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Evolutionary adaptation from hydrolytic to oxygenolytic catalysis at the α/β -hydrolase fold†

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Protein fold adaptation to novel enzymatic reactions is a fundamental evolutionary process. Cofactor-independent oxygenases degrading *N*-heteroaromatic substrates belong to the α/β -hydrolase (ABH) fold superfamily that typically does not catalyze oxygenation reactions. Here, we have integrated crystallographic analyses under normoxic and hyperoxic conditions with molecular dynamics and quantum mechanical calculations to investigate its prototypic 1-*H*-3-hydroxy-4-oxoquinaldine 2,4-dioxygenase (HOD) member. O₂ localization to the "oxyanion hole", where catalysis occurs, is an unfavorable event and the direct competition between dioxygen and water for this site is modulated by the "nucleophilic elbow" residue. A hydrophobic pocket that overlaps with the organic substrate binding site can act as a proximal dioxygen reservoir. Freeze-trap pressurization allowed the structure of the ternary complex with a substrate analogue and O₂ bound at the oxyanion hole to be determined. Theoretical calculations reveal that O₂ orientation is coupled to the charge of the bound organic ligand. When 1-*H*-3-hydroxy-4-oxoquinaldine is uncharged, O₂ binds with its molecular axis along the ligand's C2–C4 direction in full agreement with the crystal structure. Substrate activation triggered by deprotonation of its 3-OH group by the His-Asp dyad, rotates O₂ by approximately 60°. This geometry maximizes the charge transfer between the substrate and O₂, thus weakening the double bond of the latter. Electron density transfer to the O₂(π^*) orbital promotes the formation of the peroxide intermediate via intersystem crossing that is rate-determining. Our work provides a detailed picture of how evolution has repurposed the ABH-fold architecture and its simple catalytic machinery to accomplish metal-independent oxygenation.

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Introduction

The switch from a reducing to an oxidative atmosphere during our planet's history spawned the emergence of many O₂-dependent enzymes, several of which evolved from pre-existing

classes, whose original functions are unrelated to dioxygen chemistry.^{1–4} The α/β -hydrolase (ABH) fold is a common and versatile protein architecture found in all three domains of life. The comprehensive ESTHER (ESTERase, α/β -Hydrolase Enzymes and Relatives) database lists more than 70,000 family members that typically share little sequence similarity but display a common fold and catalytic machinery.⁵ Their core structure is an eight-stranded β -sheet surrounded by α -helices often decorated with additional structural elements, generally referred to as cap or lid domains, that confer functional variation (Fig. S1†).^{6,7} Mechanistically, most ABH-fold members rely on the conserved nucleophile–histidine–acidic residue proton-relay system of serine hydrolases. In keeping with this, the majority of ABHs are lipases, proteases, esterases, thioesterases, dehalogenases, and epoxide hydrolases that use H₂O as the co-substrate. However, this is not universal and other members include haloperoxidases, lyases, and even oxygenases that employ H₂O₂, HCN, or O₂, respectively, for catalysis.⁸ This underscores the functional malleability of the ABH fold and its simple catalytic triad.

Arthrobacter nitroguajacolicus R 61a 1-*H*-3-hydroxy-4-oxoquinaldine 2,4-dioxygenase (HOD) and *Pseudomonas putida*

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33/1 1-*H*-3-hydroxy-4-oxoquinoline 2,4-dioxygenase (QDO) were the first dioxygenases discovered to belong to the ABH-fold family.^{9,10} They catalyze the O₂-dependent breakdown of *N*-heteroaromatic quinolone-type substrates with concomitant carbon monoxide production (Fig. 1A). Recently, Aqdc1 and Aqdc2 from *R. erythropolis*, and Aqdc from *M. abscessus* subsp. *abscessus*, have been also identified as ABH-fold dioxygenases that catalyze the same reaction,^{11,12} and the family of quinolone-type degrading ABH-fold dioxygenases has been significantly expanded by bioinformatic analyses with the identification of more than 150 *bona fide* new members, nine of which have been validated experimentally.¹³ Atypical for oxygenases, these enzymes promote the activation of the triplet ground-state O₂ molecule without the aid of metal centers or external organic cofactors.¹⁴ Kinetic measurements indicate that their reaction mechanism relies on a fast initial step, during which the substrates' 3-hydroxyl group is deprotonated by the His-Asp

subset of the triad, thus promoting its activation for molecular oxygen attack (step 1, Fig. 1B), whilst the nucleophile is not essential.^{4,15,16} Consistent with this, in various organisms, a serine to alanine replacement is observed for the 'nucleophile' residue (Fig. S2†). The role of the His-Asp dyad as a general base is supported by the crystal structures of HOD and Aqdc that have been solved in mechanistically relevant states.^{4,17} The dispensability of the nucleophile is in stark contrast with that observed for most ABHs for which the nucleophile is required for the formation of the mandatory covalent acyl-enzyme intermediate.

Cofactor-independent ABH-fold oxygenases are intriguing from a mechanistic viewpoint as the spin-forbidden oxygenation reaction is enabled by a minimalistic catalytic toolbox. Theoretical investigations have put forward different hypotheses for the mechanism that allows the quantum chemical hurdle of the direct reaction between the singlet-state substrate

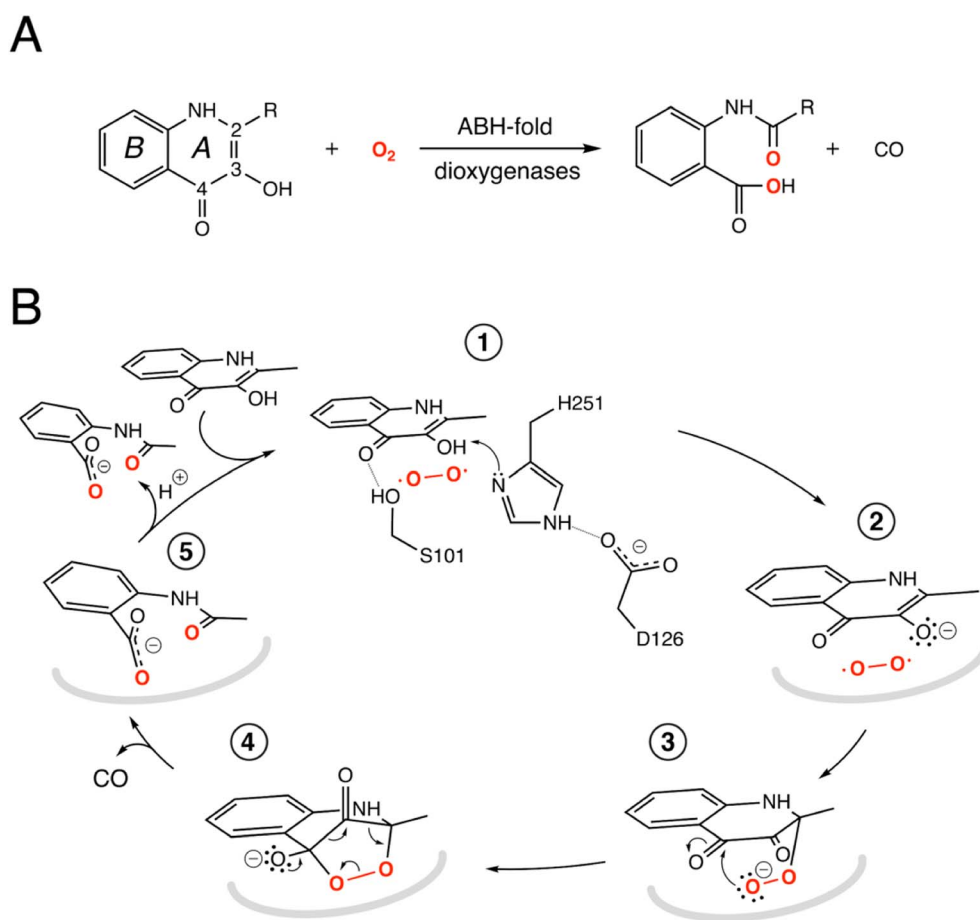


Fig. 1 ABH-fold cofactor-independent dioxygenase reaction and mechanism. (A) Scheme of the reaction catalyzed by the bacterial cofactor-independent ABH-fold dioxygenases. HOD primarily catalyzes conversion of 1-*H*-3-hydroxy-4-oxoquinoline (R=CH₃, QND) to *N*-acetyl-anthranilate (NAA). In the reaction, the A heterocycle is disrupted by the formation of carbon monoxide as a by-product. The compound 2-methyl-quinolin-4(1*H*)-one (MQO) used in this work features –H instead of –OH at position 3; (B) reaction mechanism as exemplified by HOD. The structure of the anaerobic HOD–QND complex revealed that the 3-OH group of QND is deprotonated by the H251–D126 pair of the 'nucleophile–histidine–acidic' ABH-fold triad (steps 1 and 2). S101 at the sharp structural turn known as the 'nucleophilic elbow' further stabilizes the substrate at the active site. In some ABH-fold dioxygenases an alanine residue replaces the serine at the 'nucleophilic elbow'. The reaction is then assumed to proceed *via* the peroxide intermediates (3 and 4) to the formation of the product NAA (5). Hydrogen bonds are represented by dotted lines. In reaction steps (2–5) the protein environment is shown in a simplified manner as a grey curved line.



anion and the triplet-state O₂ to be overcome.^{18,19} According to one QM/MM study the rate limiting step of the reaction is the addition of O₂ to C2 of QND leading to the formation of peroxide (steps 2–3, Fig. 1B) that has been suggested to occur on the triplet-state potential energy surface with a 17 kcal mol⁻¹ barrier, followed by an intersystem crossing leading to a singlet state.¹⁸ Another study instead suggested that the triplet-state peroxide intermediate is unlikely to play a role in the mechanism and that catalysis could proceed *via* a direct electron transfer from the QND anion to O₂, followed by radical recombination yielding the peroxide intermediate.¹⁹ Once the peroxide **3** is formed, subsequent ring closure (step 4) followed by release of CO and formation of the product (step 5) are uncontroversial.

The most elusive and interesting aspects of cofactor-independent oxygenation at the ABH-fold are those that involve molecular oxygen for which no direct structural information is currently available. In this work, we have addressed this gap using an integrated computational and experimental framework. This approach has allowed a complete understanding of the reaction mechanism to be obtained, providing a rationale for how this versatile protein architecture and its simple catalytic machinery tuned for hydrolytic reactions has been successfully redeployed during evolution to carry out O₂-dependent catalysis.

Results and discussion

HOD displays an O₂ pocket that partly overlaps with that of the organic substrate.

To identify possible dioxygen pockets within HOD we initially performed molecular dynamics (MD) simulations. Two independent replica simulations, each 1 μs-long, were carried out for water-solvated HOD in the presence of dioxygen (ten O₂ molecules in a box of approximately 74 × 74 × 74 Å³). Identical simulations were also performed without explicit dioxygen. Hereafter, we will refer to simulations with and without explicit O₂ as OXY-MD and WAT-MD, respectively. In all runs, we observed only minor deviations from the starting crystallographic model (PDB code 2WM2) as expected from the stable ABH fold (Fig. S3†). In OXY-MD runs, we sampled O₂ positions every 50 ps by mapping the locations most frequently visited sites on this time scale (Fig. 2A). These tend to be confined to the core domain and are typically transient pockets afforded by sidechain movements. Amongst the top sites there is however also a portion of the active site (B-site in Fig. 2A) that is substantially more accessible. This location overlaps with the binding site of the natural substrate, specifically that of its hydrophobic B-ring portion (Fig. 1A). We also compared root-mean-square fluctuations (r.m.s.f.) of the C α coordinates from the average structures in OXY-MD and WAT-MD runs (Fig. S4†), a measure of the structural changes during the simulations. This suggests that O₂ tends to reduce the mobility of the β 5- α C ‘nucleophilic elbow’ centered at S101, the portion of the cap domain overhanging it, and the β 8- α F’ C-terminal loop hosting the catalytic H251 residue. These regions (highlighted in yellow in Fig. 2A) define the interface between the core and cap

domains where the active site is located. The basin-shaped portion of the active site that is expected to host O₂ for its attack on the activated substrate (Fig. 2B) is not amongst the most frequented locations on the 50 ps time scale. This O₂ ‘reactive-site’ (R-site) was identified in the crystal structure of the anaerobic HOD–QND complex as a ~15 Å³ cavity opening in front of S101 underneath the substrate’s A-ring⁴ and corresponds topologically to the ‘oxyanion hole’ that stabilizes the negatively charged tetrahedral transition state in hydrolytic reactions.^{7,20}

We next employed xenon pressurization in the crystal state to probe O₂ sites experimentally. Electron-rich Xe is quite easily detected by crystallographic methods and has been successfully used to visualize hydrophobic O₂ binding sites in proteins.^{21–23} Pressurization of HOD crystals in a quartz capillary at room-temperature under a constant 30 bar Xe atmosphere allowed us to measure X-ray data at 2.9 Å using synchrotron radiation (Table S1†). Fourier difference density maps revealed strong peaks consistent with xenon binding at the B-site (as it overlaps with the substrate’s B-ring in the HOD–QND complex) of all four HOD molecules present in the asymmetric unit (Fig. 2C). Modelling these peaks as water molecules resulted in significant positive residual density post-refinement, supporting the notion that Xe occupies this site (Fig. S5A†). Anomalous maps are also consistent with this assignment (Fig. S5B†). Occupancy refinement of Xe atoms gives values in the 40–60% range. Xe is stabilized at the B-site by the hydrophobic environment of the H102, F136, L140, L143, L156, W160, W185, I192 side chains with the closest atoms 3.8–5.0 Å from the gas atom (Fig. 2C). Although we performed our experiment at room temperature, which compared to cryo-conditions, allows for enhanced protein mobility we have not identified additional Xe binding sites in our maps.

Overall, experiments and simulations agree that the ABH-fold HOD dioxygenase features an O₂ pocket (B-site) within the active site at a location that overlaps with the most hydrophobic portion of its aromatic substrate.

S101 at the nucleophilic elbow modulates O₂/H₂O stability at the R-site

HOD’s active site shape suggests that O₂ must be located at the R-site to react with the bound substrate (Fig. 2B). Unlike the hydrophobic B-site, the R-site is polar and crystallographic studies have shown that a water molecule is often bound at this position with variable occupancy.^{4,16} In the presence of NaCl at high concentration the R-site also stabilizes a chloride ion.⁴ Halide and dioxygen binding sites have been shown to be shared in other O₂-dependent enzymes.^{24–27}

As dioxygen must be in strong competition with H₂O for the R-site and considering the proximity of S101, we wondered whether this residue might play a role in modulating the O₂/H₂O preference for this location. Serine residues are characterized by three main rotational conformers (rotamers) for their (N-CA-CB-OG1) χ 1 torsion angle defined as *plus* (χ 1 = +60°), *trans* (χ 1 = 180°), and *minus* (χ 1 = -60°). Statistical analysis shows that *trans* and *plus* are the least and most frequently



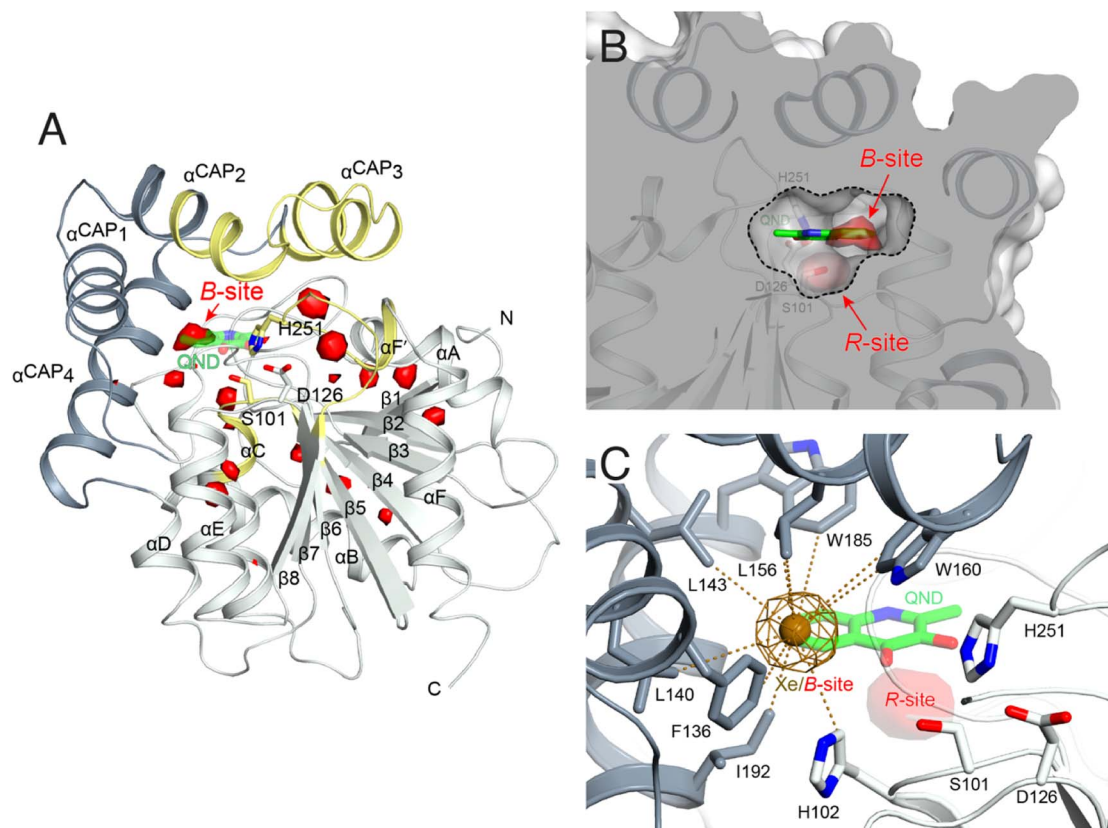


Fig. 2 Dioxygen binding sites mapped by MD simulations and xenon pressurization. (A) Cartoon representation of HOD (core and cap domains in light and dark grey, respectively) with high-probability O_2 -sites (isosurface representation in red) as revealed by MD simulations. Structural regions whose dynamics is reduced by dioxygen are highlighted in yellow. Most O_2 -sites are temporary pockets generated by protein dynamics within the core domain. One site (B-site) is much more accessible and overlaps with the location occupied by the B-ring of the QND substrate in the E–S complex. The latter is shown for reference as a stick representation with carbon, nitrogen and oxygen atoms colored green, blue and red, respectively. Residues of the catalytic triad are also shown as sticks; (B) sliced-surface back-view (roughly rotated by 180° around the vertical axis compared to the view in A) showing a cut-through of the active site cavity in the E–S complex (PDB code 2WJ4) with its largest section highlighted by a dotted line. The B-site lies within the flat horizontal portion of the active site that hosts the QND substrate. Underneath the substrate's A-ring heterocycle, the active site is shaped into a basin. The R-site (R for reactive) that measures approximately 15 \AA^3 , positioned in front of the S101 sidechain, is expected to host O_2 during catalysis; (C) active site of HOD following xenon pressurization. A Xe atom, shown as a gold sphere with its $2mF_o - DF_c$ map at the $+1.0\sigma$ level as a chicken-wire representation, localizes at the B-site. Stabilizing sidechains are shown as sticks with atoms within 5.0 \AA of Xe highlighted by dotted lines. Residues of the catalytic triad are also shown. All residues are colored according to the domain to which they belong. The bound QND substrate and the R-site underneath it are shown for reference.

observed, respectively, particularly when this residue is not part of either α -helices or β -strands.²⁸ We analyzed S101 rotamer preferences in our $1 \mu\text{s}$ -long substrate-free WAT-MD and OXY-MD simulations and found that O_2 shifts the distribution from bimodal, in which both *plus* (S101^P) and *trans* (S101^T) rotamers are sampled with equal frequency (Fig. 3A), to unimodal in which S101^T dominates (Fig. 3B). The *minus* rotamer is never frequently populated. Structurally, the transition from S101^P to S101^T causes the CB-OG bond to reorient itself, switching from the direction pointing directly toward the R-site to one 'running' tangentially to it following the backbone (Fig. 3 inset).

Next, we turned our attention to the enzyme–substrate (E–S) complex. As expected, a $1 \mu\text{s}$ -long MD run starting from the crystallographic HOD–QND structure⁴ revealed only limited deviations from the experimental model (Fig. S6†). To assess if a possible correlation exists between the S101 rotamer

preference and O_2 stability at the R-site we then performed multiple independent simulations in which a single O_2 molecule was positioned at this location underneath the substrate and monitored its residence times with S101 restrained either to its *trans* or *plus* rotamer. A total of 50 simulations were carried out for each rotamer. We find that the *trans* rotamer stabilizes O_2 at the R-site better than the *plus* rotamer with mean residence times of $2.61 \pm 0.34 \text{ ns}$ and $0.55 \pm 0.08 \text{ ns}$, respectively (Fig. 3C). Also, a wide time distribution is observed when S101 is restrained to the *trans* rotamer, including times up to 8 ns, whilst times are short and tightly distributed for the *plus* rotamer. Alanine substitution (S101A) results in a behavior like S101^T (mean \pm s.e.m. of $3.95 \pm 0.40 \text{ ns}$) albeit with generally longer residence times. We have also estimated residence times (k_{off}^{-1}) from the numerical fitting of the cumulative time distribution of individual events that is expected to follow



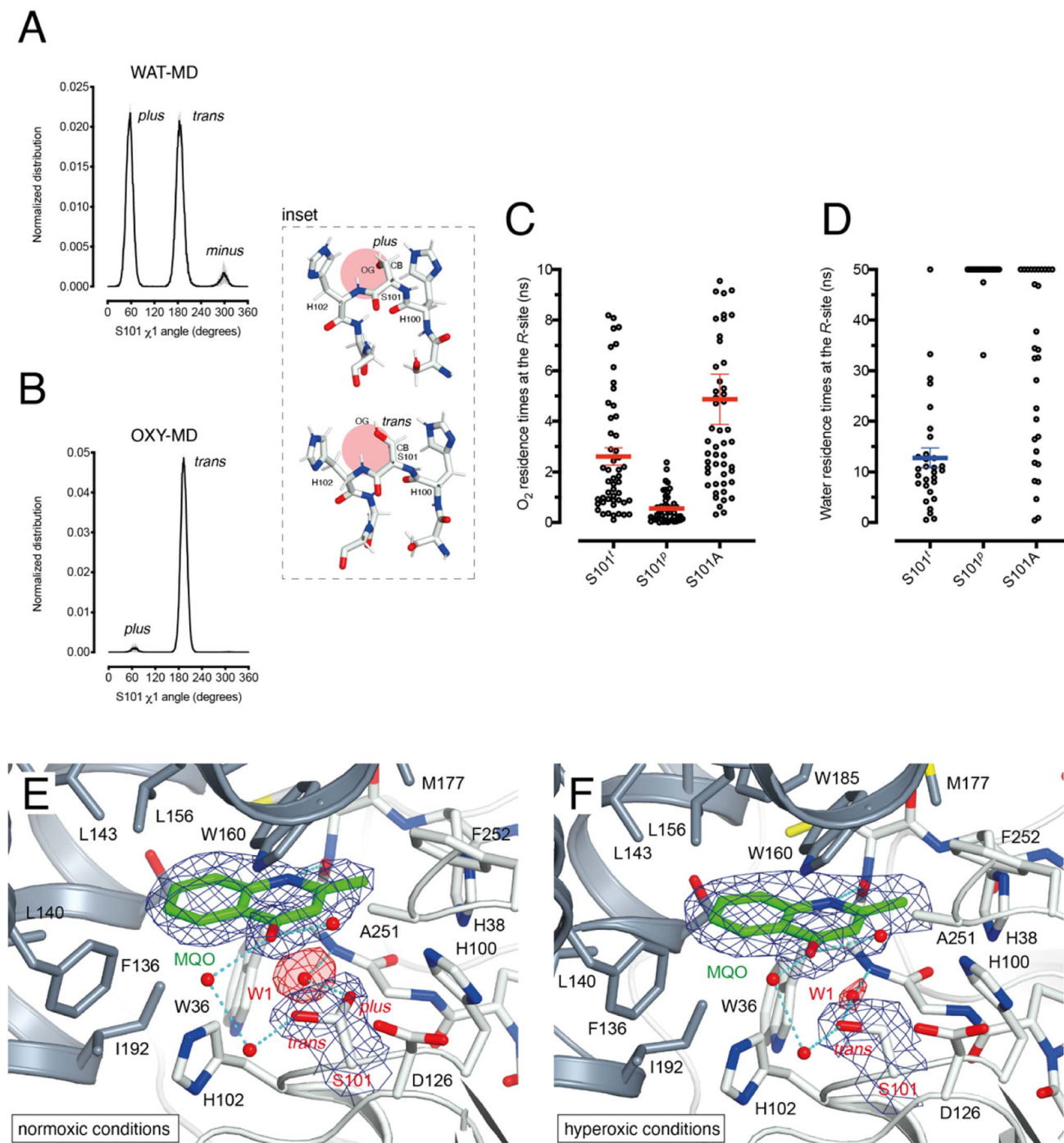


Fig. 3 S101 as the O_2/H_2O modulator at the R-site. (A) Normalized distribution of χ_1 (N-CA-CB-OG) angles for S101 in MD simulations carried out for HOD without explicit dioxygen. S101 rotamers are defined as: *plus* ($\chi_1 = +60^\circ$), *trans* ($\chi_1 = 180^\circ$), and *minus* ($\chi_1 = -60^\circ = 300^\circ$). Values are the average of two independent MD replicas each 1 μ s-long sampled every 50 ps. Standard deviation is shown as light grey bars; (B) same as A in the presence of explicit dioxygen. The inset provides snapshots during the simulations with the R-site represented by the red circle. The CB-OG bond lines the R-site for the *trans* rotamer whilst it roughly points into it for the *plus* rotamer; (C) distribution of residence times for an O_2 molecule at the R-site of the HOD-QND complex with S101 restrained to the *trans* (S101^t) and *plus* (S101^p) rotamers or with S101 replaced by an alanine (S101A). A total of 50 independent MD simulations were performed for each system. Mean and s.e.m. values are highlighted; (D) like C but for a water molecule at the R-site. A total of 30 independent MD simulations were performed for each system with a cutoff time of 50 ns. For S101^t, S101^p, and S101A water was observed leaving the R-site within the cutoff (29/30), (2/30), and (20/30) times, respectively. Mean and s.e.m. values for S101^t are highlighted; (E and F) active site of the HOD^{H251A}-MQO complex under normoxic (E) or hyperoxic (F) conditions. Electron density maps ($2mF_o - DF_c$) are shown at the $+1.0\sigma$ level as a chicken-wire representation for MQO, and the water molecule at the R-site (W1) and S101. The latter displays a mixture of *plus* (40%)/*trans* (60%) rotamers under normoxic conditions whilst only the *trans* rotamer is observed under hyperoxic conditions. Electron density for W1 is displayed in red for clarity. The H-bond network involving MQO, W1 and S101 is shown as cyan dotted lines.



Poisson statistics. This procedure gives similar $k_{\text{off}}^{-1}(\text{O}_2)$ values (Fig. S7†).

Next, we carried out 30 more independent simulations and performed a similar analysis for a H_2O molecule located at the R-site (Fig. 3D). As expected, water is retained at this location longer than O_2 . However, whilst for *plus*-restrained S101, water was observed to leave the R-site only twice within a cutoff of 50 ns (6.7% escapes, with individual residence times of 33.1 and 47.5 ns), in the case of *trans*-restrained S101 water stability is significantly reduced (96.6% escapes, mean \pm s.e.m. of 13 ± 2 ns). In the case of the S101A substitution, H_2O stability is somewhat intermediate between the behavior promoted by the two S101 rotamers (67% escapes) with a large time distribution. Numerical fitting gives a $k_{\text{off}}^{-1}(\text{H}_2\text{O})$ value of approximately 15 ns for S101^t (Fig. S7†), while it is greater than 50 ns for both S101^l and S101A.

Overall, our analysis suggests a role for S101 as a modulator of $\text{O}_2/\text{H}_2\text{O}$ stability at the R-site. Transient O_2 binding within the HOD architecture favors the switch to the *trans* rotamer which, in turn, promotes both O_2 stabilization and H_2O destabilization at the R-site. The S101A replacement at the nucleophilic elbow appears to be a useful strategy to further stabilize O_2 at the R-site at the cost, however, of less effective water clearing.

Normoxic and hyperoxic HOD^{H251A}-MQO structures validate S101 as an $\text{O}_2/\text{H}_2\text{O}$ modulator.

To visualize O_2 at the R-site, we turned to pressurization experiments in the crystal state using the 'soak-and-freeze' method that allows crystal cryocooling under pressure.²⁹ This is a significant advantage over methods that require pressure release before cryocooling as it minimizes the escape of gas molecules that often have high k_{off} rates.

Initially, we employed the near-inactive HOD^{H251A} variant that under normoxic conditions affords the visualization of a stable HOD^{H251A}-QND complex.¹⁶ However, O_2 -pressurization of the complex (40 bar O_2 for 2 minutes) leads to the complete conversion of the bound QND into the product (see the ESI and Fig. S8†). This clearly indicates that in the crystal state under hyperoxic conditions, O_2 can reach the R-site in the preformed E-S complex. To prevent turnover, we next synthesized the substrate analogue 2-methyl-quinolin-4(1H)-one (MQO). MQO, a non-reactive molecule structurally closest to the natural substrate, in which the hydroxyl group at position 3 is replaced by the -H substituent. As crystals of HOD^{H251A} typically diffract better than those of wild-type HOD we solved the X-ray structure of this variant in complex MQO under normoxic and hyperoxic conditions at the 2.0 Å and 2.1 Å, resolution, respectively (Table S1†). MQO binds in the active site similarly to QND indicating that the lack of the 3OH group does not result in substantial changes. MQO is held in place by a single H-bond between its NH group and the carbonyl oxygen of W36 at 2.8 Å. Several residues contributed both by the core domain (Gly35, Trp36, Cys37, His38, His100, Ser101, His102, Gly103, and Phe252) and the cap domain (Phe136, Leu140, Leu143, Leu156, Trp160, Met177, Trp185, Ser188, Gly189, and Ile192) further stabilize the ligand with hydrophobic interactions (Fig. 3E). A water

molecule (W1) is present at full occupancy at the R-site underneath the A-ring of MQO 2.8 Å from its molecular plane. It is stabilized by an interaction with the main chain amide of W36 and the side chain of S101 which is observed in double conformation in both molecules present in the a.u. with *trans* and *plus* rotamers refining at occupancies of 0.60 and 0.40, respectively. W1 is H-bonded to S101^p 2.7 Å from its OG atom, whilst S101^l (OG) is more than 3.8 Å away.

We next inspected electron density maps following the O_2 -pressurization experiment (40 bar O_2 for 2 minutes) (Fig. 3F). These do not reveal changes that we could positively ascribe to O_2 bound at the R-site. Instead of the elongated electron-rich density observed for bound dioxygen in other systems,^{29,30} O_2 -pressurization led to a decrease in electron density at the R-site compared to that under normoxic conditions. Occupancy refinement of a water molecule at this location gives values of 0.67 and 0.48 in the two independent molecules present in the a.u. Remarkably, S101 shifts completely to the *trans* rotamer (χ^2 values of 180.08 and 180.66°) with no indication of the alternative *plus* rotamer observed under normoxic conditions.

Overall, these observations indicate that O_2 most likely interferes with H_2O binding at the R-site, and although disorder negates its positive identification, this nevertheless leads to the shift of S101 to the *trans* rotamer that, in agreement with the simulations, appears to be positively correlated with the presence of dioxygen at the R-site.

O_2 access to the R-site in the pre-formed E-S complex

As *in crystallo* O_2 pressurization promoted turnover of the HOD^{H251A}-QND complex we again employed MD simulations to gain an understanding of possible access O_2 routes to the R-site in the pre-formed E-S complex. We performed a total of 80 independent 200 ns standard MD runs either in the presence of ten O_2 molecules placed in a box of approximately $74 \times 74 \times 74$ Å³ (OXY10-S-MD runs) or using a single O_2 molecule placed initially near W37 and constrained within a sphere of 17 Å radius from S101 (OXY1-S-MD runs). The latter condition decreases the sampling space while still allowing O_2 to diffuse outside the protein.

The simulations reveal that O_2 entry at the R-site is not a frequent event. Out of the 80 standard MD simulations we observed spontaneous O_2 entry at the R-site only once in either OXY10-S-MD or OXY1-S-MD runs. These events occurred when S101 was restrained to its *trans* rotamer. O_2 access to the R-site followed the same trajectory in both productive runs (Fig. 4A). Dioxygen entered the ABH-fold near P132 (cluster I in Fig. 4A) and reached the R-site (cluster II) taking advantage of a hydrophobic path lined by residues belonging to α_{CAP1} (F136 and L140) and α_{CAP3} (I192 and G189). O_2 then escaped from the R-site highlighting clusters III and IV near the nucleophilic elbow. These were recurrently visited prior to reaching clusters V and VI close to W36 and V71, respectively. Cluster V matches very well a high-probability O_2 pocket seen in MD simulations in the absence of the substrate.

In a second set of calculations, we employed biased metadynamics simulations with a single O_2 molecule placed in the



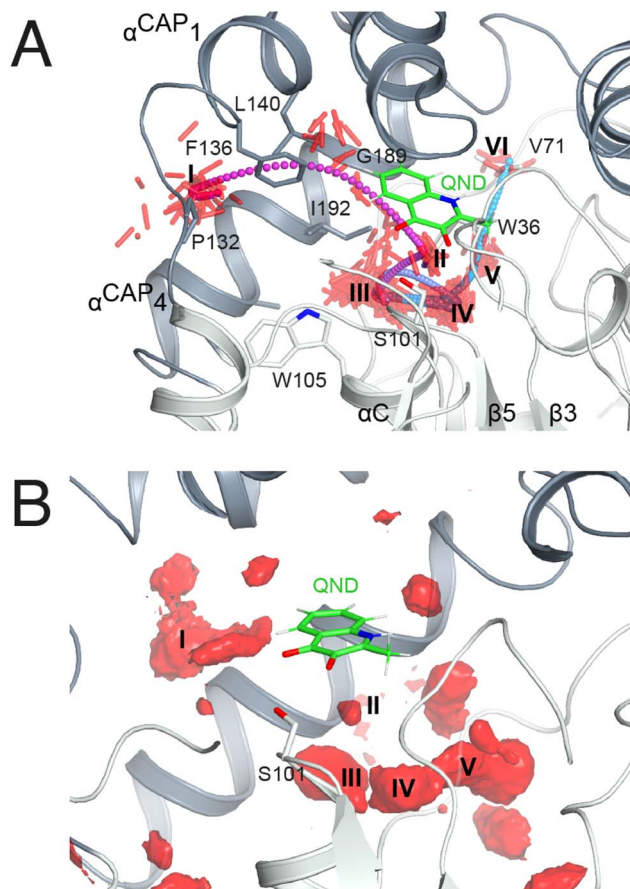


Fig. 4 O_2 access to the R-site in the E-S complex. (A) Dioxxygen trajectory extracted from a standard MD simulation. Roman numbers indicate clusters of dioxxygen molecules, shown as red sticks, sampled every 50 ps. The O_2 trajectory, highlighted by small spheres colored using a cyan-magenta gradient (entry-exit), shows that dioxxygen reaches the R-site (cluster II) from cluster I, and exits the ABH-fold via clusters (III-VI) as described in the main text. (B) Regions of high O_2 occupancy probability (isosurface representation in red) obtained reconstructing the free energy in the Cartesian space from metadynamics simulations. Roman numbers indicate the same clusters of panel A.

bulk and free to visit any location inside the box (OXY1-S-metaD runs). The use of a bias (see the Methods section) allows the sampling of the entire 3D Cartesian space with simulations restricted to 1.8 μ s each to be accelerated. OXY1-S-metaD runs highlight the same clusters I-V seen in the standard simulations in addition to a few additional clusters (Fig. 4B). Energy calculations confirm a lower barrier for O_2 entry when S101 is in the *trans* rotamer (4.1 ± 1.1 kcal mol⁻¹) compared to the *plus* rotamer (6.5 ± 1.5 kcal mol⁻¹). This corresponds roughly to a five-fold increased probability in the case of the former.

Visualization of O_2 at the R-site

As the simulations suggested that the S101A substitution decreases $k_{\text{off}}(O_2)$, we then solved the structure of HOD^{S101A} in a complex with MQO under normoxic and hyperoxic conditions at 2.0 Å resolution (Table S1†). The HOD^{S101A} variant was also

considered interesting as the AqdC ABH-fold dioxxygenase that catalyzes a chemical reaction identical to HOD naturally possesses an alanine instead of a serine at the nucleophilic elbow (hydrophobic elbow).

MQO binds to HOD^{S101A} as observed with HOD^{H251A}. Although the S101A replacement causes the loss of the H-bond between S101(OG) and MQO(O4) this is compensated by the interaction with H251(NE2) resulting in an essentially identical binding mode. Under normoxic conditions, electron density at the R-site of HOD^{S101A} is consistent with the presence of a water molecule (Fig. S9†). However, in contrast with what was observed for the HOD^{H251A}-MQO complex, following O_2 -pressurization (40 bar O_2 for 2 minutes) Fourier difference maps revealed a strong elongated peak at the R-site of one of the two HOD^{S101A} chains in the asymmetric unit (Fig. 5A). This peak ranked third overall for height ($+7.3\sigma$) with the first two corresponding to sodium ions. Modelling of this peak as an O_2 molecule did not result in negative difference density, and occupancy refinement converged to unity with atomic displacement parameters for both oxygen atoms consistent with those of the surrounding atoms (average B -value MQO = 21.3 Å², B -value $O_2(O1)$ = 20.5 Å² and B -value $O_2(O2)$ = 19.7 Å²). Moreover, omit maps for the individual oxygen atoms produced difference Fourier peaks, supporting the assignment as O_2 (insets of Fig. 5A). The equivalent peak in the other protein chain was less prominent and more spherical in shape. We have modelled this peak also as O_2 and occupancy refinement converged at 0.80 with an average B -value of 30.3 Å². This value is marginally higher than that of the MQO molecule in its proximity (average B -value = 21.9 Å²) suggesting higher rotational disorder.

Using the fully occupied O_2 molecule as a reference, we find that dioxxygen binds underneath the A-ring of MQO approximately parallel to its plane at an average distance of 3.2 Å (Fig. 5B). The ligand's atoms closest to $O_2(O2)$ and $O_2(O1)$ are C2 at 3.1 Å and C4 at 3.4 Å, respectively, and dioxxygen further interacts with the main chain amide groups of W36 and H102 as well as with the side chain of the latter residue. Next to the R-site, a water molecule (W3) is also present at a H-bond distance from O1 (2.7 Å) stabilized by the side chain of H100 that is flipped compared to the HOD^{H251A} variant. W3 and H100 flipping are also seen under normoxic conditions in two of the four molecules present in the a.u. The O_2 molecular axis is roughly parallel to the MQO(C2-C4) direction as quantified by the dihedral angle $\phi(C4-C2-O2-O1)$ of -2.2° . Thus, in the HOD^{S101A}-MQO- O_2 ternary complex, dioxxygen appears to preferentially adopt an orientation that mimics that of the endoperoxide during the catalytic cycle (step 4 in Fig. 1B).

The geometry and electronic properties of the ligand- O_2 complex depend on the ligand's charge.

To seek further insight into the geometry and electronic properties of the QND- O_2 and MQO- O_2 complexes, we performed *ab initio* quantum mechanics (QM) calculations. Initially, we considered two systems constituted only by the ligand (either QND or MQO) and O_2 . As QND is deprotonated by the His-Asp



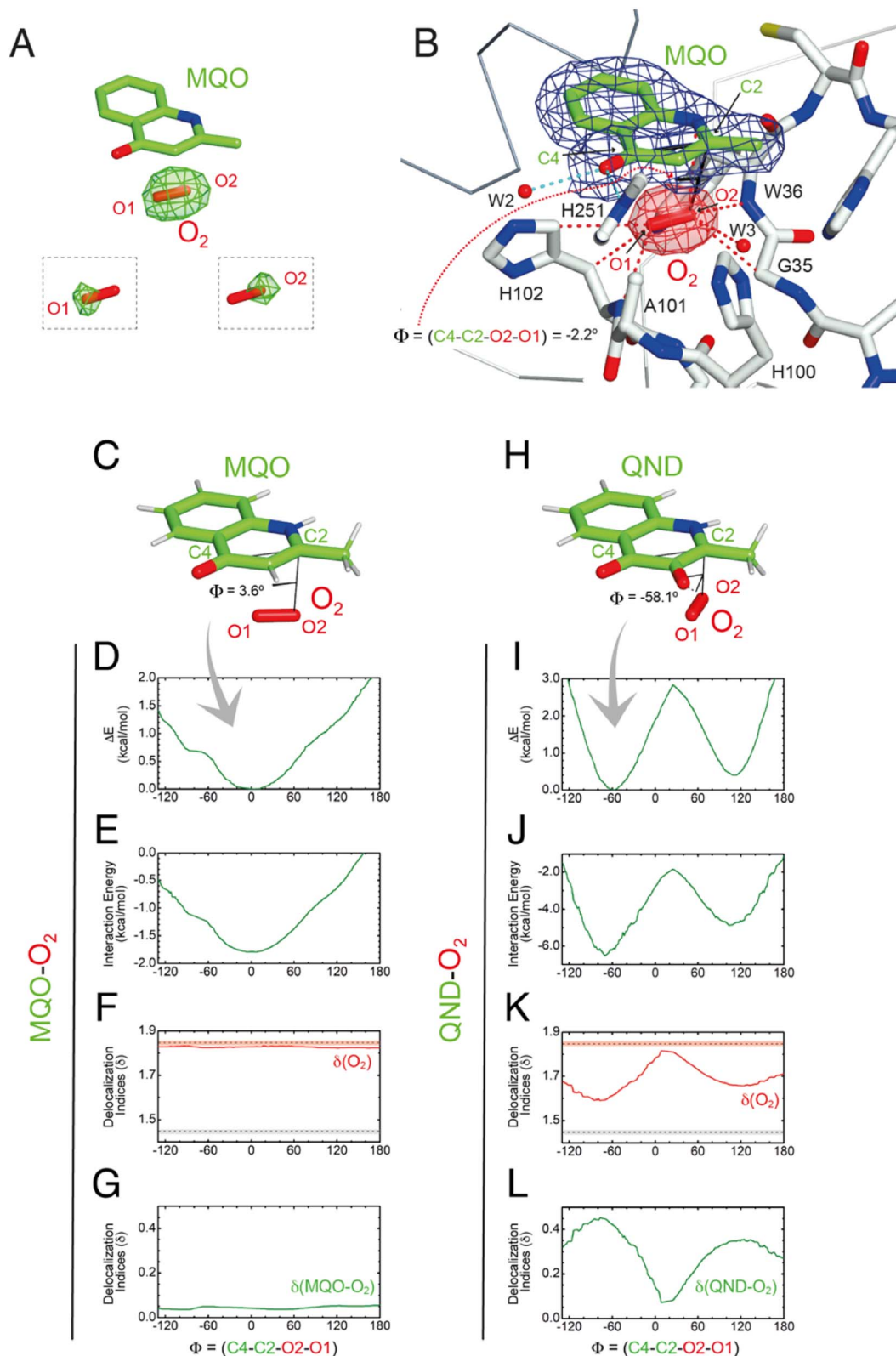


Fig. 5 Visualization of dioxygen at the R-site and QM validation. (A) Following O₂ pressurization (40 bar for 2 minutes) of the HOD^{S101A}-MQO complex, mF_o-DF_c Fourier difference maps at 2.0 Å-resolution (in green at the $+3.0\sigma$ contour level) reveal a strong elongated peak at the R-site underneath the MQO ligand that is consistent with dioxygen. O₂ from the refined model is shown for reference as a red stick. The insets show the mF_o-DF_c electron density map ($+3\sigma$) with O1 and O2 atoms selectively removed from crystallographic refinement. (B) $2mF_o-DF_c$ electron density of the HOD^{S101A}-MQO-O₂ complex. Electron density maps are shown at the $+1.0\sigma$ level as a chicken-wire representation for MQO (blue) and O₂ at the R-site. The latter is displayed in red for clarity. The dihedral angle $\phi = (C4-C2-O2-O1)$ in the refined crystallographic model is -2.2° . (C-L) QM geometry restrained optimization and delocalization indices (δ) for the MQO-O₂ (C-G) and QND-O₂ (H-L) complexes as a function



dyad, we assumed that the QND–O₂ system bears a single negative charge whilst MQO–O₂ is neutral. In these calculations, we restrained the sum of the (C2–O1) and (C4–O2) distances to the crystallographic values. However, around the equilibrium geometry, our results are insensitive to small changes in the restraint value (Fig. S10†).

The total energy of the complexes as a function of Φ is shown in Fig. 5D and I. Despite the structural similarity between QND and MQO, the QM calculations revealed that the orientation of O₂ with respect to the ligand is different. Whilst in the MQO–O₂ complex, dioxygen orients its axis along the MQO(C2–C4) direction with a dihedral angle Φ of 3.6° (Fig. 5C) in excellent agreement with our crystallographic structure, in the QND–O₂ complex, O₂ is aligned with the QND(N1–C3) bond at an angle of about –58.1° (Fig. 5H). We observe that an essentially identical angular dependence is observed for the “interaction energy”-only component of the total energy (Fig. 5E and J), indicating that molecular distortions are not important in defining the equilibrium geometry of these systems. Energy decomposition analysis allowed the further investigation of the contribution of the different terms (Pauli repulsion, charge transfer, electrostatic, dispersion, and polarization) to the interaction energy (Fig. S11†).^{31,32} For the MQO–O₂ complex, the most stable orientation is that which minimizes the Pauli repulsive energy between the electronic clouds of MQO and O₂, that is, in this case, associated with π – π stacking. Differently, for the QND–O₂ complex, the equilibrium geometry is that which maximizes the charge-transfer from the QND anion to O₂. We hypothesized that the difference in the relative O₂ orientation between the two complexes is caused by the negative charge of QND. This was confirmed by calculations for the protonated (neutral) QNDH–O₂ complex, that closely mirror those obtained for MQO–O₂ (Fig. S12†).

Next, we analysed the electron density shared between the ligand and O₂ using delocalization indices (δ), which are intimately related to bond order.³³ We calculated these values as a function of Φ for dioxygen $\delta(\text{O1–O2})$, and for the sum of all the pair- δ values between the ligand and O₂, $\delta(\text{QND–O}_2)$ and $\delta(\text{MQO–O}_2)$. For the MQO–O₂ complex, we find that independently of Φ , $\delta(\text{MQO–O}_2)$ is close to zero whilst $\delta(\text{O1–O2})$ is close to the value for isolated O₂, $\delta(\text{O1–O2}) = 1.84$ (Fig. 5F and K). This indicates that there is essentially no electron density transfer between MQO and O₂. For the QND–O₂ complex, both $\delta(\text{QND–O}_2)$ and $\delta(\text{O1–O2})$ are no longer independent of Φ . Moreover, at the equilibrium geometry, where charge-transfer is maximized, $\delta(\text{QND–O}_2)$ has a maximum at ~ 0.35 while $\delta(\text{O1–O2})$ has a minimum at ~ 1.67 , which is intermediate between the value for isolated O₂ and that of an aromatic C–C bond (Fig. 5G and L). In this complex therefore electron density is shared between QND and O₂ that results from the electron transfer from the π cloud of QND to the π^* orbital of O₂, decreasing its bond strength.

To test the effect of the protein matrix we have repeated the same analysis above using a QM/MM model in which the QM region includes the substrate, O₂ and key active site residues (Fig. S13A and E†). In these calculations that explicitly include the active site environment, no restraints were employed. These more sophisticated calculations do not alter our previous conclusions and the results are entirely consistent with that obtained for the isolated complexes (Fig. S13B and F†). We note, however, that for MQO–O₂, the energy difference associated with O₂ rotation is smaller, with a barrier height lower than 0.5 kcal mol^{–1}, which could be easily overcome at thermal energies. This flat energy profile therefore allows rotational averaging which likely contributes to an increase in the sphericity of O₂ electron density at the R-site as observed in one of the two active sites. For the QND–O₂ complex, the barrier is about 2 kcal mol^{–1}, like that obtained from the isolated complex. Analysis of the delocalization indices also produced results like those obtained for the isolated complexes (Fig. S13C, D, G and H†).

Reaction mechanism

To investigate the reaction mechanism, we calculated the QM/MM energy profile along the reaction coordinate α (Fig. 6A). The reaction can be conveniently rationalized as composed of two stages: (i) an intersystem crossing (ISC) stage in which the system ‘hops’ from the triplet to the singlet state with the formation of C2-peroxide (2) and (ii) the CO-release stage, in which the peroxide breaks down *via* endoperoxide (3 and 4) leading to products (5). For the ISC stage, the energy profile was calculated as a function of the energy difference between the singlet and triplet states (Fig. S14†) and then represented as a function of the reaction coordinate α as shown in Fig. 6A.

Reactants (1) are stable in the triplet state arising from the combination of the electronic ground state of O₂(³ Σ_g^-), and the ground state of QND (singlet, all electrons are paired). However, as QND and O₂ move closer to each other, the energy of the system increases and any attempt to optimize C2-peroxide in the triplet state led back to the reactants. In contrast, formation of the peroxide in the singlet state proceeds barrierless leading to a stable moiety. Analysis of the singly occupied molecular orbitals (Fig. S15†) suggests that stabilization and destabilization of the singlet state and of the triplet state, respectively, originate from the combination of one π^* orbital of O₂ with the π cloud of QND. This leads to an orbital with antibonding character along the C2–O2 and O1–O2 bonds, which is singly occupied in the triplet state and unoccupied in the singlet state. Singlet and triplet states must cross at some point along the reaction path and the minimum energy crossing point (MECP, highlighted by a black dot in Fig. 6A) acts as the effective transition state. Our calculations locate the MECP (2) at 10.2 kcal mol^{–1} above the energy of the activated QND–O₂ complex (1). At the MECP, $d(\text{C2–O2})$ is 1.62 Å, and QND is no longer

of the dihedral angle Φ . (C and H) Stick representations of the minimized structures, (D and I) total energy, (E and J) interaction energy-only, (F and K) δ for O₂, and (G) sum of the pair- δ between the ligand and O₂ for the MQO–O₂ and the QND–O₂ complexes, respectively. In (H and J) δ values for isolated O₂ (dashed red line) and the aromatic C–C bond in benzene (dashed grey line) are given for reference. Experiment and theory are in excellent agreement that the MQO–O₂ complex displays an energy minimum close to $\Phi = 0^\circ$.



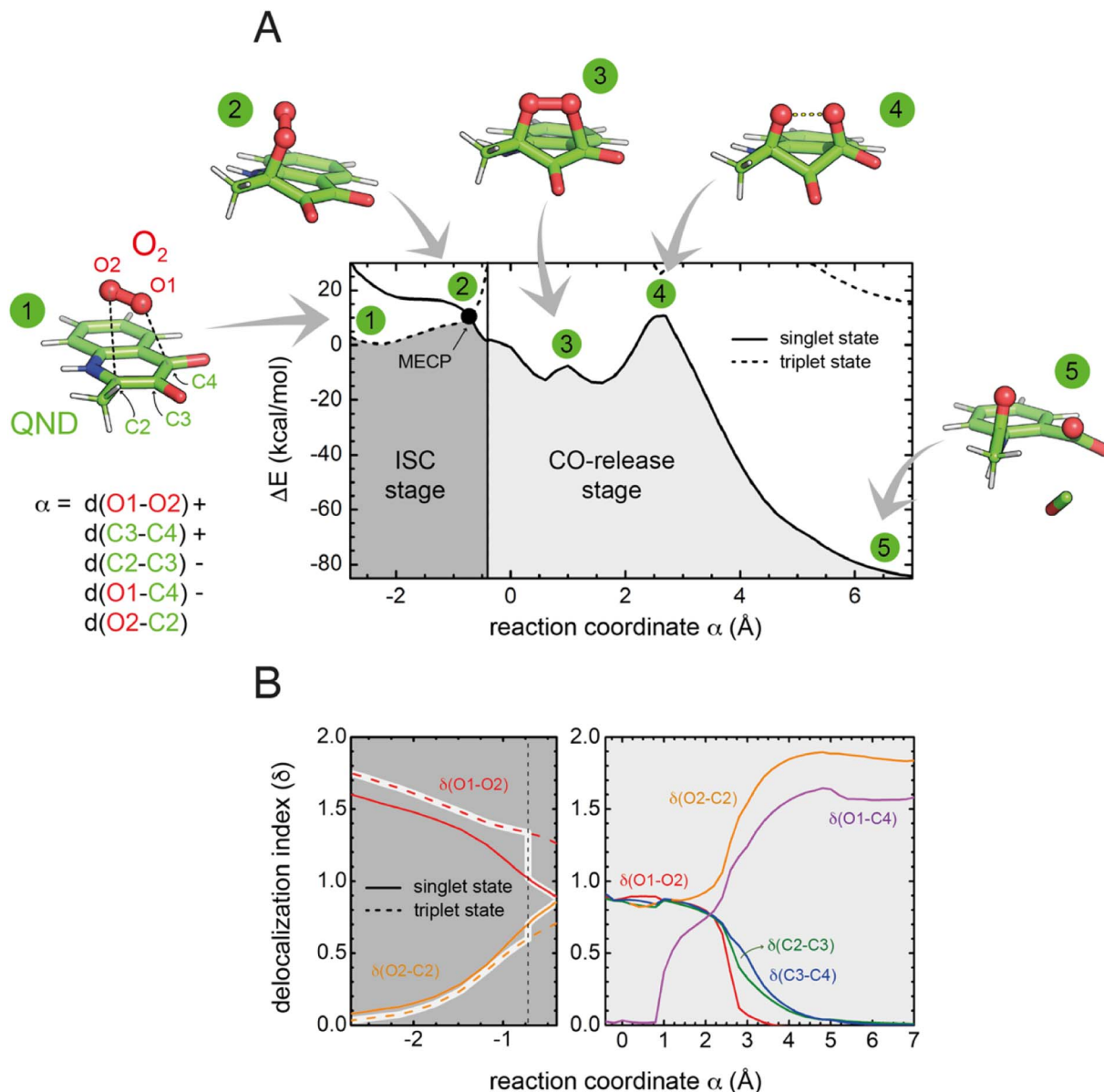


Fig. 6 Energetics and charge transfer along the reaction path. (A) Energy profile calculated at the M06-2X/6-31+G(d,p) level of theory as a function of the reaction coordinate α . The lowest energy singlet and triplet states are shown as continuous and dashed lines, respectively. Numbered key stages of the reaction are visualized as ball-and-stick representations. In all of them, the protein is not drawn for simplicity. The vertical line (at $\alpha = -0.4$) distinguishes the two reaction stages. The left-hand region defines the intersystem crossing (ISC) stage, in which the system switches from the triplet to the singlet state leading to the formation of the peroxide through the minimum energy crossing point (MECP, highlighted by a black dot). The CO-release stage on the singlet potential energy surface is strongly exothermic. (B) Evolution of the delocalization indices (δ). The left panel shows δ indices for the ISC stage where only $\delta(\text{O1-O2})$ and $\delta(\text{O2-C2})$ change significantly. The DIs are shown for the triplet (dashed lines) and singlet (solid line) states, and those corresponding to the most stable state are highlighted. The vertical dashed line represents the position of the MECP. Differences between δ values for the singlet and triplet states at the MECP indicate that ISC is associated with a significant transfer of electronic density from the substrate to O_2 . In the right-hand side panel, δ indices are shown only for the singlet state, as the triplet state is very high in energy.

planar. To get further insight into the ISC stage, we evaluated how $\delta(\text{O2-C2})$ and $\delta(\text{O1-O2})$ change along the reaction path in both the singlet and triplet states (left-hand panel in Fig. 6B). We observed that shortening of $d(\text{C2-O2})$, results in an increase in the electronic density shared between the substrate and dioxygen, which leads to the weakening of the O_2 bond, a consequence of the larger charge transfer from QND to O_2 .

Moreover, at every point along the reaction path, charge transfer is stronger in the singlet than in the triplet state, so the spin-flip involves a significant transfer of electronic density between QND and O_2 .

Once the peroxide is formed, the next stage of the reaction proceeds *via* the formation of endoperoxide (3) and subsequent rupture of the C2-C3 and C3-C4 bonds, followed by CO release



that is very exothermic. The process is again well described by the delocalization indices (right-hand panel in Fig. 6B). Once the system is in the singlet state, the peroxide attacks the carbonyl group establishing the C4–O1 bond of endoperoxide. At the barrier (4), concerted breaking of endoperoxide and CO release takes place. As shown in Fig. 6B, the energy of the transition state for the CO-release stage lies 10.8 kcal mol⁻¹ above the reactants, which is slightly above the energy of the MECP. However, to compare the barrier heights it is necessary to correct the ISC energy barrier for the probability of hopping from the triplet to the singlet state,³⁴ which depends on the value of the spin-orbit coupling (SOC). At the MECP, we calculated a SOC of 22 cm⁻¹, leading to a swapping probability of 0.005, *i.e.* an increase of 3.1 kcal mol⁻¹ of the barrier (see the ESI† for further details). Therefore, the ISC stage is the rate limiting step of the overall process.

We also considered the possibility of a direct electron transfer within the complex by calculating the energy difference between [³O₂ + ¹QND⁻] and [²O₂[·] + ²QND[·]] at the reactants' asymptote, based on separate calculations for the donor and the acceptor. We obtained a value of 17 kcal mol⁻¹, which is significantly larger than the energy of the MECP (even when corrected by the swapping probability). This result is compatible with our findings that, along the reaction path, the energy of the singlet state monotonically decreases. Although such an endothermicity does not allow the possibility of a direct electron transfer to be completely ruled out,¹⁹ our calculations predict a scenario where spin-flipping is more favorable. Consistent with this, no significant amounts of radical species were detected in spin-trapping experiments with wild-type HOD.³⁵

Concluding remarks

The “great oxidation event” 2.4 billion years ago led to the permanent accumulation of O₂ in the atmosphere, thus exerting evolutionary pressure on enzyme systems that could take advantage of such a strong oxidant. A very recent KEGG (Kyoto Encyclopedia of Genes and Genomes) analysis of the 136 Pfam families of all known O₂-dependent enzymes, found that only ~60% of these can be classified as having a function primarily related to O₂ whilst the remaining ~40% represent families featuring only sporadic O₂ utilizers.² The latter group has therefore likely evolved O₂-metabolizing capabilities from a different set of original catalytic competences. Hydrolases, particularly metallo-β-lactamases and those of the ABH-fold, appear to be the most common progenitors.² ABH-fold dioxygenases are particularly intriguing as they accomplish O₂ redox chemistry without external cofactors.

Here, using HOD as a paradigm for the growing family of ABH-fold dioxygenases, we have shown how evolution has taken advantage of multiple structural elements of the ABH-fold to enable the switch from hydrolytic to oxygenolytic reactions. A key requirement for such catalytic repositioning is the ability to host O₂ sufficiently close to its *N*-heteroaromatic substrate for the reaction to take place. Using *in crystallo* O₂-pressure freeze-trapping we have visualized the ternary complex with a substrate analog at high resolution, thus providing conclusive

evidence that the oxyanion hole fulfills the role of an O₂ receptor. Computational analyses of dioxygen diffusion pathways in various non-ABH-fold enzyme systems including flavoenzymes,³⁶ the nonheme iron-containing 12/15-lipoxygenase,³⁷ and also the metal-independent DpgC dioxygenase²¹ have uncovered some common themes in O₂ access strategies, often involving a network of transient pathways that are mostly lined with hydrophobic residues. In HOD, we have identified a set of pockets that provide a route for O₂ access to the oxyanion hole in the E–S complex. However, compared to other systems, O₂ entry appears a much rarer event arguing in favor of additional strategies to increase the local O₂ concentration. In urate oxidase, a hydrophobic cavity adjacent to the active site has been suggested to act as a transient O₂ reservoir.³⁸ We propose that in HOD, and likely in all ABH-fold dioxygenases that operate on similar organic substrates, the active site itself acts as a temporary O₂ binding site. Transient O₂ storage at the xenon-validated B-site that we have identified overlapping with the hydrophobic portion of the *N*-heteroaromatic binding site represents a convenient solution to the problem of O₂ localization, as one might envisage a mechanism of displacement that upon substrate binding transfers O₂ from the B-site to the R-site that is only about 5 Å away, thus organizing both reactants for catalysis.

ABH-fold enzymes often employ their typical Ser-His-Asp triad to catalyze reactions following an esterase-type mechanism. Departures from this have however been observed in some non-hydrolytic reactions. *Manihot esculenta* (cassava) and *Hevea brasiliensis* (rubber tree) hydroxynitrile lyases that break the C–C bond in cyanohydrin although featuring the classical Ser-His-Asp triad, they use their active site serine as a general base. Even more dramatic is tomato methyl ketone (MK) synthase (MKS1) involved in MK 2-tridecanone synthesis that lacks the canonical triad whilst retaining a conserved histidine residue that acts as the catalytic base with a neighboring threonine providing the H-bond bond pattern required for the decarboxylation step of β-keto acid substrates.³⁹ For oxygenation, the ABH-fold catalytic machinery is also employed in an unconventional manner and substrates' activation by deprotonation of their 3-hydroxyl group only requires the His-Asp subset (H251-D126 in HOD) that acts as a general base.^{4,16} Although HOD's serine residue (S101) has been shown to retain nucleophilic properties in some hydrolase-like reactions,⁴⁰ this capability is irrelevant for oxygenation.⁴ Instead, our results indicate that in addition to a contribution in the stabilization of the organic reactant,⁴ S101 plays a role as an O₂/H₂O modulator with its *trans* rotamer promoting both O₂ stabilization and H₂O destabilization in the oxyanion hole. Substitution of the cryptic serine nucleophile with an alanine, a replacement naturally present in some ABH-fold dioxygenases that further strengthens the argument of nucleophile dispensability, also promotes O₂ stabilization at the catalytic site. In HOD, S101A replacement, however, has a negative impact on the *K_M* value of its organic substrate (60-fold increase compared to that of wild-type HOD) whilst *k_{cat}* is not affected.⁴ This suggests that the evolutionary choice of the specific amino acid at the ‘nucleophile elbow’ is likely the result of a combination of factors aimed at



maximizing the stability of the organic substrate and O₂ whilst achieving destabilization of H₂O.

Deprotonation of the substrate is a common step in cofactor-independent oxygenation^{44,41} and our QM and QM/MM analyses reveal that the charge of the substrate determines the geometry of the O₂-substrate complex. Cofactorless addition of O₂ to QND requires swapping from the triplet to the singlet state, a “spin-forbidden” process. The degree of spin-forbiddenness is determined by the magnitude of the SOC term which correlates with the energy difference between the two π -orbitals of O₂. When QND is deprotonated, the geometry of the O₂-QND complex is that which maximizes electron density transfer from QND (π) to O₂ (π^*) orbitals, thus perturbing the π -symmetry of O₂ and leading to values of SOC large enough to make the reaction feasible,⁴² even in the absence of any metal cofactor. For spin-forbidden reactions, the MECF acts as the main dynamical bottleneck for the reaction,³⁴ and we calculated that the additional burden caused by the spin-forbidden character of the reaction is 3.1 kcal mol⁻¹. Similarly to what was obtained for DpgC,⁴³ vitamin K-dependent glutamate carboxylase,⁴⁴ or nogalamycin monooxygenase,⁴⁵ O₂ is not protonated at the MECF, in contrast to what was obtained for *p*-hydroxyphenylacetate hydroxylase, a flavin-dependent monooxygenase.⁴⁶

To summarize, evolution has repurposed the ABH-fold architecture and its simple catalytic machinery to accomplish metal-independent oxygenation. This is achieved by dioxygen entrapment at the oxyanion hole in proximity to the organic ligand with its stabilization and concomitant H₂O destabilization modulated by the nucleophile/hydrophobic elbow residue (Ser/Ala). Substrate deprotonation mediated by the His-Asp subset of the catalytic triad elicits a ligand-O₂ geometry that maximizes electron transfer such that the spin-forbiddenness of the direct reaction is relaxed.

Data availability

Crystallographic structures have been deposited with the Protein Data Bank with accession codes 7OJM, 7OKZ, 8A97, 8ORO, 8OXN, 8OXT.

Author contributions

RAS conceived the project. SB performed the experimental work. SGG, MC, and PGJ carried out the computational work. PvdL and PC contributed to the crystal pressurization experiments. SB, SGG, MC, PGJ, and RAS analyzed the data. The manuscript was written by RAS with contributions from MC and PGJ and commented on by all authors.

Conflicts of interest

The authors declare no competing interests.

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