv. Microb. Physiol.*, 2007, **52**, 107–227.
- 18 E. M. Johnston, S. Dell'Acqua, S. R. Pauleta, I. Moura and E. I. Solomon, *Chem. Sci.*, 2015, **6**, 5670–5679.
- 19 W. G. Zumft, *Microbiol. Mol. Biol. Rev.*, 1997, **61**, 533–616.
- 20 L. K. Schneider, A. Wüst, A. Pomowski, L. Zhang and O. Einsle, *Met. Ions Life Sci.*, 2014, **14**, 177–210.
- 21 S. R. Pauleta, M. S. P. Carepo and I. Moura, *Coord. Chem. Rev.*, 2019, **387**, 436–449.
- 22 M. A. McGuirl, J. A. Bollinger, N. Cosper, R. A. Scott and D. M. Dooley, *J. Biol. Inorg. Chem.*, 2001, **6**, 189–195.
- 23 S. P. Bennett, M. J. Soriano-Laguna, J. Bradley, D. A. Svistunenko, D. J. Richardson, A. J. Gates and N. E. Le Brun, *Chem. Sci.*, 2019, **10**, 4985–4993.
- 24 L. Zhang, A. Wüst, B. Prasser, C. Müller and O. Einsle, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 12822–12827.
- 25 S. Dell'Acqua, S. R. Pauleta, J. J. Moura and I. Moura, *Philos. Trans. R. Soc., B*, 2012, **367**, 1204–1212.
- 26 J. M. Charnock, A. Dreusch, H. Körner, F. Neese, J. Nelson, A. Kannt, H. Michel, C. D. Garner, P. M. Kroneck and W. G. Zumft, *Eur. J. Biochem.*, 2000, **267**, 1368–1381.
- 27 P. Chen, I. Cabrito, J. J. Moura, I. Moura and E. I. Solomon, *J. Am. Chem. Soc.*, 2002, **124**, 10497–10507.
- 28 P. M. H. Kroneck, *J. Biol. Inorg. Chem.*, 2018, **23**, 27–39.
- 29 P. A. Williams, N. J. Blackburn, D. Sanders, H. Bellamy, E. A. Stura, J. A. Fee and D. E. McRee, *Nat. Struct. Biol.*, 1999, **6**, 509–516.
- 30 A. I. Graham, S. Hunt, S. L. Stokes, N. Bramall, J. Bunch, A. G. Cox, C. W. McLeod and R. K. Poole, *J. Biol. Chem.*, 2009, **284**, 18377–18389.



- 31 I. Petit-Hartlein, K. Rome, E. de Rosny, F. Molton, C. Duboc, E. Gueguen, A. Rodrigue and J. Coves, *Biochem. J.*, 2015, **472**, 205–216.
- 32 C. A. Blindauer, *Chem. Commun.*, 2015, **51**, 4544–4563.
- 33 I. Dokmanic, M. Sikic and S. Tomic, *Acta Crystallogr., Sect. D: Biol. Crystallogr.*, 2008, **64**, 257–263.
- 34 S. Ghosh, S. I. Gorelsky, S. DeBeer George, J. M. Chan, I. Cabrito, D. M. Dooley, J. J. G. Moura, I. Moura and E. I. Solomon, *J. Am. Chem. Soc.*, 2007, **129**, 3955–3965.
- 35 S. I. Gorelsky, S. Ghosh and E. I. Solomon, *J. Am. Chem. Soc.*, 2006, **128**, 278–290.
- 36 E. I. Solomon and R. G. Hadt, *Coord. Chem. Rev.*, 2011, **255**, 774–789.
- 37 E. I. Solomon, R. K. Szilagy, S. D. George and L. Basumallick, *Chem. Rev.*, 2004, **104**, 419–458.
- 38 J. M. Guss and H. C. Freeman, *J. Mol. Biol.*, 1983, **169**, 521–563.
- 39 H. B. Gray, B. G. Malmström and R. J. P. Williams, *J. Biol. Inorg. Chem.*, 2000, **5**, 551–559.
- 40 H. Nar, A. Messerschmidt, R. Huber, M. Vandekamp and G. W. Canters, *J. Mol. Biol.*, 1991, **218**, 427–447.
- 41 C. M. Groeneveld, R. Aasa, B. Reinhammar and G. W. Canters, *J. Inorg. Biochem.*, 1987, **31**, 143–154.
- 42 L. B. LaCroix, D. W. Randall, A. M. Nersissian, C. W. G. Hoitink, G. W. Canters, J. S. Valentine and E. I. Solomon, *J. Am. Chem. Soc.*, 1998, **120**, 9621–9631.
- 43 L. B. LaCroix, S. E. Shadle, Y. N. Wang, B. A. Averill, B. Hedman, K. O. Hodgson and E. I. Solomon, *J. Am. Chem. Soc.*, 1996, **118**, 7755–7768.
- 44 K. Olesen, A. Veselov, Y. W. Zhao, Y. S. Wang, B. Danner, C. P. Scholes and J. P. Shapleigh, *Biochemistry*, 1998, **37**, 6086–6094.
- 45 H. Iwasaki, S. Noji and S. Shidara, *J. Biochem.*, 1975, **78**, 355–361.
- 46 A. E. Palmer, D. W. Randall, F. Xu and E. I. Solomon, *J. Am. Chem. Soc.*, 1999, **121**, 7138–7149.
- 47 A. Messerschmidt and R. Huber, *Eur. J. Biochem.*, 1990, **187**, 341–352.
- 48 D. M. Dooley, J. Rawlings, J. H. Dawson, P. J. Stephens, L. E. Andreasson, B. G. Malmström and H. B. Gray, *J. Am. Chem. Soc.*, 1979, **101**, 5038–5046.
- 49 T. E. Machonkin, L. Quintanar, A. E. Palmer, R. Hassett, S. Severance, D. J. Kosman and E. I. Solomon, *J. Am. Chem. Soc.*, 2001, **123**, 5507–5517.
- 50 A. B. Taylor, C. S. Stoj, L. Ziegler, D. J. Kosman and P. J. Hart, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 15459–15464.
- 51 S. Bagherzadeh and N. P. Mankad, *Chem. Commun.*, 2018, **54**, 1097–1100.
- 52 E. M. Johnston, C. Carreira, S. Dell'Acqua, S. G. Dey, S. R. Pauleta, I. Moura and E. I. Solomon, *J. Am. Chem. Soc.*, 2017, **139**, 4462–4476.
- 53 C. Carreira, R. F. Nunes, O. Mestre, I. Moura and S. R. Pauleta, *J. Biol. Inorg. Chem.*, 2020, **25**, 927–940.
- 54 S. Ghosh, S. I. Gorelsky, P. Chen, I. Cabrito, J. J. G. Moura, I. Moura and E. I. Solomon, *J. Am. Chem. Soc.*, 2003, **125**, 15708–15709.

