Interaction of methanol with the oxygen-evolving complex: atomistic models, channel identification, species dependence, and mechanistic implications†

Marius Retegan‡ and Dimitrios A. Pantazis*

Methanol has long being used as a substrate analogue to probe access pathways and investigate water delivery at the oxygen-evolving complex (OEC) of photosystem-II. In this contribution we study the interaction of methanol with the OEC by assembling available spectroscopic data into a quantum mechanical treatment that takes into account the local channel architecture of the active site. The effect on the magnetic energy levels of the Mn$_4$Ca cluster in the S$_2$ state of the catalytic cycle can be explained equally well by two models that involve either methanol binding to the calcium ion of the cluster, or a second-sphere interaction in the vicinity of the “danger” Mn$_4$ ion. However, consideration of the latest $^{13}$C hyperfine interaction data shows that only one model is fully consistent with experiment. In contrast to previous hypotheses, methanol is not a direct ligand to the OEC, but is situated at the end-point of a water channel associated with the O4 bridge. Its effect on magnetic properties of plant PS-II results from disruption of hydrogen bonding between O4 and proximal channel water molecules, thus enhancing superexchange (antiferromagnetic coupling) between the Mn3 and Mn4 ions. The same interaction mode applies to the dark-stable S$_1$ state and possibly to all other states of the complex. Comparison of protein sequences from cyanobacteria and plants reveals a channel-altering substitution (D1-Asn87 versus D1-Ala87) in the proximity of the methanol binding pocket, explaining the species-dependence of the methanol effect. The water channel established as the methanol access pathway is the same that delivers ammonia to the Mn$_4$ ion, supporting the notion that this is the only directly solvent-accessible manganese site of the OEC. The results support the pivot mechanism for water binding at a component of the S$_3$ state and would be consistent with partial inhibition of water delivery by methanol. Mechanistic implications for enzymatic regulation and catalytic progression are discussed.

The water-oxidizing tetramanganese-calcium cluster of photosystem-II (PS-II) that forms the inorganic core of the oxygen-evolving complex (OEC, Fig. 1)\textsuperscript{11-14} is a prime example of a metallocofactor embedded in a membrane enzyme of such complexity\textsuperscript{15-18} that many structural aspects of substrate delivery, product evolution, and proton removal remain insufficiently defined. Driven by photo-induced charge separation at the chlorophyll complex P680\textsuperscript{+}, the OEC cycles through five redox states denoted S$_i$, where $i$ indicates the number of accumulated oxidizing equivalents ($i = 0$–4).\textsuperscript{11,12} Stored in the dark the catalytic centers adopt predominantly the S$_1$ state, whereas the S$_0$, S$_2$ and S$_1$ states are metastable. In these intermediates the Mn oxidation states evolve from Mn$^{[iii]}$,Mn$^{[ii]}$ in S$_0$ to Mn$^{[iv]}$, in S$_1$ (for a recent review on oxidation state assignments, see Krewald et al.\textsuperscript{13}). O–O bond formation\textsuperscript{14,15} occurs upon advancement to the reactive and still unobserved S$_3$ state; following O$_2$ release the cluster is reset to the most reduced S$_0$ state. Strict control of accessibility and ordered substrate binding at the active site, buried into the membrane-embedded part of PS-II, have long been recognized to be functionally critical in regulating the complex chemistry of water oxidation and avoiding side effects. A fundamental aspect of mechanistic regulation in metalloenzymes is how the active site, where catalytic transformation of substrates occurs, is connected to the protein environment. Connections may include solvent-permeable channels for the transport of substrates and products, pathways for the uptake or release of protons, and—in the case of redox transformations—electron transfer pathways to other sites of the enzyme or to peripheral cofactors. Such channels and pathways are not always easily identifiable from crystallographic models, and when they are (for example, in the form of ordered water chains) their function with respect to catalytic activity is not necessarily obvious.

\textsuperscript{†} Electronic supplementary information (ESI) available: Additional geometric and energetic data, sequence alignments, and Cartesian coordinates of all models. See DOI: 10.1039/c6sc02340a

\textsuperscript{‡} Present address: European Synchrotron Radiation Facility, 71 avenue des Martyrs, 38000 Grenoble, France.
Methanol is a key molecule in this respect, because it affects the electronic structure and modifies the EPR signatures\(^{27}\) of all states.\(^{47-64}\) A major effect is that it increases the energy separation between the lowest magnetic levels of the OEC.\(^{48,51,60,62}\) stabilizing the \(S = 1/2\) ground spin states of the \(S_0\) and \(S_2\) states and the diamagnetic ground state of \(S_0\). Other effects relate to the amplitude enhancement of specific spectral forms, for example of the \(g \approx 2\) multilinear signal of the \(S_0\) state in plants over the \(g \approx 4\) component. It is not clear whether a common interaction mode is valid for all catalytic states. Interestingly, higher plants and cyanobacteria behave differently to addition of methanol.\(^{62-76,77}\) the latter appearing less sensitive in its spectroscopic response than the former, but the origin of this species dependence is not understood. Efforts to identify possible methanol binding modes employed deuterated methanol (CD\(_3\)OH) in electron spin-echo envelope modulation (ESEEM) experiments for the \(S_2\) and \(S_0\) states.\(^{49,59,78}\) For the \(S_2\)-state studies\(^{49,59}\), the inferred Mn–H distances were interpreted as consistent with direct ligation of methanol to a Mn center, but the two ESEEM data sets and their interpretation in the context of a structural model of the OEC differ substantially, being consistent either with methanol displacing a water-derived ligand bound to Mn4, or with methanol displacing Glu189, an amino acid residue directly coordinated to Mn1.\(^{64}\) The concept of direct ligation to the Mn1 and Mn4 sites with different affinities was recently used also in the interpretation of D1-Tyr161 radical “split signals” that can be generated from the \(S_0\) state under different illumination conditions.\(^{65}\) On the other hand, the ESEEM study of the \(S_0\) state did not result in observation of specific interactions, disfavoring direct methanol binding to the OEC.\(^{78}\) It is clear that neither the data can be reconciled, nor any of the interpretations supported with confidence. Beyond the intrinsic limitations in the treatment of deuteron dipolar couplings, the involvement of Mn1 as a potential binding site\(^{49,59,63}\) conflicts with channel accessibility studies,\(^{20,25}\) while explanations that invoke drastic reorganizations of first-sphere ligands seem incompatible with the small observed spectroscopic perturbations and the retention of catalytic activity. Similar ambiguities pertain to structural interpretations of methanol effects on the miss parameter for S-state advancement.\(^{64}\)

A significant advance in this line of study was achieved recently by Opala et al.,\(^{65}\) who used \(^{13}\)C-labeled methanol in pulse EPR studies of spinach PS-II poised in the \(S_0\) state. This is the most direct study of methanol interaction with the OEC to date. The measured \(^{13}\)C hyperfine interactions, compared with a reference Mn(m,m,n) complex, argued very strongly against direct binding of methanol to a Mn ion. Maps of the \(^{13}\)C dipolar hyperfine coupleings identified two plausible regions for methanol interaction, either in the cluster of water molecules surrounding Ca\(^{2+}\), possibly by directly replacing one of the calcium-bound waters, or in the vicinity of Mn4 where water molecules can form hydrogen bonds with oxygen bridges, or a combination of the above. Two of the channels shown in Fig. 1 may therefore be implicated in methanol delivery, but no further differentiation can be made and no unique structural models can be proposed. This leaves still open the crucial

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**Fig. 1** The OEC and the proximal branches of associated water channels. Selected residues are shown for orientation. Coordinates are taken from QM/MM models of the \(S_2\) state.\(^{27}\) Inset: schematic diagram and labelling of the inorganic core.
question of substrate delivery and identification; for example, if MeOH indeed replaces the Ca-bound W3 and remains bound in later stages of the catalytic cycle, this would likely exclude the possibility of a calcium-bound water acting as a substrate in a nucleophilic attack scenario for O–O bond formation.\textsuperscript{79–82}

In this study we use available experimental data, including the valuable dataset on \textsuperscript{13}C hyperfine interactions,\textsuperscript{83} to evaluate a series of quantum chemical models that include explicit binding of MeOH at different sites in either protonated or deprotonated form, as well as many second-sphere interaction models. Based on the magnetic and spectroscopic properties of these models, we conclude that the experimental observations have a unique structural interpretation, identifying a second-sphere interaction site for MeOH along a water channel associated with the O4 bridge. The atomistic details of the model provide a novel explanation of how the energy level splittings associated with the O4 bridge. The labeling used for the water molecules is shown in Fig. 2.

The correspondence of these labels with the identification numbers of the most recent crystallographic models with PDB codes 3ARC, 3WU2, and 4UB6 is provided in Table S1 of the ESL.\textsuperscript{†} For each of the twelve positions, after replacing the H\textsubscript{2}O for MeOH several conformations and protonation isomers were generated either by automatically scanning dihedral angles or by manually reorienting specific groups. Of the several tens of conformers thus generated and geometry-optimized, eventually we selected twenty “best-case” models that we consider representative of all conceivable MeOH arrangements. Given that MeOH is only mildly perturbing the spectroscopic signature of the OEC without significantly altering its electronic structure, any models in which the initial positioning of MeOH resulted in extensive structural rearrangements upon optimization such as decoordination of first-sphere residues with concomitant large changes in electronic structure, were discarded.

Geometries were optimized with the ORCA program package,\textsuperscript{84} using the BP86 functional\textsuperscript{85,86} combined with the D3 model for dispersion corrections,\textsuperscript{87} the COSMO model\textsuperscript{88} with a dielectric of 8 to simulate the effect of the protein environment, and the zeroth-order regular approximation (ZORA) Hamiltonian\textsuperscript{89–91} for scalar relativistic effects. ZORA-recontracted\textsuperscript{92} Karlsruhe basis sets of polarized valence triple-\(\zeta\) (Mn, Ca, O, N) and double-\(\zeta\) (C, H) quality were used,\textsuperscript{93,94} with increased integration grids (Grid6 and IntAcc 6.0 in ORCA notation) and tight SCF convergence criteria. Fully decontracted def2-TZVP/J basis sets\textsuperscript{95} were employed in density fitting.

### Methodology

#### Generation of structural models

Our approach for creating atomistic models to describe the interaction of methanol with the OEC is based on systematically substituting first and second sphere water molecules by MeOH and exploring the conformational space for each substitution. The “open cubane” form of the cluster that corresponds to the multiline \(S = 1/2\) signal of the S\textsubscript{2} state\textsuperscript{79} was used as starting point. We employed our previously described QM/MM models of the S\textsubscript{2} state\textsuperscript{97} for preliminary screening and set the size of the final QM-only models to sufficiently include all first-sphere hydrogen bonding interactions. The final models consist of 240–242 atoms. The formally “open” coordination site of the five-coordinate Mn(n) ion Mn1 was also considered as a possible interaction site. However, after multiple attempts we conclude that this site is both inaccessible, in line with previous studies of channel architecture,\textsuperscript{79} and unsuitable for methanol binding as it lies along the Jahn–Teller axis of the Mn(n) ion.

Twelve water molecule positions were considered as candidates for MeOH substitution: four are direct Mn or Ca ligands (W1–W4) and eight are second-sphere crystallographically characterized water molecules (W5–W12) that participate in hydrogen-bonding interactions with first-sphere residues.\textsuperscript{7,8} The W1 and W2 ligands to Mn4 were considered as H\textsubscript{2}O and OH.\textsuperscript{69,72} The labeling used for the water molecules is shown in Fig. 2.

#### Spin states and magnetic properties

For each model a complete set of broken-symmetry (BS) DFT calculations was carried out in terms of single and double spin flips to compute the six pairwise exchange coupling constants \(J_{ij}\) that parameterize the magnetic coupling between the four Mn ions in an Ising-type Hamiltonian. These calculations employed the hybrid meta-GGA functional TPSSH,\textsuperscript{96} which has been...
shown to perform extremely well in the prediction of magnetic properties for high-valent Mn clusters. These BS solutions are not spin eigenfunctions and it is meaningless to attempt any direct comparison with experiment, either with respect to an observed ground spin state or with respect to measured energy differences between magnetic sublevels. It is unfortunate that this crucial point is missed in part of the literature. The exchange coupling constants obtained from the BS solutions serve to form and diagonalize the Heisenberg–Dirac–van Vleck (HDvV) Hamiltonian

$$\hat{H}_{\text{HDvV}} = -2 \sum_{i<j} J_{ij} \hat{S}_i \hat{S}_j$$

thereby obtaining the complete spectrum of spin eigenstates (e.g. 320 magnetic levels for the $S_2$ state including spin multiplets) and their relative energies. These results constitute a valid basis for comparisons with experiment; we use the spin states and energy differences between HDvV states thus identified to evaluate all models.

**Spectroscopic properties**

The hyperfine coupling results from the interaction of the spin of the unpaired electron(s) $S$ and the magnetic nuclear moments $I$:

$$\hat{H} = S \hat{A}^{(A)} \hat{I}^{(A)} = \sum_{k \neq 1,2} \sum_{l \neq 1,2} A_{kl}^{(A)} \hat{S}_k \hat{I}_l^{(A)}$$

The tensor parameterizing the interaction, $A^{(A)}$, consists of three contributions: the Fermi contact, the spin dipolar, and the spin–orbit coupling contribution:

$$A_{kl}^{(A)} = A_{kl}^{(A, FC)} + A_{kl}^{(A, SD)} + A_{kl}^{(A, SO)}$$

In the framework of density functional methods analytical solutions for the three terms have been derived according to the following equations:

$$A_{kl}^{(A, FC)} = \delta_{kl} \frac{8 \pi}{3} \frac{\mu B_{\text{N}}}{2 S} \sum_{\mu \nu} p_{\mu \nu}^{x, \beta} \langle \phi_{\mu} | \delta(r_{A}) | \phi_{\nu} \rangle$$

$$A_{kl}^{(A, SD)} = \frac{\mu B_{\text{N}}}{2 S} \sum_{\mu \nu} p_{\mu \nu}^{x, \beta} \langle \phi_{\mu} | r_{A}^{-5} (r_{A}^{-\delta_{kl}} - 3r_{A} \delta_{kl}) | \phi_{\nu} \rangle$$

$$A_{kl}^{(A, SO)} = -\frac{\mu B_{\text{N}}}{2 S} \sum_{\mu \nu \kappa} \frac{\partial p_{\mu \nu}^{x, \beta}}{\partial \delta_{k}} \langle \phi_{\mu} | z_{\text{SOMF}} | \phi_{\nu} \rangle$$

where $r_i$ is the position vector of electron $i$ and $R_A$ is the position vector of nucleus $A$. The summation runs over the basis functions $\phi$. The first two terms are expectation values over the spin density; the isotropic Fermi contact term arises from the spin density on the nucleus under investigation, while the spin dipolar part is due to the through-space interaction between the electron and nuclear spin magnetic moments. The third term is a second-order spin–orbit contribution, which is important mainly for heavier elements. In the present work, the $^{13}$C hyperfine parameters were computed with the TPSSh functional combined with triple-$\zeta$ ZORA-recontracted basis sets on carbon, locally increased integration grids (SpecialGridIntAcc = 9.0), and inclusion of relativistic picture-change effects. For each model we report the isotropic hyperfine value $(A_{\text{iso}})$, defined as the sum of the Fermi contact term and the isotropic part of the spin–orbit coupling contribution, and the dipolar term $T$ defined as

$$T = \frac{1}{2} A_{kl}^{(A, SD)}.$$

**Sequence alignment**

The sequences of the PS-II core proteins D1 (PsbA), D2 (PsbD), CP43 (PsbC) and CP47 (PsbB) from T. vulcanus, T. elongatus, and S. oleracea were aligned with ClustalW2.†

**Results and discussion**

**Atomistic models for methanol interaction**

Among the many structural models that were optimized and evaluated in terms of their properties, we have selected twenty representative structures (Fig. 3) that fully cover the range of possibilities with respect to methanol interaction with the OEC. The models are labeled according to the water molecule that is substituted by methanol, followed by an additional number that refers to the conformational variant. The models include structures where methanol is deprotonated when it binds as a first-sphere ligand; these are indicated with the letter “d” as a label suffix.

In all cases there is minimal structural perturbation of the inorganic core, even in models where methanol ligates directly to a metal ion of the OEC. The relative energies of these models, computed with three different approaches, are provided in Table S2.† It is stressed however that comparison of these energies is only partially informative for four important reasons. First, a fundamental limitation is that not all models are isomers (e.g. W2-substituted models are unique, and a subset of models contain deprotonated methanol). Second, several models span a narrow energy range; this is particularly true for positions remote from the inorganic cluster that are not associated with large energetic perturbations (e.g. models W5-1, W7-1, and W10-1/2 are isoenergetic even though the access and interaction sites of methanol are fundamentally different). Third, for several structures the truncation of interactions that would propagate beyond the computational model employed in this study implies unknown uncertainties (for example, a high-energy
model might be stabilized by hydrogen bonding or a low-energy model might be destabilized by steric interactions with parts of the protein missing from the QM models). And fourth, relative energies are irrelevant when comparing models that involve different access channels. To appreciate this, let us assume that methanol in reality can only access the calcium site of the cluster; then it is of no relevance that \( W4-1 \) is 3–4 kcal mol\(^{-1} \) higher in energy than \( W7-1 \) (Table S2†), because the latter would be impossible to exist. The only relevant energetic quantities in this case would be the relative permeation barriers through different channels of the enzyme, an aspect that is entirely missing from a cluster QM approach. Therefore, although relative energies are crucial in elucidating and comparing reaction pathways,\(^{14,15,161-167} \) they are of limited utility for the present study and make sense only when comparing closely related subsets, for example pairs of rotamers such as \( W6-1 \) and \( W6-2 \) (see Table S2†). A meaningful evaluation of models in the present case can be based only on comparison of the values of observable properties against experiment, so in the following we focus on the magnetism and spectroscopy of these twenty structures.

**Magnetic properties**

An important effect of methanol on the multiline signal of the \( S_z \) state is the increase of the energy gap \( \Delta E \) between the ground \((S = 1/2)\) and first excited \((S = 3/2)\) states.\(^{62} \) For \( T. \) elongatus it is estimated that the native energy separation of \( 13 \text{ cm}^{-1} \) is increased to \( 22 \text{ cm}^{-1} \) upon addition of methanol, while for spinach the most recent study reported a change from \( 3 \text{ cm}^{-1} \) to \( 25 \text{ cm}^{-1} \).\(^{62} \) Older estimates in the literature place the native \( \Delta E \) of spinach to \( 6 \text{ cm}^{-1} \),\(^{47} \) while for methanol-treated samples previous estimates included 12 cm\(^{-1} \),\(^{50} \) 30 cm\(^{-1} \),\(^{47} \) and 36.5 cm\(^{-1} \).\(^{46} \) Therefore, our principal target here is to identify methanol interaction sites that correlate with an increase of \( \Delta E \).

Table 1 lists all exchange coupling constants for all models of the present study. The spin of the ground and first excited states as well as their energy difference \( \Delta E \), obtained from diagonalization of the HDvV Hamiltonian, are also reported in Table 1 for each model. The results computed for the “native” \( S_z \) state model of the OEC are provided as reference. Before we proceed, it is important to clarify a methodological point: given that the present model of the unperturbed, “native” state of the OEC does not extend too far from the inorganic cluster, it cannot capture differences between species that originate in protein

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Fig. 3 Position of methanol (in blue) and of water molecules around the OEC cluster in selected optimized models. All protein residues are omitted for clarity.
sequence variations at sites that are not included in the model. The structural nature of these remote differences is in any case undefined, as no atomic-resolution structure of higher plant PS-II is still available. In this sense, the unperturbed S2 state model used as the departure point in this study can be viewed in principle as “species-agnostic” and serves as a platform to study relative changes, i.e. the sign and magnitude of \(\Delta E\) expected upon displacement of specific ligated or proximal waters by methanol.

In all cases the nature of the ground spin state \((S = 1/2)\) and the first excited state \((S = 3/2)\) remain the same, regardless of the type of methanol interaction. The exchange coupling constants and the energy splitting between the ground and the first excited states proved in general to be rather insensitive to water displacement by methanol, but there are also a few models where \(\Delta E\) increases or decreases significantly. Focusing on the models that show a large decrease as required for agreement with experiment, we can see that they belong to two groups: the first involves direct binding of methanol to calcium, but only in deprotonated (methoxy) form. These are the models W3-1d and W4-1d, the latter showing (together with W6-1) the largest increase in \(\Delta E\) compared with the native system. The second group includes the two water sites that form part of the O4-related channel and is represented by models W5-1 and W6-1. Both groups correspond to two different channels, the calcium channel and the O4 channel of Fig. 1 respectively. By contrast, there is no water position associated with the third channel (W7, W8) that can reproduce to any extent the phenomenology when occupied by methanol. Direct ligation to Mn4 either has minimal effect on the predicted \(\Delta E\), or it has the opposite effect than what is experimentally observed, when MeOH binds as methoxy in the W1 position (W1-1d).

Therefore, from the perspective of the effect on magnetism, direct ligation to Mn is disfavored and the two models W4-1d and W6-1 stand out from the rest as most consistent with experiment, even though they represent fundamentally different binding modes.

The fact that W4-1d reproduces the large positive change in \(\Delta E\) is somewhat unexpected because there is no direct interaction with a magnetic site. Nevertheless, the values in Table 1 show that coordination of MeO− to Ca2+ does affect the exchange coupling constants, particularly within the cuboidal Mn4CaO4 unit. The major cause for the change in \(\Delta E\) is the diminished value of the ferromagnetic coupling constant \(J_{23}\) compared to the native system. The changes in exchange coupling constants in this model as well as in W3-1d show that such alterations are not necessarily local and cannot always be properly analyzed as such. This serves as a warning that the interpretation of magnetic and spectroscopic responses under the assumption of locality of perturbations may sometimes be oversimplified.

The other model that emerged as a candidate for MeOH interaction is W6-1. Here the major change is observed for \(J_{34}\).

To understand how the presence and orientation of methanol at the W6 site regulates the magnetic coupling between Mn3 and Mn4, it is necessary to examine both W6-1, which shows a shift in \(\Delta E\) that agrees best with experiment, and W6-2, which behaves in exactly the opposite way and shows the poorest agreement among all models in terms of \(\Delta E\). In the native system W6 forms a hydrogen bond with O4. When methanol replaces W6 it can be oriented in a way that a hydrogen bond is also established between its hydroxy group and O4; this case is represented by W6-2. An additional hydrogen bonding interaction is established between W5 and MeOH. However, methanol can also be oriented in a way that abolishes hydrogen...
bonding with O4 and this is what happens in W6-1, where the OH group of methanol acts instead as a hydrogen bond donor to D1-Asp61 and as an acceptor from both W5 and CP43-Arg357 (Fig. 4). These differences have two implications. First, the orientation of methanol that is associated with a higher degree of hydrogen bonding, W6-1, is energetically favorable by ca. 7 kcal mol\(^{-1}\) compared to W6-2, because the positioning of methanol without hydrogen bonding to O4 but with three hydrogen-bonding interactions in total better stabilizes MeOH at the W6 site (Fig. 4, middle). Second, as expected\(^{99,109-111}\) the enhanced superexchange over O4 in the absence of the hydrogen bonding interaction compared to the native system leads to a more negative (antiferromagnetic) value for \(J_{ab}\), stabilization of the low-spin \(S = 1/2\) state of the OEC, and increase in \(\Delta E\). The differences in exchange coupling constants mirror the trend in optimized Mn3–Mn4 distances, which are 2.753 Å for the “native” model, 2.735 Å for W6-1, and 2.777 Å for W6-2.

This atomistic description of the methanol effect on the magnetic structure of the cluster in the case of model W6-1 is almost entirely local compared to W4-1d. As such, it has obvious parallels with our previous qualitative description of the methanol effect on the \(S_2\) state that was cast in terms of an effective interaction \(J_{eff}\) between the terminal Mn4 and the Mn3,Ca subunit of the OEC in a monomer–trimer approximation of the magnetic topology.\(^{62}\) Model W6-1 provides for the first time a structural realization of this mode of action. As an aside, it is worth mentioning that the above analysis can clarify an interesting technical point. Earlier theoretical work employed smaller models that did not include the crystallographic water W6 which hydrogen-bonds to O4.\(^{66}\) It was already noticed that the computed \(\Delta E\) values for these models compared well with experimental values from methanol-treated samples, even though methanol was obviously not included in the models. If we assume that W6-1 is a correct representation of how methanol interacts with the OEC, an explanation for this curious result becomes immediately obvious: a computational model that lacks W6 mimics the effect of methanol on the hydrogen bonding environment of O4.

Evaluation in terms of magnetic interactions allows us to eliminate most models, including all those that involve direct MeOH ligation to Mn. Still, this criterion is not sufficient to definitively distinguish between direct binding of methanol to Ca (model W4-1d) or second-sphere interaction in the vicinity of O4 (model W6-1). In the following section this distinction is achieved conclusively by comparison of the computed \(^{13}\)C hyperfine coupling parameters with the data of Oyala et al.\(^{65}\)

\(^{13}\)C hyperfine interactions

After confirming that our predictions for \(^{13}\)C hyperfine interactions are reliable by comparing with the same reference mixed-valence synthetic complex\(^{111}\) as the one used in the experimental study of Oyala et al.\(^{65}\) (Fig. S1†), we have computed the relevant parameters for all models of Fig. 3 and Table 2 lists the isotropic coupling \(A_{iso}\) and the dipolar coupling \(T\) for the \(^{13}\)C nucleus of methanol in all structures. Even though most models were already disfavored based on the energy level splittings in the previous section, it is important to discuss the \(^{13}\)C hyperfine parameters of the group of models that involve methanol ligation to Mn4, because the two properties lead to identical conclusions and strongly reinforce each other. The \(^{13}\)C \(A_{iso}\) values for MeO\(^-\) binding are two orders of magnitude larger than the experimental values (5.73 MHz and 4.06 MHz for W1-1d and W2-2d), and even the “best-case” scenario represented by model W1-1 is off by an order of magnitude. The deviation of computed dipolar coupling \(T\) values from experiment is equally dramatic, with values ranging from 1.53 MHz (W2-1) up to 2.96 MHz (W1-1d). In view of the above, the conclusion that methanol does not coordinate to Mn4 is definitive.

With respect to the calcium site, models W3-1 and W4-1 agree very well with the \(^{13}\)C hyperfine data, and this observation is in agreement with the conclusions of Oyala et al., who favored calcium as one of the likely methanol interaction sites.\(^{65}\) W3-1, together with W6-1 discussed below, show the best overall agreement with the experimental values. On the other hand, both W3-1 and W4-1 were excluded in the previous section because they cannot explain the increase in the energy gap \(\Delta E\), thus these models cannot satisfy both constraints. The two models that were favored instead were those that involved

![Fig. 4](https://example.com/fig4.png)

**Fig. 4** Comparison of the local hydrogen-bonding environment of O4 in the native system (left) and in the two models W6-1 and W6-2 with methanol occupation of the W6 site.

<table>
<thead>
<tr>
<th>Model</th>
<th>(A_{iso}) (MHz)</th>
<th>(T) (MHz)</th>
<th>Model</th>
<th>(A_{iso}) (MHz)</th>
<th>(T) (MHz)</th>
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<td>0.65</td>
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<td>0.17</td>
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<tr>
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<td>2.54</td>
<td>W8-2</td>
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<td>0.59</td>
</tr>
<tr>
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<td>W12-2</td>
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</tr>
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<td>Exp.</td>
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<td>0.27 ± 0.05</td>
<td>Exp.</td>
<td>0.05 ± 0.02</td>
<td>0.27 ± 0.05</td>
</tr>
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</table>
binding of deprotonated methanol, W3-1d and W4-1d, the latter being one of the two strong candidates. However, the computed values are compelling: this type of methanol binding cannot possibly reproduce the experimental $^{13}$C hyperfine data, because both $A_{iso}$ and $T$ are orders of magnitude too large.

Combined with the results of the previous section this leaves only one candidate, model W6-1. In stark contrast to W4-1d, the agreement with experiment in this case is excellent, with $A_{iso} = 0.02$ versus the experimental value of 0.05 ± 0.02 and $T = 0.24$ versus the experimental value of 0.27 ± 0.05. Therefore, there is only one model, W6-1, requiring methanol delivery through the water channel associated with O4, that simultaneously accounts for the effect on the magnetic couplings of the OEC in the S2 state and reproduces the experimental $^{13}$C hyperfine data.

**Possibility of a secondary binding site**

Although the identification of W6 as the site occupied by methanol is definitive, one may consider whether a second molecule of methanol can be present at a secondary site, potentially depending on concentration. Based on the experimental data and our calculations, binding as a first-sphere ligand to Mn and as a methoxy to calcium are excluded, while occupation of the W6 site is required to explain simultaneously the effect on magnetism and the $^{13}$C hyperfine data, so if there is only one way that methanol interacts with the OEC, this is represented by model W6-1. On the other hand, the presence of a second molecule of methanol cannot be excluded if it occupies a site that satisfies the $^{13}$C hyperfine data while not adversely affecting the energy level splitting. A hypothesis regarding the presence of two methanol binding sites was previously advanced by Sjöholm et al. to account for the different methanol concentration dependence in the induction of “split” EPR signals of the OEC, that is, of states that contain a tyrosyl radical interacting with the manganese cluster.62 The postulated interaction modes (direct binding of methanol to Mn1 and/or Mn4) can no longer be considered viable, so it might be worth revisiting that analysis in light of the present results and of the required occupation of the W6 site.

**S-state invariance of the methanol binding site**

The S1 state of the OEC is diamagnetic ($S = 0$). Parallel-mode EPR however probes the first excited state ($S = 1$)18,114,115 which is very low in energy, less than 2 cm$^{-1}$ higher than the ground state.48 In terms of methanol interaction, it was shown that the parallel-mode S1 state signal at $g = 4.9$ is no longer observable upon addition of methanol,48 suggesting that the triplet state is depopulated, i.e. the energy splitting $\Delta E$ between the ground and first excited states is increased. If we assume that the methanol binding pocket in the S1 state is the same as in S2 and is represented well by the S1 analogue of model W6-1, these observations obtain an obvious structure-based rationalization that mirrors the analysis we presented for the S2 state. The antiferromagnetic coupling between Mn3 and Mn4 in the S1 state24 would be enhanced by the presence of methanol via the same mechanism discussed above, stabilizing the $S = 0$ ground state and resulting in disappearance of the excited-state $S = 1$ signal.

Given that the oxidation states of the manganese cluster cannot influence the channel architecture, methanol should access the OEC through the same channel regardless of the catalytic state of the cluster, so we also expect the same methanol site in the S0 and S3 states. The phenomenology of split EPR signals of both the S1YZ and the S0YZ states has been rationalized in terms of an increase in $\Delta E$ upon addition of methanol,48 so a similar mode of interaction as for the S1 and S2 states (operating on the Mn3-Mn4 antiferromagnetic coupling) would be valid for the S0 state as well. We prefer not to propose specific S0 models at this point because of the more complex EPR phenomenology of this state35,77,116,117 and remaining uncertainties in the protonation states of terminal water-derived ligands and oxo bridges.35,72,118-120 EPR studies of the S3 state reveal a rich and complex phenomenology.25,54,56,71,113 S3 is an integer spin state and interpretation of spectroscopic data is additionally complicated by heterogeneity, related at least in part to structural polymorphism that has only recently begun to be understood.73,134 A more complete understanding of the S3 state is necessary before one can confidently discuss methanol interaction in terms of atomic structure. Nevertheless, the present work constrains the methanol interaction site to W6 and restricts the question only to how methanol might affect the different possible S1 forms, or stabilize one of these forms over the others, for example by inhibiting water delivery.

In conclusion, methanol likely occupies the same site in all states, adopts approximately the same orientation and, at least in the S0-S2 states, affects the electronic structure in the same fashion. Finally, we would like to point out that since direct ligation to a Mn ion is excluded, the apparent S state dependent sensitivity to methanol cannot be attributed to interaction with Mn ions in different oxidation states. An alternative and simpler interpretation of such observations emerges from the present work: if methanol can occupy the same site in all S states and have a similar effect on the electronic structure by modulating the Mn3-O4-Mn4 interaction, the apparent differences in sensitivity between catalytic states arise simply from the different weight of $f_{ij}$ in determining the magnetic properties of the cluster in each S state.

**Effect on the two components of the S2 state**

Two EPR signals are associated with the S2 state of the OEC in plants, a multline signal centered at $g = 2$ that corresponds to an $S = 1/2$ ground state species, and a signal at $g = 4.1$ that arises from an $S = 5/2$ form of the cluster. In 2012 we demonstrated that these two signals correspond to two geometric and valence isomers that are interconvertible and lie very close in energy, within 1 kcal mol$^{-1}$ or less.77 The $S = 1/2$ signal arises from the most stable open-cubane form of the cluster (with respect to the position of the O5 bridge) that has the oxidation state distribution III–IV–IV–IV, i.e. the unique Mn(III) ion is Mn1. The $S = 5/2$ species adopts instead a closed-cubane form with the Mn(III) ion located at the “danger” Mn4 site (Scheme 1).
A crucial experimental observation is that in PS-II from plants, methanol prevents formation of the $g \approx 4.1$ signal, that is, it destabilizes the valence isomer that contains a Mn(II) at the Mn4 site. The reorganization of the hydrogen bonding described above for model W6-1 provides a possible structure-based explanation of the phenomenology: the energetically favorable removal of the H-bonding interaction with O4 when methanol occupies the W6 site results in a more negatively favorable conformation that corresponds to model W6-1. In this sense, W6-1 most probably reflects better how methanol interacts in spinach, something that explains why our QM models for methanol interaction agree better with experimental data in spinach than in cyanobacteria. By contrast, in the D1-Ala87 analogue the larger side chain of asparagine may sterically interact with the W6 pocket, while the formation of transmembrane helices of the CP43/CP47 proteins and the luminal side of CP47 (Fig. S2†). Consistent with the strictly conserved nature of the OEC and the identical chemistry across all oxygenic photosynthetic organisms, no variants exist in the first coordination sphere (associated with the D1 and CP43 proteins), in the environment of the redox active tyrosine YZ (D1-Y161), or in any parts of the protein that either bond covalently or interact via direct or water-mediated hydrogen bonding with the immediate environment of the OEC.

Within a sphere of 10 Å radius around any atom of the Mn$_4$CaO$_5$ cluster, we find two differences, both in the D1 protein: cyanobacterial Ser85 is threonine in spinach PS-II (a conservative substitution that has no relevance to any of the methanol interaction models), and cyanobacterial Asn87 is alanine in spinach (note that T. vulcanus and T. elongatus are identical in this respect). The latter substitution is distinctly non-conservative: it involves side chains of markedly different size and properties. No other such difference between the two organisms could be found along any of the channels in a radius of ca. 20 Å around the OEC for the core proteins. The unique importance of the N87A substitution is that it occurs right at the final turn of the O4-related channel we identified as the delivery channel for methanol.

Focusing on the local structure of this site (Fig. 5), it is apparent that cyanobacterial D1-Asn87 forms a hydrogen bond with the backbone of CP43-Glu354 (the residue that bridges Mn2 and Mn3), whereas in spinach D1-Ala87 cannot form such a bond. The difference in the formation of this hydrogen bond and the difference in the size of the side chains means that the diameter and local architecture of the channel at this point are also different: in the D1-Ala87 analogue methanol can access the interaction site unimpeded and adopt the energetically favorable conformation that corresponds to model W6-1. In this sense, W6-1 should apply equally well to all organisms. It is improbable that fundamentally different access modes exists between species, therefore we consider the following explanation as most likely: the model favored in the present work must be equally valid for both cyanobacteria and higher plants, if methanol can reach this site and adopt this position in all organisms. In the following we identify the structural origin of the species-dependent response and we demonstrate that it strongly supports the proposed delivery channel.

Sequence alignment for the four core proteins of T. vulcanus and S. oleracea (spinach) PS-II reveals in total 41 differences in D1 (Psba), 34 in D2 (Psbd), 73 in CP43 (Psbc), and 114 in CP47 (Psbb). The vast majority are conservative substitutions; non-conservative substitutions are found mostly in the CP proteins (detailed sequence alignments are provided in the ESI†). Overall homology is approximately 88% and 91% for the D1 and D2 chains, while if conservative substitutions are not taken into account, the sequence similarity rises to 96%. Inspection of the sites with respect to the crystallographic model of PS-II from T. vulcanus7,8 shows that most differences occur in the flexible outer loops of the D1/D2 proteins, or in the periphery of Fig. 5 The water channel with the critical N87A substitution, viewed from the OEC. Coordinates for the cyanobacterial model (left) are taken from a QM/MM model7 that was based on the crystallographic model of PS-II from T. vulcanus. The model for spinach (right) was constructed by local side chain replacement to demonstrate the expected structural effect.
proteins, and (b) this channel has the lowest free energy barrier of PS-II at the cavity formed by the extrinsic PsbO and PsbU to Mn₄ in the place of W₁ in the S₂ state of the OEC.

The model for methanol interaction proposed here requires access from the water channel associated with the O₄ bridge of the cluster. In the literature this channel is referred to as the “narrow” channel by Ho and Styring,⁴⁴ “E, F” in the work of Gabdulkhakov et al.,²¹ “channel 2” by Vassiliev et al.,²⁵ and “Path 3” by Ogata et al.²⁶ Consistent with the present findings, Ho and Styring had already suggested by analysis of a low-resolution static crystallographic model of PS-II that this channel would be the most permeable to methanol.²⁶ Molecular dynamics simulations by Vassiliev et al. subsequently demonstrated that: (a) this channel connects the OEC directly with the luminal surface of PS-II at the cavity formed by the extrinsic PsbO and PsbU proteins, and (b) this channel has the lowest free energy barrier for water permeation (peak activation energy ca. 9 kcal mol⁻¹) than at all other channels identified.²⁵ Moreover, recent experimental and theoretical work has demonstrated that NH₄ binds to Mn₄ in the place of W₁ in the S₂ state of the OEC.⁴⁴⁻⁴⁶,⁶₃ Although the interaction modes of NH₄ and MeOH are fundamentally different—the first one acting as a direct ligand to Mn₄, the latter interacting from the second coordination sphere—access of methanol coincides with access of ammonia. This is consistent with competition studies that showed interrelations between the binding of these substrate analogues⁶⁶ and reinforces the assignment of the O₄ channel as the one involved in delivery of substrate analogues, and, by extension, of substrate water.

The O₄ water channel was recently proposed instead to be involved in proton transfer, at least in the S₃–S₁ transition, if the critical assumption is made that O₄ is protonated in the S₀ state.⁷⁵,⁷⁷ Although this idea is not implausible,⁷⁵ it seems to us improbable for three reasons: (a) explicit calculation of protonation patterns by a wide variety of methods⁷²,¹¹⁸,¹¹⁹ suggests that O₄ protonation is unfavorable in S₀, therefore the basis for these computational constructs is questionable; (b) the involvement of this channel in the delivery of methanol (identified here), of ammonia (as a ligand to Mn₄).⁴⁴⁻⁴⁶,⁶₃ as well as of water in the S₃ state would necessitate an unlikely dual role for this channel; and (c) residue D₁-Asn₈₇ would have to be involved in the hypothetical proton transfer, so models that might be valid for T. vulcanus would not be transferable to D₁-Ala₈₇ organisms, which is implausible for a function as critical as proton removal from the water oxidation site. Proton egress instead is most likely to occur via the other channel associated with D₁-Asp₆₁, as previously assigned by Knapp and co-workers based on the monotonic increase of calculated pKₐ values of titratable residues,⁴⁴ and by FTIR and mutation studies.¹³⁶,¹³⁷ The H₂O molecule occupying the W₁ site in the S₃ state has been identified as the Mn-bound ligand that is deprotonated (by D₁-Asp₆₁)¹₀,¹⁴,¹₃₈–¹₄₁ in order to enable oxidation of Mn₄ and allow transition to the S₁ state.¹₃₂,¹₃₄,¹₄₂–¹₄₄ In this case, the influence of MeOH on proton release from W₁ would be a second-order effect, acting through perturbation of the proton acceptor. This may result in modulation of the relative stabilities of intermediates such as tyrosyl radical states⁵₃,⁵₇,⁶₃,¹₂₂,¹₄₅,¹₄₆ in a species-dependent manner and is a subject that deserves further study.

The lack of experimental or theoretical support for either NH₃ or MeOH interacting with Ca²⁺ suggests that the calcium channel system may not be involved in small molecule access to the OEC. It has been proposed instead that the channels at this side of the cluster are more hydrophobic and hence better optimized for removal of dioxygen,²⁶,²⁷ and that calcium itself may play a role in facilitating O₂ release from the cluster.¹⁴⁰ In this case, Ca²⁺ is not a likely site of substrate binding and by extension the water ligands of Ca²⁺ (W₃, W₄) are not likely to be substrates in O-O bond formation. Substrate analogues are instead found either on Mn₄ or at the endpoint of the O₄ channel, implying that this channel may be the only one involved in substrate delivery and that Mn₄ may be the only site involved in substrate binding. This is consistent with recent studies of the S₃–S₁ progression,⁷₃,¹₃₄,¹₄₂,¹₄₈ which showed that the first component of the S₁ state is a “closed-cubane” structure with a five-coordinate Mn₄(v) center.⁷₃ The approximately trigonal bipyramidal geometry of the Mn₄(v) site of this species results in properties such as absorption in the near-IR and high zero-field splitting,⁷₃ similarly to analogous synthetic Mn₄(v) complexes.¹⁴⁹,¹⁵₀ These studies established that water binding can only occur at this S₃-state species: water is delivered via the O₄ channel to the Mn₄ center and binds via a low-barrier transition state,⁷₃,¹₄₈ allowing access to the other components of the S₁ state.⁷₃ Based on the above, we propose that occupation of the W₆ site by methanol may enhance the S₃ population with the five-coordinate Mn₄(v) ion by inhibiting water delivery to the Mn₄ site.⁷₃,¹₄₈

Water binding to the Mn₄ ion⁷₃,¹₄₈ and internal structural rearrangement⁷₄,¹₃₂ would be necessary to reach the structural form of the S₃ state attributed to an all-octahedral Mn₄(v) species⁷₃ that has been proposed by Siegbahn to be the active form of the OEC in O-O bond formation,⁷₅,¹₄₁,¹₅₁ in an o xo–oxyl coupling
scenario.\textsuperscript{14,104,114,115} Given that neither methanol nor ammonia are inhibitors and that the kinetics of substrate exchange show both substrates to be already bound in the \( S_2 \) state,\textsuperscript{136–138} we believe that further evidence is required to conclusively establish whether water binding in the \( S_3 \) state\textsuperscript{73,106,161} is catalytically required for progression to the \( S_2Y' \) state and its subsequent deprotonation and transition to \( S_4 \). A better characterization of the \( S_1 \) and \( S_2Y' \) states is needed to clarify this point.\textsuperscript{131,145,146,162–167}

Conclusions

Precise structural characterization of how substrate analogues interact with the oxygen-evolving complex of photosystem-II is important for understanding regulatory and mechanistic aspects of water oxidation. We studied the interaction of methanol with the \( S_2 \) state of the OEC using extensive quantum mechanical models evaluated against experimental data. The two main criteria used were: (a) the change in the energy splitting between the \( S = 1/2 \) ground-state and the \( S = 3/2 \) first excited state of the complex,\textsuperscript{69} and (b) the recently reported \( ^{13}C \) hyperfine coupling parameters of isotopically labeled methanol.\textsuperscript{69} Our results definitively exclude direct binding of methanol to manganese and strongly disfavor direct interaction with calcium. The only interaction mode that is consistent with observations involves displacement of W6, the final water molecule of the water channel associated with the O4 bridge of the cluster, proximal to the “danger” Mn4 ion. This site reproduces the isotropic and dipolar \( ^{13}C \) hyperfine couplings for methanol and provides a physically transparent explanation for the observed ground-state stabilization: the preferred methanol orientation removes the hydrogen bond between O4 and W6, amplifying the antiferromagnetic coupling between Mn3 and Mn4. The same interaction mode can be expected in other states. The effect of removing the hydrogen bond to O4 can also explain the stabilization of the \( S = 1/2 \) form of the \( S_2 \) state in plant PS-II. Sequence alignments of PS-II core proteins substrate analogues to the OEC, with Mn4 being the only cation of the delivery site, the identification of the delivery channel and the correlation with other experimental and computational results are in line with the pivot mechanism for delivery and binding of water to the five-coordinate Mn4(\textit{iv}) ion of an \( S_2 \)-state component,\textsuperscript{73} and support the idea that the O4 channel is uniquely responsible for delivery of water and substrate analogues to the OEC, with Mn4 being the only directly solvent-accessible site of the cluster.

Notes and references


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