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Gaseous inhibition of the transsulfuration pathway by cystathionine β -synthase†

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The transsulfuration pathway plays a key role in mammals for maintaining the balance between cysteine and homocysteine, whose concentrations are critical in several biochemical processes. Human cystathionine β -synthase is a heme-containing, pyridoxal 5'-phosphate (PLP)-dependent enzyme found in this pathway. The heme group does not participate directly in catalysis, but has a regulatory function, whereby CO or NO binding inhibits the PLP-dependent reactions. In this study, we explore the detailed structural changes responsible for inhibition using quantum chemical calculations to validate the experimentally observed bonding patterns associated with heme CO and NO binding and molecular dynamics simulations to explore the medium-range structural changes triggered by gas binding and propagating to the PLP active site, which is more than 20 Å distant from the heme group. Our results support a previously proposed mechanical signaling model, whereby the cysteine decoordination associated with gas ligand binding leads to breaking of a hydrogen bond with an arginine residue on a neighbouring helix. In turn, this leads to a shift in position of the helix, and hence also of the PLP cofactor, ultimately disrupting a key hydrogen bond that stabilizes the PLP in its catalytically active form.

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Introduction

In mammals, the transsulfuration pathway plays a key role in sulfur metabolism and in intracellular redox regulation.¹ At the heart of the transsulfuration pathway is the regulation of homocysteine and cysteine concentrations. Their concentrations are crucial from two opposing perspectives: homocysteine in high concentrations is toxic in mammals,² and cysteine plays an important part in many biochemical processes including the production of the antioxidant glutathione,³ the biogenesis of iron–sulfur clusters,⁴ and in the folding and stability of cellular proteins *via* structural disulfide bonds,⁵ among many others. There is also some recent evidence that cysteine has a cytoprotective function on tumour growth.⁶

The transsulfuration pathway formally begins with homocysteine, which can react with serine to form cystathionine *via* a β -elimination reaction with catalysis by cystathionine β -synthase (CBS) (see Fig. 1).

In all organisms, CBS is a pyridoxal 5'-phosphate (PLP) dependent enzyme, but in higher organisms such as humans

(henceforth hCBS), it also contains a peripheral heme group. The broad mechanistic and kinetic features of reactions occurring in PLP-dependent enzymes are fairly well understood (see Fig. S1 for the observed reactions, ESI†).^{7–9} Besides the regulation of homocysteine concentration *via* the canonical reaction of hCBS (shown in Fig. 1), this enzyme can also catalyse various side-reactions leading to production of H₂S (shown in Fig. S1, ESI†).^{10,11} While traditionally considered as purely a toxic gas,¹² there is increasing evidence that H₂S plays a regulatory role in critical biochemical processes involved in vasodilation and neuromodulation.^{13–18} The importance of H₂S as an intracellular signalling molecule is a subject of increasing interest, potentially joining CO and NO as the third endogenous gas signalling molecule.¹⁹

hCBS exhibits various mechanisms of allosteric regulation and inhibition, based on binding of commonly-occurring intracellular molecules.²⁰ One regulatory pathway which remains challenging to understand is the binding of small gas molecules to the aforementioned peripheral heme group. In hCBS, the heme group is known to not be directly involved in

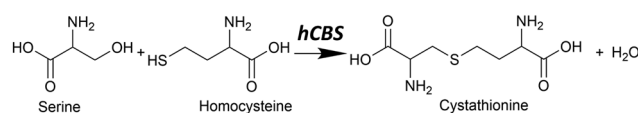


Fig. 1 The canonical reaction found in hCBS, where serine reacts with homocysteine to form cystathionine with water condensation *via* a β -elimination reaction.

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catalysis, and is located in a hydrophobic pocket near the surface of the enzyme approximately 20 Å away from the catalytic core.²¹ The heme group is 6-coordinate, and has a coordination sphere of a distal histidine and a proximal cysteine. The heme may exist in either the ferric or ferrous redox states, but is ferric in the basal state,²⁰ where ferrous heme has been found to be approximately 2-fold less active than ferric heme.²² Given the different activities of the electronic states, the heme group in hCBS has been suggested to be redox-active and has accordingly been designated as a heme redox sensor protein.^{22–24}

When reduced to the ferrous state, the heme group may bind either CO or NO, with both causing complete inhibition of hCBS.²¹ Upon CO binding, UV-vis. Spectroscopy shows a blue shift of the Soret absorption maximum from 450 nm to 422 nm.^{25,26} Similarly, NO binding causes a larger UV-vis. Blue shift from 450 nm to ~390 nm.^{27–29} Compared to those reported for CO binding, the equilibrium dissociation constants for NO binding were initially predicted to be rather a lot higher than for CO binding.^{27,28} However, more recent refinement of these values shows that the equilibrium dissociation constants for both gases are similar in magnitude, at $K_D = 0.8 \mu\text{M}$ and $K_D = 0.23 \mu\text{M}$ for CO and NO, respectively.²⁹ Relative to another heme signalling protein which binds both CO and NO, soluble guanylate cyclase (sGC), these equilibrium dissociation constants are slightly lower for CO ($K_D = 260 \mu\text{M}$ for sGC), and many orders of magnitude higher for NO ($K_D = 53 \text{ nM}–850 \text{ pM}$ for sGC).³⁰ The above dissociation constants for hCBS do not take into account the fact that it is a homodimer, with binding to the heme group in one chain potentially being able to affect the binding strength in the other chain. While there is evidence of such allosteric effects for bonding of CO, with different K_{D1} and K_{D2} being reported,^{25,26,29} we have not attempted to model these effects here. In any case, whether NO or CO binds to hCBS, the heme group is quickly oxidised back to the ferric state by O_2 , resulting in total recovery of enzymatic activity.³¹

As shown by EPR spectroscopy, CO binding is associated with decooordination of proximal cysteine in a two-step fashion ($\text{A} \rightarrow \text{D} \rightarrow \text{E}$), where the initial loss of cysteine ($\text{A} \rightarrow \text{D}$) is the rate-limiting step (see Fig. 2). The final inhibitory state, E, is found to be hexa-coordinate with proximal CO and distal histidine.^{26,32} Also based on EPR spectroscopy, the NO binding pathway has been shown to occur in two steps ($\text{A} \rightarrow \text{B} \rightarrow \text{C}$), starting with concerted displacement of the distal histidine by incoming NO ($\text{A} \rightarrow \text{B}$), followed by (again rate-limiting) decooordination of proximal cysteine ($\text{B} \rightarrow \text{C}$). In this case, the final inhibitory state, C, is determined to be penta-coordinate, with a proximal NO, and both amino acids decoordinated (see Fig. 2).^{26,27,32} Given the mechanism for CO binding, we suggest that it is feasible that D could bind to NO rather than CO equivalent, forming a different hexa-coordinate heme group with proximal NO and distal histidine (F) (see Fig. 2).

NO also binds to heme in a somewhat similar protein, soluble guanylate cyclase, and in that case, the NO-bound form is also suggested to be pentacoordinate, with NO binding being

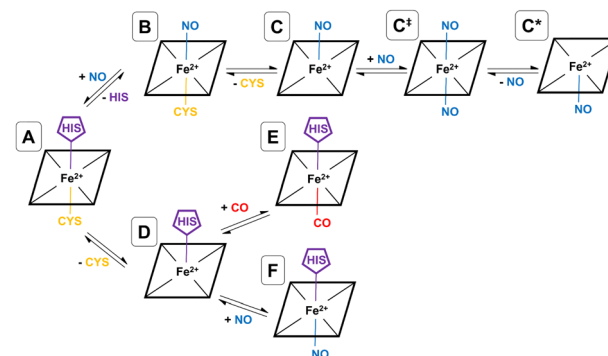


Fig. 2 Gaseous inhibitory pathways shown in hCBS. Species A, B, C, D, and E have been experimentally observed, and C and E are the final inhibitory states for NO and CO, respectively. Species C[‡], C* and F are not experimentally observed, but are computationally and biochemically suggested in this work.

accompanied by histidine sidechain decooordination. There is also some evidence that the NO-bound form involves a proximal NO ligand, *i.e.*, bound on the same side as the departing histidine sidechain. We have previously suggested³³ that this proximal NO state could be formed by reaction of a second NO equivalent with the initially formed penta-coordinate Fe(II)–NO system with a distal NO, to form a transient hexa-coordinate NO–Fe(II)–NO system (C[‡]) that would then lose the first NO to yield the isomeric penta-coordinate NO–Fe(II) (C*) (see Fig. 2). We suggested that this proximal NO binding (whose formation might be plausible during a short period of elevated NO partial pressure) could be associated with slower return to the initial state, as under low NO conditions, access to hexa-coordinate C[‡] would be difficult. It is possible that a similar mechanism occurs also for hCBS, where experiment does not allow a conclusive assignment of the bound NO state to a C or C* form. If C* is indeed formed, this could explain the reported lower rate of NO dissociation ($\sim 0.003 \text{ s}^{-1}$) compared to that of CO ($\sim 0.5 \text{ s}^{-1}$) in hCBS, as has already been suggested.²⁹

Both CO or NO binding to the heme group can inhibit hCBS, and it has been shown that this is due to conformational changes in the PLP-dependent active site upon gas binding to the heme site, ultimately leading to mechanical signalling between the two sites. However, the detailed way in which this signalling occurs and the way in which it causes the inhibition are not yet fully understood. The environment of the heme group and how it is structurally related to the PLP environment is shown in Fig. 3.

The proximal Cys52 forms a salt bridge with Arg266, where Arg266 is at one end of a short α -helix 8, and at the other end, Thr260 and Thr257 exhibit strong hydrogen bonding with the phosphate group of PLP.^{20,34–36} Mutation of both Arg266 and Thr260/Thr257 has been shown to reduce enzymatic activity.^{35,37–39} As well as the suggested mechanical signalling between the heme group and PLP cofactor, a form of electronic coupling between the heme group, α -helix 8 and the PLP cofactor has been suggested⁴⁰ to play a role in this inhibition,



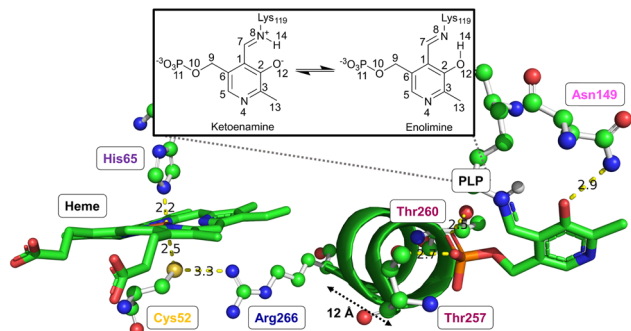


Fig. 3 Environments of the heme group and active PLP site. All relevant amino acids are shown, alongside the α -helix 8 which links the two sites. α -helix 8 is approximately 12 Å in length. For simplicity, hydrogen atoms are omitted, so intermolecular interactions (shown by dotted lines) involving hydrogen atoms are implied. The figure inset shows the tautomerisation of the PLP active site between the active ketoenamine form and inactive enolimine form.

but the nature of this coupling is unclear,[‡] and we will focus here on the mechanical signalling hypothesis.

In the mechanical signalling model,⁴¹ the active ketoenamine form of PLP and its protonated Schiff base nitrogen is stabilised by hydrogen bonding between Asn149 and the pyridoxine ring oxygen atom, as shown in Fig. 3. The protonation of the Schiff base nitrogen atom makes the imine carbon atom more susceptible to nucleophilic attack.^{7,8,41} In case the PLP centre undergoes tautomerisation, an inactive enolimine form with a neutral imine nitrogen is obtained.⁴¹ Fluorescence spectroscopy experiments show that upon binding of CO/NO, the PLP cofactor changes character from the ketoenamine to enolimine form.⁴¹ This was suggested to be due to helix 8 motion caused by breaking of the Cys52–Arg266 salt bridge, with the strong hydrogen bonds between Thr257/Thr260 and the phosphate group of PLP leading to movement of the PLP group that cause disruption of the Asn149–O[−](PLP) hydrogen bond and hence the shift of the tautomerisation equilibrium towards the inactive enolimine form. The experimental technique does not however provide access to detailed atomistic insight into the structural changes in helix 8 and the PLP site.

The goal of this study is to investigate the structural changes occurring upon CO or NO binding to the heme group. To do so, we first explore the free energy landscape associated with CO and NO binding, using quantum-chemical methods, to obtain accurate microscopic models of the C, E and F inhibited forms of the heme group. Based on these models, extended MD simulations of the inhibited forms are performed to study the motion of α -helix 8 following breakage of the Cys52–Arg266 salt

bridge, and what effect this has on the structure surrounding the PLP active site.

Computational details

Quantum chemical calculations

In all calculations, the heme ring was modelled as an Fe-porphin ring,^{42,43} and cysteine and histidine were modelled as hydrosulfide and imidazole, respectively. To provide adiabatic spin-state energetics,⁴² each of the heme derivatives shown in Fig. 2 (excluding C[‡] and C*, which is identical to C with the modelling approach used) were geometry optimised in their low-, medium-, and high-spin states using the hybrid B3LYP functional^{44,45} where Fe(II) was described using the def2-TZVP basis set and all other atoms with the def2-SVP basis set⁴⁶ (TZVP/SVP basis in Table S1, ESI[†]). The D3 dispersion correction with the Becke–Johnson damping function was used for all optimisations.^{47,48} Using second-derivative calculations, no imaginary frequencies were found, thus all heme derivatives were found to be stable minima on their corresponding potential energy surface. These optimisations and frequency analyses were performed with the Gaussian 16 program (see Table S6 for convergence criteria for these optimisations, ESI[†]).⁴⁹ See the data availability statement for the geometries for the optimised species for the various spin states.

To identify the electronic ground states, single-point DLPNO-CCSD(T₁) calculations were performed on the optimised geometries of the heme derivatives using the ORCA quantum chemistry program package (see Table S7 (ESI[†]) for the accuracy settings for all DLPNO-CCSD(T₁) calculations).⁵⁰ The coupled-cluster expansion was based on hybrid B3LYP-D3 Kohn–Sham orbitals^{44,45} obtained using the TZVP/SVP basis set combination (see Table S1, ESI[†]).⁴⁶ The density-fitting approximation was used to solve Coulomb integrals, and numerical chain-of-sphere integration was used for the Hartree–Fock exchange integrals.⁵¹ As previously shown,⁵² test calculations using Hartree–Fock orbitals for the coupled-cluster expansion showed very similar relative energies. Scalar relativistic effects were also considered through the zero-order regular approximation (ZORA).^{53–55} The ZORA recontracted versions of the def2 basis sets were used within this approximation.⁵⁶ As auxiliary basis sets, the normal def2 family of basis sets and the segmented all-electron relativistically contracted (SARC) were used.^{46,56}

On the calculated electronic ground state for each species, a higher-level DLPNO-CCSD(T₁) calculation was performed, where two-point complete basis set limit extrapolation⁵⁷ (details in ESI[†]) was applied with respect to the heme iron using two basis set combinations, TZVPP(Fe(II))/TZVP(N,O,S,C)/SVP(H) & QZVPP(Fe(II))/TZVP(N,O,S,C)/SVP(H) (see Table S1, ESI[†]).⁴⁶ All other parameters for the DLPNO-CCSD(T₁) calculations remained the same as already outlined above.

To evaluate the Gibbs free energy changes associated with the steps of the reaction pathways, two terms were added to the complete basis set extrapolated DLPNO-CCSD(T₁) electronic energy. The first is the statistical mechanics Gibbs free energy correction for each of the species at standard state (1 M) taken

[‡] Ref. 40 indeed shows very large changes in electron density distribution when performing comparative density functional theory calculations on the heme group, α -helix 8 and the PLP cofactor as separate fragments and in structures as found in hCBS, but these large changes seem to be due to artefacts associated with the gas-phase nature of the DFT calculations performed, which involve a fragment with large negative charge (the PLP cofactor) being brought into close proximity of the model for α -helix 8.



from a frequency analysis of a hybrid B3LYP-D3 calculation performed using the TZVPP/TZVP/SVP basis set combination. The translational contribution to the Gibbs free energy was removed for deoxygenated amino acid models, hydrosulfide and imidazole, since these remain bound to the protein even upon dissociation from the heme group. The second is the Gibbs free energy of solvation for each of the species, which was calculated by the difference in hybrid B3LYP-D3 electronic energy between a vacuum and implicitly solvated model using a CPCM implicit solvent model with a protein-representative dielectric constant of $\epsilon = 4$. We have also evaluated Gibbs free energy changes associated with the steps of the reaction pathways using a selection of DFT functionals – see Table S2 (ESI†) and the accompanying text.

Molecular dynamics simulations

To the best of our knowledge, there exist 8 crystal structures of hCBS in the basal state (pdb-id: 1JBQ, 1M54, 4COO, 4L0D, 4L3V, 4L27, 4L28, 5MMS).^{20,24,34,58,59} On the basis of having the best resolution (2.00 Å), being the wild-type form of hCBS, and having all residues modelled in the relevant domains, 4COO was used as the starting point in this work. For a detailed comparison of all available crystal structures for hCBS, see Table S8 (ESI†). hCBS is a biological homodimer and was modelled as such. 4COO contains all three domains of hCBS: the N-terminal heme domain, the central catalytic domain, and the C-terminal regulatory domain.³⁴ Since the C-terminal regulatory domain is not the main focus of this work and its inclusion would surely increase computational expense, it was excluded from modelling – a previously successful methodology for catalytic domain focused modelling work.⁶⁰ There are 64 missing residues at the N-termini for both chains which were not added due to uncertainty in their secondary structures. With these considerations in mind, both chains, residues 43–381, were used for modelling throughout, giving a total of 676 residues. Within this selection, there are a few residues with missing atoms – these were added back using SCWRL.⁶¹

The parameters for the covalently-bound PLP cofactor were generated using CGenFF⁶² and deemed to be satisfactory for the purposes of this work (see ESI† for parameters). To include the heme group in the system topology, the native support within CHARMM36 was used.⁶³ Where relevant, the distal His65 and proximal Cys52 were bonded to the heme group by inclusion of a `specbond.dat` file. All other atoms also used the native support within CHARMM36.⁶³ The protonation states of all titratable amino acid side-chains were assigned based on their pK_a predicted by PropKa⁶⁴ and set accordingly within the system topology (see ESI† for the protonation states of all amino acids).

Prior to production MD, equilibration was performed for species **A** (see Fig. 2), using the TIP3P water model,⁶⁵ and periodic boundary conditions with a rhombic dodecahedron unit cell, where the protein was at least 20 Å from the box edge. The total charge of the system was neutralised and set to a realistic physiological concentration of NaCl (0.15 M).⁶⁶ To remove any initial bad contacts, an energy minimisation was

performed and NVT and NPT equilibrations were performed for 1000 ps each, using the velocity-rescaling thermostat⁶⁷ and Parrinello–Rahman barostat,⁶⁸ thus equilibrating the system at 300 K and 1 bar. Lastly, positional restraints on all atoms were sequentially relaxed, and two 1000 ns simulations were performed using different starting velocities in the NPT ensemble with Parrinello–Rahman pressure coupling.⁶⁸

In order to simulate a given inhibitory pathway (**A** → **C**, **A** → **F**, and **A** → **E** – see Fig. 2), the final geometry from the trajectory of **A** was extracted and structurally modified to represent the relevant bonding for **C**, **E**, or **F**, and the `specbond.dat` file was updated appropriately. Any CO or NO molecules were introduced appropriately by making slight structural modifications to the amino acids they displace to avoid any severe steric clashes and were bonded to the heme group using the `specbond.dat` file. These modifications were performed for both monomers, so for each trajectory, there are two heme and PLP groups to be modified. In all cases, the proximal Cys52 was protonated upon deoxygenation from the heme iron. An energy minimisation was used to mitigate any bad contacts caused by the crude structural modifications. Following this, NVT and NPT equilibrations were performed for 1000 ps each, using the velocity-rescaling thermostat⁶⁷ and Parrinello–Rahman barostat.⁶⁸ Once each of the systems were sufficiently equilibrated, all positional restraints were sequentially relaxed, and simulations for each inhibitory state in the NPT ensemble with Parrinello–Rahman pressure coupling⁶⁸ were performed. For each pathway, two 1000 ns simulations were performed using different starting velocities. All MD simulations were performed with GROMACS.^{69,70}

Results and discussion

Quantum chemical calculations

Heme derivatives ground spin states. Many of the heme derivatives which have been studied in this work have previously been investigated in detail, in some cases with more accurate methods as compared to DLPNO-CCSD(T1). Nevertheless, for the sake of having a consistent dataset, and to identify the ground spin states of species **A** and **B** (which have not been studied previously, to the best of our knowledge), the ground spin states of all species **A–F** have been studied here (details in Table S3, ESI†).

In good agreement with previous *ab initio* and DFT studies,^{71–74} **F** has a clear doublet ground state, and this is also the case for **C**, again in agreement with previous work.⁷⁵ As already mentioned, there are no computational studies of species **B** to the best of our knowledge, though the corresponding ferric species has been studied, and found to have a clear low-spin form (in that case, a singlet).⁷⁶ This suggests that **B** should be low-spin (*i.e.* a doublet), which we indeed find, with a large spin state gap to the closest-lying spin state (sextet) of 20.5 kcal mol⁻¹. We calculate a stable singlet ground state for species **E**, in positive agreement with numerous previous computational studies.^{71,74,77}



Moving now to the gas-free heme adducts, it is well established from previous experimental⁷⁸ and computational⁴² work that **D** has a quintet ground state with a close-lying triplet, and this is also found here, albeit with a larger quintet-triplet gap than in some previous studies,⁴² perhaps due to the use of local coupled-cluster with a relatively small basis set. For species **A** there is, to the best of our knowledge, no previous experimental or computational work on the electronic nature of the ground state. We find the ground spin state of **A** to be a singlet, albeit with a relatively small spin state gap to the triplet of 4.2 kcal mol⁻¹.

Gibbs free energy of reactions in hCBS. Having established the ground spin states of the various heme adducts, we wish to check that the Gibbs free energy changes associated with the inhibitory pathways calculated using the ground spin states for the electronic structure calculations are consistent with experimental observations. As mentioned above, the free energies calculated herein are composed of complete basis set limit extrapolated DLPNO-CCSD(T1) electronic energies and a Gibbs free energy correction. Before discussing the results, we want to mention possible errors in our calculated values. First, the local CCSD(T) approach used is not exact,⁷⁹ especially for transition metal complexes.⁸⁰ We have tried to minimise these errors by including the improved iterative T₁ method as well as complete basis set extrapolation with respect to heme iron.⁸¹ Pantazis *et al.* recommend the use of complete PNO space extrapolation to approach the limit of CASPT2/CC calculations,⁸¹ however this was computationally unfeasible in the present study, so some uncertainty remains.

As well as the inherent uncertainties coming from the level of electronic structure theory used, our computational approach is limited in accuracy due to three aspects, which we discuss here. First, dissociation of histidine and/or cysteine sidechains from the heme group does not involve formation of a separate molecule, since the sidechains remain part of the whole protein-heme system. Hence, within the ideal-gas-like treatment of statistical mechanics, it is not clear how to treat the dissociated species, as this will not have a large relative translational partition function. In our calculations, we (somewhat arbitrarily) treat it as having no translational freedom. The rotational and vibrational parts of the partition function are computed using the usual rigid-rotor/harmonic oscillator approaches. Since the dissociated sidechain does have some range of possible motion, this may underestimate the entropy (and hence overestimate the Gibbs energy) of the dissociated states, but this may be partly compensated for by including the full rotational partition function (the sidechain will in reality have some degree of constraint on its rotational motion).

A second approximation involves the use of a continuum solvent with a generic dielectric permittivity of 4 to describe the medium surrounding the heme model used in our quantum chemical calculations. The initial species **A** is an anion, but dissociation of cysteine leads to neutral species, so solvation does affect our calculated Gibbs energies to some extent, and the polarity of the medium may not be well captured by a single dielectric constant. We have checked that varying the dielectric constant to that of water does not impact the calculated relative

Gibbs energies in a major way (for example, species **C** moves from -0.4 to -2.9 kcal mol⁻¹, with the difference being of the order of the estimated error in the quantum-chemical method used), but this approach does still introduce some additional error in our estimated Gibbs energies. We note that the departing cysteine sidechain is treated as anionic HS⁻ in our calculations. The cysteine sidechain has a pK_a near 7, so that when using a standard state for the proton concentration corresponding to pH 7 (*i.e.* using ΔG^{o'} rather than ΔG^o, see ref. 82) the standard Gibbs energies of the anionic thiolate and neutral thiol forms are almost equal, and we therefore do not need to account for the differential solvation energies of the two forms. As an additional test for the accuracy of this assumption and of the solvation treatment, we did however compute the pK_a of H₂S at the same level of theory used here, obtaining a value of 4.7, reasonably close to the experimental value of 7. Cysteine residues have pK_a values ranging from 3.5 to 12 depending on the protein environment and reactivity,⁸³ so our prediction is in reasonable qualitative agreement.

A third aspect relates to the conformational rearrangement of the overall protein structure, α-helix 8, and the PLP cofactor following CO or NO coordination, that will be discussed below. This rearrangement is spontaneous and will hence lead to a favourable contribution to the reaction free energy. As the inhibition by CO and NO is observed to be total, the free energy gain associated with relaxation of the structure after CO or NO coordination must be significant, since otherwise a small portion of the protein would remain in its active form. Assuming inhibition by a factor of 100, the Gibbs energy change should be -RT ln 100 or 2.7 kcal mol⁻¹, but the precise value is unclear. Of course, relaxation cannot be too favourable, because otherwise the protein would not be stable in the active form, even in the absence of CO or NO binding.

These provisos noted, we can compare our results with experimental observations. The pathway leading to inhibition by CO is believed to follow the route **A** → **D** → **E**. According to our calculations, the first step is endoergic in Gibbs energy terms (+11.2 kcal mol⁻¹), and the second step is exoergic (-15.1 kcal mol⁻¹) (see Fig. 4). We note that our calculations do not address energy barriers. As previously argued,⁷² gaseous ligand binding, as in the second step, can be expected to be quite fast so will have a low barrier. Decoordination of cysteine (with concomitant protonation) in the first step will likely have a larger barrier, consistent with the slow ($k \cong 10^{-3} \text{ s}^{-1}$, equivalent to a free energy barrier of 21.5 kcal mol⁻¹ at 298 K in terms of the Eyring equation) observed binding of CO to ferrous hCBS.³² The net free energy change from **A** to **E** is calculated to be -3.9 kcal mol⁻¹. Given the many sources of uncertainty mentioned above, this is consistent with the observed binding affinity and K_D = 0.8 μM, which implies ΔG^{o'} = -8.3 kcal mol⁻¹.

Experimentally, NO has been observed to inhibit hCBS by binding through the pathway, **A** → **B** → **C**. The alternative pathway *via* intermediate **D** as for CO is presumably much less favourable, given the difference in the observed kinetics. The first step **A** ≥ **B** is calculated to be exoergic (-5.2 kcal mol⁻¹) and will have a barrier which we have not studied here. Presumably, given the observed fast binding of NO, this barrier



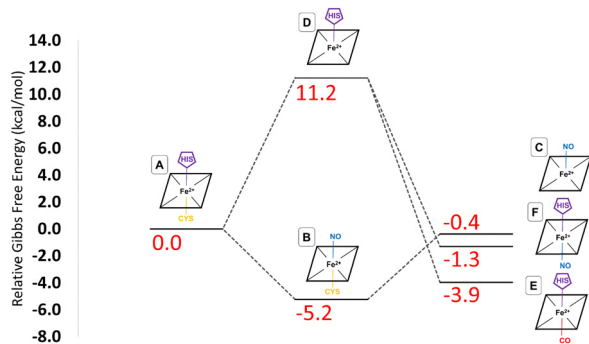


Fig. 4 Gibbs Free energy changes associated with the inhibitory pathways caused by binding of CO/NO to hCBS. Relative free energies are calculated relative to the basal species, A. All values are given in kcal mol⁻¹.

is relatively low, but the reason for the difference with CO, for which this pathway is not observed, is less clear. The second step is predicted as endoergic (+4.8 kcal mol⁻¹), leading to an overall predicted binding free energy of -0.4 kcal mol⁻¹, which differs by 8.7 kcal mol⁻¹ from the value indicated by the experimental dissociation constant of $K_D = 0.23 \mu\text{M}$, which means that $\Delta G^{\circ} = -9.1 \text{ kcal mol}^{-1}$. Formation of C requires dissociation of both the cysteine and histidine sidechains, so is twice subject to the uncertainty mentioned above relating to the translational entropy of the dissociated ligand. In case some portion of the translational entropy does still contribute to the dissociation Gibbs energy, this would account for some of the mismatch between the calculated and experimental ΔG° . The conformational change would also account for some of the difference and bearing in mind possible inaccuracies local CCSD(T), the computed values are overall roughly consistent with experiment.

The suggested endpoint of NO binding based on experiment is species C, while the *a priori* plausible F species not experimentally observed. Our calculations suggest that F would be more stable (see Fig. 4), but given the errors in our computational approach, and the small difference in computed Gibbs energies, this is not a major inconsistency.

To conclude this section, the quantum chemical calculations are consistent with the experimental observations and provide us with a solid basis for assigning the bonding pattern to be used in the MD simulations addressing the conformational changes in structure upon NO or CO coordination, which will be discussed next.

Molecular dynamics simulations. The goal of our MD simulations is to understand the changes in structure around the heme group, α -helix 8 and the PLP cofactor upon gas binding to the heme group. To the best of our knowledge, no experimental structural data on this inhibited form of hCBS is available. Based on the atomistic structures for the heme group in the inhibited form, together with extensive MD simulation, exploratory studies can be performed to predict the nature of the structural changes. Note that as hCBS is a homodimer, all simulations provide duplicate structures for the two chains. In principle, some coupling between motion in the two monomers could occur, and could account for the mentioned difference in affinity

between the first and second CO binding affinity,^{25,26,29} but we have not tried to model such allosteric effects here.

To confirm the quality of the CGenFF parameters for PLP generated herein with respect to their validity for studying the inhibitory pathways A \rightarrow C, A \rightarrow E, and A \rightarrow F, simulations of the resting state, A, were performed first. The core of our hypothesis is that when Cys52 decoordinates from the heme group, it is quickly protonated, and the strong salt bridge between Cys52 and Arg266 (see Fig. 3) would be broken as a direct consequence. Rupture of this salt bridge may cause some mechanically induced signalling pathway to PLP *via* α -helix 8 and enable the tautomerisation to the enolimine form of PLP, thus inhibiting the enzyme's activity. For this hypothesis to be supported, the salt bridge must of course be stable in the basal state A, so two separate simulations (1000 ns each) were performed for this state, with the Cys52 still coordinated to heme. These trajectories indeed show stable structures, as evidenced by a flat overall root mean square deviation (RMSD) for the protein as a whole (Fig. S2a & b, ESI[†]), an intact Cys52-Arg266 hydrogen bond throughout the simulation (Fig. S4, ESI[†]), and a largely unbroken hydrogen-bonding network around the PLP cofactor. For this latter aspect, we will first define a criterion for a hydrogen bond: the distance between the heavy atoms of the donor and acceptor must be <3.5 Å, and the donor-hydrogen-acceptor angle must be >100°. ⁸⁴ Using this criterion, returning to the simulations of the basal state A, we find a stable Asn149-O⁻(PLP) hydrogen bond throughout the simulations, with the hydrogen bond being observed for an average of 84% of the simulation time (Fig. S8, S9, S16, S17 and Table S5, ESI[†]). We do note some slight weakening occurring during one of the replica simulations for one of the homodimeric chains, where the hydrogen bond is observed for 73% of simulation time (Fig. S8, S9, S17 and Table S5, ESI[†]). Besides this key hydrogen bond, we also expect that strong hydrogen bonds are formed between the PLP phosphate group and the residues at the N-terminus of α -helix 8, ultimately coupling the positioning of α -helix 8 to the PLP cofactor. For these hydrogen bonds, we do not use the criteria described above due to the large number of possible donor-acceptor pairs complicating analysis and instead use only the donor-acceptor distances. However, the stability and minimal fluctuation of these donor-acceptor distances indicates that these hydrogen bonds are, as expected, very stable throughout the simulations (Fig. S24-S27, ESI[†]).

We now turn to simulations of the inhibited forms C and E resulting from binding of NO and CO, respectively, as well as of the possible modified inhibited form F, in which NO binds to the heme group but the histidine residue also remains bonded to the heme iron. We have performed two 1000 ns simulations of each of the final inhibitory states (C, E, and F), representing the pathways A \rightarrow C, A \rightarrow E, and A \rightarrow F. From a MD standpoint, the topologies of E and F are very similar and are generated by modification of the topology of A, such that heme-Cys52 bond is broken, and a distal CO or NO (for E and F, respectively) molecule is introduced. While similar, the forcefield parameters are also slightly different for the hexa-coordinate heme groups. Again, from modification of a topology of A, generation of a topology for C requires that both the heme-Cys52 and heme-His65



bonds are broken, and that a distal NO molecule is introduced, resulting in a penta-coordinate heme group. For all simulations of C, E, and F, Cys52 has been protonated as we expect this would be the case given the proximity to bulk solvent and the typical pK_a of cysteine sidechains. Moreover, the experimental study of the coupling between the heme group and the PLP cofactor suggested that cysteine protonation is an essential step in the inhibition pathway.⁴¹

As no structures are available for the inhibited form, it was not clear which structural changes compared to species A would be likely to be observed, nor on which timescale they could be expected to occur, though minor conformational change could certainly be expected to be visible on the micro-second timescale of the MD simulations. We monitored a range of structural coordinates relating to α -helix 8, the heme group and the PLP cofactor for each of the species C, E and F, each of the chains in the homodimers, and each of the independent simulations.

In all simulations, the Cys52-Arg266 interaction is found to break almost immediately upon modifying the heme group topology. This is expected since the cysteine is protonated as part of the change in topology. (see Fig. S5, S6 & S7 for C, E, and F, respectively, ESI[†]). The forcefield partial charge on the coordinated sulfur atom in species A is -0.8 but is only -0.23 in the neutral cysteine form, leading to much weaker electrostatic interactions. After the initial instantaneous increase in $d(S\gamma_{Cys52}-N1\eta_{Arg266})$, slower and higher-amplitude variations are also observed, associated with the fact that the proximal Cys52 is part of a loop region of the secondary protein structure. Consequently, upon decooordination from heme, Cys52 (and His65 for the case of C) is free to move as its position is no longer defined by the coordination sphere of the heme group. On the other hand, the position of the heme group is in some way locked by interaction of the propionate sidechains with Arg51 and Arg224,⁸⁵ which are not part of the flexible loop region, so that despite the loss of the ligand, the heme group remains reasonably well anchored. Motion of Cys52 following decooordination occurs in all simulations of C, E, and F, but the direction of translation of Cys52 is not systematic, presumably due to the flexibility of the loop region. Similar freedom of movement is seen for His65 for simulations of C, again with no discernible pattern. Fig. 5 shows the types of large-scale motions which are observed for both Cys52 and His65 when they have decoordinated from heme to form C.

None of the simulations of species C, E and F show rapid or major changes in the positions of the heme group, α -helix 8 or the PLP cofactor, as shown by the moderate overall RMSD values for the trajectories, with respect to their species A-derived reference points (Fig. S2 & S3, ESI[†]). Nevertheless, close inspection of these trajectories does show evidence for structural change consistent with the mechanical signalling hypothesis. The main metric showing this is the previously mentioned Asn149-O⁻(PLP) hydrogen bond. Considering the three species C, E and F, the two chains of the homodimer, and the two independent MD simulations for each case, there are 12 time series to be analysed. Again, using the criteria defined

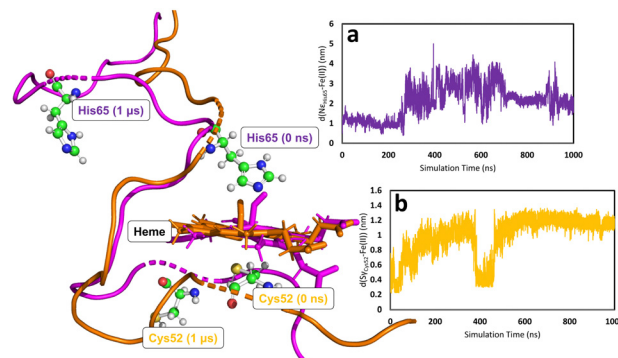


Fig. 5 Large scale motions of Cys52 and His65 during one of the 1000 ns simulations for species C for chain B. The heme group and flexible loop region is shown in orange and purple for 0 and 1000 ns, respectively. Figure insets (a) and (b) show the motion of His65 and Cys52, respectively, relative to their initial coordination with heme Fe(II).

above, in a way that is not seen for the simulations of A, in three of these time series, the Asn149-O⁻(PLP) hydrogen bond is unambiguously disrupted in a major way, with it being observed for 38, 25, and 13% of simulation times (Fig. S10, S11, S14, S15, S19, S22, S23 and Table S5, ESI[†]). In a further three time series, the hydrogen bond shows some weakening in a similar fashion to that which was observed in one of the replica simulations of A, where the hydrogen bond is observed for 75, 71, and 79% of simulation times (Fig. S10, S11, S14, S15, S18, S19, S22 and Table S5, ESI[†]). The remaining six simulations do not show convincing evidence of hydrogen bond elongation. Since our MD simulations cannot account for the tautomeric change in the PLP cofactor, any weakening of the hydrogen bonding must be due to the mechanical signalling through α -helix 8. We note that the lack of clear changes in hydrogen bonding in some of the timeseries may be due to insufficient sampling, evidenced by the fact that there is no clear pattern concerning the differences between species C, E and F. In fact, the species showing the least evidence for weakening of the interaction, E (Fig. S12, S13, S20, S21 and Table S5, ESI[†]), is treated in an extremely similar way within the forcefield to the related species F, which shows clear elongation of the Asn149-O⁻(PLP) hydrogen bond in some of the time series (Fig. S14 and S15, ESI[†]), strengthening the suggestion that the differences between simulations are indeed due to insufficient sampling.

Other structural metrics relating to the PLP cofactor are stable throughout the simulation. For example, the strong hydrogen bonds between the PLP phosphate group and the α -helix 8 N-terminus are largely unbroken throughout (Fig. S24–S39, ESI[†]) (some of the changes in distance shown in these plots arise due to switching of hydrogen bonds between the different phosphate oxygens). The α -helix is thereby firmly attached to the PLP cofactor, and motion of the α -helix leads to the above-mentioned weakening of the interaction of the PLP cofactor with Asn149. See the data availability statement for initial GROMACS geometry files for each of the species A, C, E, F as well as selected forcefield parameters.



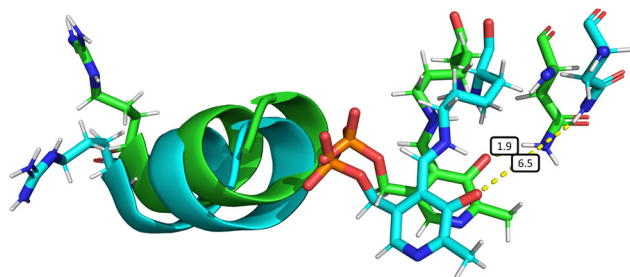


Fig. 6 Overall structural changes following decooordination of Cys52 and rotation of Arg266. The initial structure at 0 ns is shown in green and a snapshot after 1000 ns is shown in blue (species F, chain A). The bond length $d(\text{N}\delta_{\text{Asn149}}-\text{O}12_{\text{PLP}})$ is highlighted in the figure, increasing from 1.9 Å (at 0 ns) to 6.5 Å (at 1000 ns). This figure was generated by α -carbon backbone alignment in PyMOL.⁸⁶

The elongation of the Asn149–O[−](PLP) hydrogen bond is associated with other slight structural changes relating to the position of α -helix 8 and the PLP cofactor. Some insight into these changes can be obtained by comparing a typical species A structure and a typical structure in which the hydrogen bond is surely broken (Fig. 6). This comparison is somewhat artificial since the choice of the two structures is necessarily arbitrary, but it has the benefit of providing an overall picture of the changes arising with the mechanical coupling mechanism. In Fig. 6, the two structures are aligned based on the positions of the α -carbons of all amino acids, so that their protein backbones are maximally close to each other. As can be seen, there is a clear change in the relative position of α -helix 8 and the PLP cofactor with respect to Asn149, confirming that indeed α -helix 8 does play a role in mechanically induced signalling between the heme group and PLP cofactor.

In principle, biased simulation techniques together with free energy evaluation methods could be used to quantify the differences arising when switching from the Cys-heme bound species A to the free Cys species C, E and F. This would however be rather challenging to model, given the extensive small structural differences between the various species, and the lack of a clearly identified reaction coordinate associated with the mechanical coupling. Hence this has not been attempted here.

Conclusions

In this work, we have aimed to investigate in more detail the role of a peripheral heme group found in hCBS that does not directly engage in catalytic activities, but instead exerts regulatory control over the enzyme. This regulatory function manifests when CO or NO gas molecules bind to the heme group, leading to complete inhibition of PLP-dependent reactions.

Initially, we validated the CO/NO bonding models proposed by EPR spectroscopy through detailed quantum chemical studies, analysing the Gibbs free energy changes associated with the inhibitory pathways. Despite noted limitations in computational accuracy, our findings demonstrated positive agreement with previous calculations on related heme derivatives, often employing more sophisticated quantum chemical methods,

and our calculated Gibbs energies are consistent with observed thermodynamics, giving confidence in the atomistic models for the basal and ligand-bound states.

Using forcefields with bonding patterns identified using the quantum chemical calculations, several 1000 ns molecular dynamics simulations on the various inhibitory species were performed. Through these simulations, we sought to investigate in greater detail the proposed mechanical signalling model, which suggests that the rupture of the Cys52–Arg266 salt bridge, triggered by gas binding, initiates a cascade of allosteric changes. We find that motion of α -helix 8 (which Arg266 is a part of) which happens following breakage of the Cys52–Arg266 salt bridge leads to destabilisation of the active ketoenamine form of PLP. This destabilization arises from the elongation of a crucial hydrogen bond between Asn149 and a pyridoxine ring oxygen atom, suggesting a lowered barrier to tautomerization which produces the inactive enolimine form of PLP.

Our findings underscore the intriguing regulatory role of CO and NO in hCBS, indicating that both gases have the potential to inhibit the enzyme both experimentally and theoretically. Moreover, reactions catalysed by hCBS at the PLP active site cause production of H₂S, yet further indicating the emerging theme of complex interplay between CO, NO and H₂S as signalling molecules.

Data availability

The data referenced throughout this have been uploaded as part of the electronic supplementary material and files for replicating quantum chemical calculations and molecular dynamics simulations are available *via* the KU Leuven Research Data Repository, see <https://doi.org/10.48804/EBWSWB>.

Author contributions

Neil R. McFarlane: conceptualisation, data curation, formal analysis, visualisation, writing – original draft, writing – review & editing; Jiāngli Gui: data curation, formal analysis, visualisation; Julianna Oláh: conceptualisation, supervision, writing – review & editing; Jeremy N. Harvey: conceptualisation, supervision, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 J. I. Sbodio, S. H. Snyder and B. D. Paul, *Br. J. Pharmacol.*, 2019, **176**, 583–593.
- 2 J. Perla-Kaján, T. Twardowski and H. Jakubowski, *Amino Acids*, 2007, **32**, 561–572.
- 3 Y. Chen, H. G. Shertzer, S. N. Schneider, D. W. Nebert and T. P. Dalton, *J. Biol. Chem.*, 2005, **280**, 33766–33774.
- 4 R. Lill and U. Mühlenhoff, *Annu. Rev. Cell Dev. Biol.*, 2006, **22**, 457–486.
- 5 C. S. Sevier and C. A. Kaiser, *Nat. Rev. Mol. Cell Biol.*, 2002, **3**, 836–847.
- 6 H.-F. Zhang, R. I. K. Geltink, S. J. Parker and P. H. Sorensen, *Trends Cell Biol.*, 2022, **32**, 800–814.
- 7 A. Salvà, J. Donoso, J. Frau and F. Muñoz, *J. Phys. Chem. A*, 2003, **107**, 9409–9414.
- 8 E. F. Oliveira, N. M. F. S. A. Cerqueira, P. A. Fernandes and M. J. Ramos, *J. Am. Chem. Soc.*, 2011, **133**, 15496–15505.
- 9 J. Liang, Q. Han, Y. Tan, H. Ding and J. Li, *Front. Mol. Biosci.*, 2019, **6**, 4.
- 10 A. P. Landry, J. Roman and R. Banerjee, *Curr. Opin. Struct. Biol.*, 2021, **71**, 27–35.
- 11 T. Chiku, D. Padovani, W. Zhu, S. Singh, V. Vitvitsky and R. Banerjee, *J. Biol. Chem.*, 2009, **284**, 11601–11612.
- 12 R. J. Reiffenstein, W. C. Hulbert and S. H. Roth, *Annu. Rev. Pharmacol. Toxicol.*, 1992, **32**, 109–134.
- 13 K. Abe and H. Kimura, *J. Neurosci.*, 1996, **16**, 1066–1071.
- 14 R. Hosoki, N. Matsuki and H. Kimura, *Biochem. Biophys. Res. Commun.*, 1997, **237**, 527–531.
- 15 W. Zhao, J. Zhang, Y. Lu and R. Wang, *EMBO J.*, 2001, **20**, 6008–6016.
- 16 W. Zhao and R. Wang, *Am. J. Physiol.: Heart Circ. Physiol.*, 2002, **283**, H474–H480.
- 17 M. Whiteman, J. S. Armstrong, S. H. Chu, S. Jia-Ling, B.-S. Wong, N. S. Cheung, B. Halliwell and P. K. Moore, *J. Neurochem.*, 2004, **90**, 765–768.
- 18 C. Szabo, *Biochem. Pharmacol.*, 2018, **149**, 5–19.
- 19 R. Wang, *FASEB J.*, 2002, **16**, 1792–1798.
- 20 M. Meier, M. Janosik, V. Kery, J. P. Kraus and P. Burkhard, *EMBO J.*, 2001, **20**, 3910–3916.
- 21 S. Singh, P. Madzelan and R. Banerjee, *Nat. Prod. Rep.*, 2007, **24**, 631–639.
- 22 S. Taoka, S. Ohja, X. Shan, W. D. Kruger and R. Banerjee, *J. Biol. Chem.*, 1998, **273**, 25179–25184.
- 23 R. Banerjee and C. Zou, *Arch. Biochem. Biophys.*, 2005, **433**, 144–156.
- 24 S. Taoka, B. W. Lepore, Ö. Kabil, S. Ojha, D. Ringe and R. Banerjee, *Biochemistry*, 2002, **41**, 10454–10461.
- 25 S. Taoka, M. West and R. Banerjee, *Biochemistry*, 1999, **38**, 2738–2744.
- 26 M. Puranik, C. L. Weeks, D. Lahaye, Ö. Kabil, S. Taoka, S. B. Nielsen, J. T. Groves, R. Banerjee and T. G. Spiro, *J. Biol. Chem.*, 2006, **281**, 13433–13438.
- 27 S. Taoka and R. Banerjee, *J. Inorg. Biochem.*, 2001, **87**, 245–251.
- 28 C. Gherasim, P. K. Yadav, O. Kabil, W.-N. Niu and R. Banerjee, *PLoS One*, 2014, **9**, e85544.
- 29 J. B. Vicente, H. G. Colaço, M. I. S. Mendes, P. Sarti, P. Leandro and A. Giuffrè, *J. Biol. Chem.*, 2014, **289**, 8579–8587.
- 30 G. Wu, I. Sharina and E. Martin, *Front. Mol. Biosci.*, 2022, **9**, 1007768.
- 31 S. Carballal, P. Madzelan, C. F. Zinola, M. Graña, R. Radi, R. Banerjee and B. Alvarez, *Biochemistry*, 2008, **47**, 3194–3201.
- 32 S. Carballal, E. Cuevasanta, I. Marmisolle, O. Kabil, C. Gherasim, D. P. Ballou, R. Banerjee and B. Alvarez, *Biochemistry*, 2013, **52**, 4553–4562.
- 33 A. M. Rozza, M. Papp, N. R. McFarlane, J. N. Harvey and J. Oláh, *Chem. – Eur. J.*, 2022, **28**, e202200930.
- 34 T. J. McCorvie, J. Kopec, S.-J. Hyung, F. Fitzpatrick, X. Feng, D. Termine, C. Strain-Damerell, M. Vollmar, J. Fleming, J. M. Janz, C. Bulawa and W. W. Yue, *J. Biol. Chem.*, 2014, **289**, 36018–36030.
- 35 P. K. Yadav, P. Xie and R. Banerjee, *J. Biol. Chem.*, 2012, **287**, 37611–37620.
- 36 R. Evande, S. Ojha and R. Banerjee, *Arch. Biochem. Biophys.*, 2004, **427**, 188–196.
- 37 S. Singh, P. Madzelan, J. Stasser, C. L. Weeks, D. Becker, T. G. Spiro, J. Penner-Hahn and R. Banerjee, *J. Inorg. Biochem.*, 2009, **103**, 689–697.
- 38 A. T. Smith, Y. Su, D. J. Stevens, T. Majtan, J. P. Kraus and J. N. Burstyn, *Biochemistry*, 2012, **51**, 6360–6370.
- 39 A. Bhatt and Md. E. Ali, *J. Biomol. Struct. Dyn.*, 2022, **40**, 12690–12698.
- 40 A. Bhatt, A. Mukhopadhyaya and Md. E. Ali, *J. Phys. Chem. B*, 2022, **126**, 4754–4760.
- 41 C. L. Weeks, S. Singh, P. Madzelan, R. Banerjee and T. G. Spiro, *J. Am. Chem. Soc.*, 2009, **131**, 12809–12816.
- 42 M. Radoń, *J. Chem. Theory Comput.*, 2014, **10**, 2306–2321.
- 43 Md. E. Ali, B. Sanyal and P. M. Oppeneer, *J. Phys. Chem. B*, 2012, **116**, 5849–5859.
- 44 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 45 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
- 46 F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305.
- 47 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.
- 48 S. Grimme, S. Ehrlich and L. Goerigk, *J. Comput. Chem.*, 2011, **32**, 1456–1465.
- 49 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark,



- J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision C.01*, Gaussian, Inc., Wallingford CT, 2016.
- 50 F. Neese, F. Wennmohs, U. Becker and C. Riplinger, *J. Chem. Phys.*, 2020, **152**, 224108.
- 51 B. Helmich-Paris, B. de Souza, F. Neese and R. Izsák, *J. Chem. Phys.*, 2021, **155**, 104109.
- 52 Z. Benedek, P. Timár, T. Szilvási and G. Barcza, *J. Comput. Chem.*, 2022, **43**, 2103–2120.
- 53 E. van Lenthe, J. G. Snijders and E. J. Baerends, *J. Chem. Phys.*, 1996, **105**, 6505–6516.
- 54 E. van Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1993, **99**, 4597–4610.
- 55 E. van Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1994, **101**, 9783–9792.
- 56 D. A. Pantazis, X.-Y. Chen, C. R. Landis and F. Neese, *J. Chem. Theory Comput.*, 2008, **4**, 908–919.
- 57 F. Neese and E. F. Valeev, *J. Chem. Theory Comput.*, 2011, **7**, 33–43.
- 58 J. B. Vicente, H. G. Colaço, F. Malagrinò, P. E. Santo, A. Gutierrez, T. M. Bandejas, P. Leandro, J. A. Brito and A. Giuffrè, *Oxid. Med. Cell. Longevity.*, 2017, **2017**, e8940321.
- 59 J. Ereño-Orbea, T. Majtan, I. Oyenarte, J. P. Kraus and L. A. Martínez-Cruz, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, E3790–E3799.
- 60 S. Gupta, S. Kelow, L. Wang, M. D. Andrade, R. L. Dunbrack and W. D. Kruger, *J. Biol. Chem.*, 2018, **293**, 13921–13931.
- 61 G. G. Krivov, M. V. Shapovalov and R. L. Dunbrack Jr., *Proteins: Struct., Funct., Bioinf.*, 2009, **77**, 778–795.
- 62 K. Vanommeslaeghe, E. Hatcher, C. Acharya, S. Kundu, S. Zhong, J. Shim, E. Darian, O. Guvench, P. Lopes, I. Vorobyov and A. D. MacKerell, *J. Comput. Chem.*, 2010, **31**, 671–690.
- 63 J. Huang and A. D. MacKerell, *J. Comput. Chem.*, 2013, **34**, 2135–2145.
- 64 M. H. M. Olsson, C. R. Søndergaard, M. Rostkowski and J. H. Jensen, *J. Chem. Theory Comput.*, 2011, **7**, 525–537.
- 65 P. Mark and L. Nilsson, *J. Phys. Chem. A*, 2001, **105**, 9954–9960.
- 66 G. A. Ross, A. S. Rustenburg, P. B. Grinaway, J. Fass and J. D. Chodera, *J. Phys. Chem. B*, 2018, **122**, 5466–5486.
- 67 G. Bussi, D. Donadio and M. Parrinello, *J. Chem. Phys.*, 2007, **126**, 014101.
- 68 M. Parrinello and A. Rahman, *J. Appl. Phys.*, 1981, **52**, 7182–7190.
- 69 M. J. Abraham, T. Murtola, R. Schulz, S. Páll, J. C. Smith, B. Hess and E. Lindahl, *SoftwareX*, 2015, **1–2**, 19–25.
- 70 S. Pronk, S. Páll, R. Schulz, P. Larsson, P. Bjelkmar, R. Apostolov, M. R. Shirts, J. C. Smith, P. M. Kasson, D. van der Spoel, B. Hess and E. Lindahl, *Bioinformatics*, 2013, **29**, 845–854.
- 71 N. Strickland and J. N. Harvey, *J. Phys. Chem. B*, 2007, **111**, 841–852.
- 72 A. Lábás, D. K. Menyhárd, J. N. Harvey and J. Oláh, *Chem. – Eur. J.*, 2018, **24**, 5350–5358.
- 73 J. Oláh and J. N. Harvey, *J. Phys. Chem. A*, 2009, **113**, 7338–7345.
- 74 M. Radoń and K. Pierloot, *J. Phys. Chem. A*, 2008, **112**, 11824–11832.
- 75 C. Rovira, K. Kunc, J. Hutter, P. Ballone and M. Parrinello, *J. Phys. Chem. A*, 1997, **101**, 8914–8925.
- 76 D. A. Scherlis, C. B. Cymeryng and D. A. Estrin, *Inorg. Chem.*, 2000, **39**, 2352–2359.
- 77 S. Liu, S. Xia, D. Yue, H. Sun and H. Hirao, *Inorg. Chem.*, 2022, **61**, 17494–17504.
- 78 L. Pauling and C. D. Coryell, *Proc. Natl. Acad. Sci. U. S. A.*, 1936, **22**, 210–216.
- 79 D. G. Liakos, Y. Guo and F. Neese, *J. Phys. Chem. A*, 2020, **124**, 90–100.
- 80 M. Feldt, Q. M. Phung, K. Pierloot, R. A. Mata and J. N. Harvey, *J. Chem. Theory Comput.*, 2019, **15**, 922–937.
- 81 M. Drosou, C. A. Mitsopoulou and D. A. Pantazis, *J. Chem. Theory Comput.*, 2022, **18**, 3538–3548.
- 82 M. R. Gunner, T. Murakami, A. S. Rustenburg, M. Işık and J. D. Chodera, *J. Comput.-Aided Mol. Des.*, 2020, **34**, 561–573.
- 83 D. W. Bak, T. J. Bechtel, J. A. Falco and E. Weerapana, *Curr. Opin. Chem. Biol.*, 2019, **48**, 96–105.
- 84 E. N. Baker and R. E. Hubbard, *Prog. Biophys. Mol. Biol.*, 1984, **44**, 97–179.
- 85 V. Guallar and B. Olsen, *J. Inorg. Biochem.*, 2006, **100**, 755–760.
- 86 L. Schrödinger and W. DeLano, *PyMOL (version 2.4.0)* <https://www.pymol.org/pymol>, 2020.

