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**PDB References:** MauG–TTQ<sub>OX</sub> MADH, 3sww;  
 MauG–TTQ<sub>OQ</sub> MADH, space group *P*<sub>2</sub><sub>1</sub>, 3sxt;  
 MauG–TTQ<sub>OQ</sub> MADH, space group *C*<sub>2</sub>, 4k3i

## Structures of MauG in complex with quinol and quinone MADH

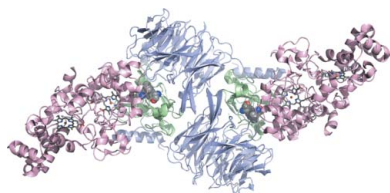
MauG has been cocrystallized with methylamine dehydrogenase (MADH) with its TTQ cofactor in the *o*-quinol (TTQ<sub>OQ</sub>) and quinone (TTQ<sub>OX</sub>) forms and the structures of the resulting complexes have been solved. The TTQ<sub>OQ</sub> structure crystallized in either space group *P*<sub>2</sub><sub>1</sub> or *C*<sub>2</sub>, while the TTQ<sub>OX</sub> structure crystallized in space group *P*<sub>1</sub>. The previously solved structure of MauG in complex with MADH bearing an incompletely formed TTQ cofactor (preMADH) also crystallized in space group *P*<sub>1</sub>, although with different unit-cell parameters. Despite the changes in crystal form, the structures are virtually identical, with only very minor changes at the protein–protein interface. The relevance of these structures with respect to the measured changes in affinity between MauG and various forms of MADH is discussed.

### 1. Introduction

Methylamine dehydrogenase (MADH) utilizes a catalytic tryptophan tryptophylquinone (TTQ) cofactor to catalyze the oxidative deamination of methylamine to ammonia and formaldehyde (McIntire *et al.*, 1991). The biosynthesis of TTQ requires post-translational modification of Trp57 ( $\beta$ Trp57) and Trp108 ( $\beta$ Trp108) of the small  $\beta$ -subunit of the  $\alpha_2\beta_2$  MADH heterotetramer (Davidson, 2011). MauG is a *c*-type di-heme enzyme that is required to complete this post-translational modification. Expression of MADH in a background bearing a deleterious mutation of MauG results in an inactive precursor (preMADH) bearing an incompletely formed cofactor with only a single hydroxyl group inserted into  $\beta$ Trp57 (preTTQ; Pearson *et al.*, 2004). MauG catalyzes the final six-electron oxidation of preTTQ *in vitro* using either H<sub>2</sub>O<sub>2</sub> or reducing equivalents and O<sub>2</sub> to yield the mature TTQ cofactor (Wang *et al.*, 2005).

The MauG–preMADH complex has been crystallized in a triclinic unit cell and its structure has been solved (PDB entry 3l4m; Jensen *et al.*, 2010). Two MauG molecules are found associated with the  $\alpha_2\beta_2$  preMADH tetramer in the asymmetric unit. Despite the fact that the preTTQ site is over 40 Å away from the heme that binds H<sub>2</sub>O<sub>2</sub>, these crystals are catalytically active upon addition of H<sub>2</sub>O<sub>2</sub>. The reaction proceeds by long-range electron transfer from the preTTQ site to the hemes of MauG *via* an interface tryptophan residue, MauG Trp199 (Tarboush *et al.*, 2011). The active form of MauG is an unusual bis-iron(IV) heme species which extracts two electrons from its preMADH substrate to reform the resting di-iron(III) state (Li *et al.*, 2008). However, since MauG-dependent TTQ biosynthesis is a six-electron oxidation, biosynthetic intermediates are clearly formed (Fig. 1). The first two-electron oxidation is the formation of the C–C bond between the tryptophan residues (Yukl *et al.*, 2013) followed by the second hydroxylation of  $\beta$ Trp57 to yield an *o*-quinol species (TTQ<sub>OQ</sub>). The final two-electron oxidation then yields the quinone form of the cofactor (TTQ<sub>OX</sub>) (Li *et al.*, 2006). The only significant conformational change during the entire process is an approximately 20° rotation of the  $\beta$ Trp57 side chain which accompanies cross-link formation (Yukl *et al.*, 2013).

The final step in TTQ formation is fully reversible, and reduction of TTQ<sub>OX</sub> to TTQ<sub>OQ</sub> can be achieved by the addition of dithionite (Husain *et al.*, 1987). Here, we present structures of both TTQ<sub>OQ</sub> and TTQ<sub>OX</sub> MADH cocrystallized with MauG. Interestingly, although the structures are virtually identical to that of the MauG–preMADH

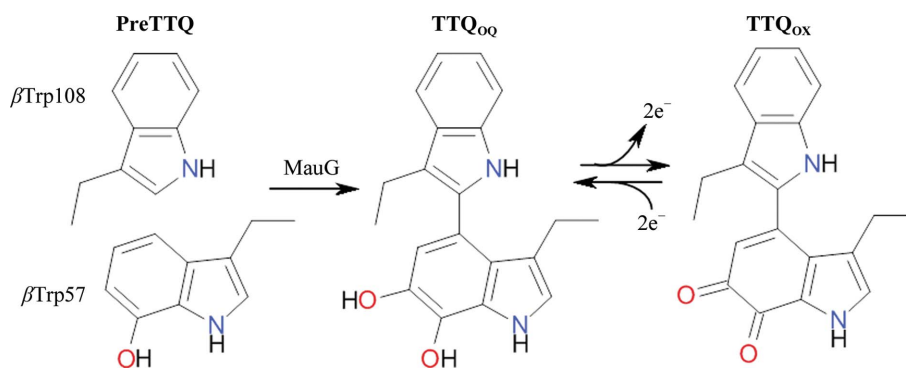


complex, neither is in the same unit cell. Furthermore, the TTQ<sub>OO</sub> MADH complex crystallizes in two different monoclinic space groups as opposed to the triclinic MauG–preMADH structure. The impact of the oxidation state of TTQ on the MauG–MADH interface and crystal packing is discussed.

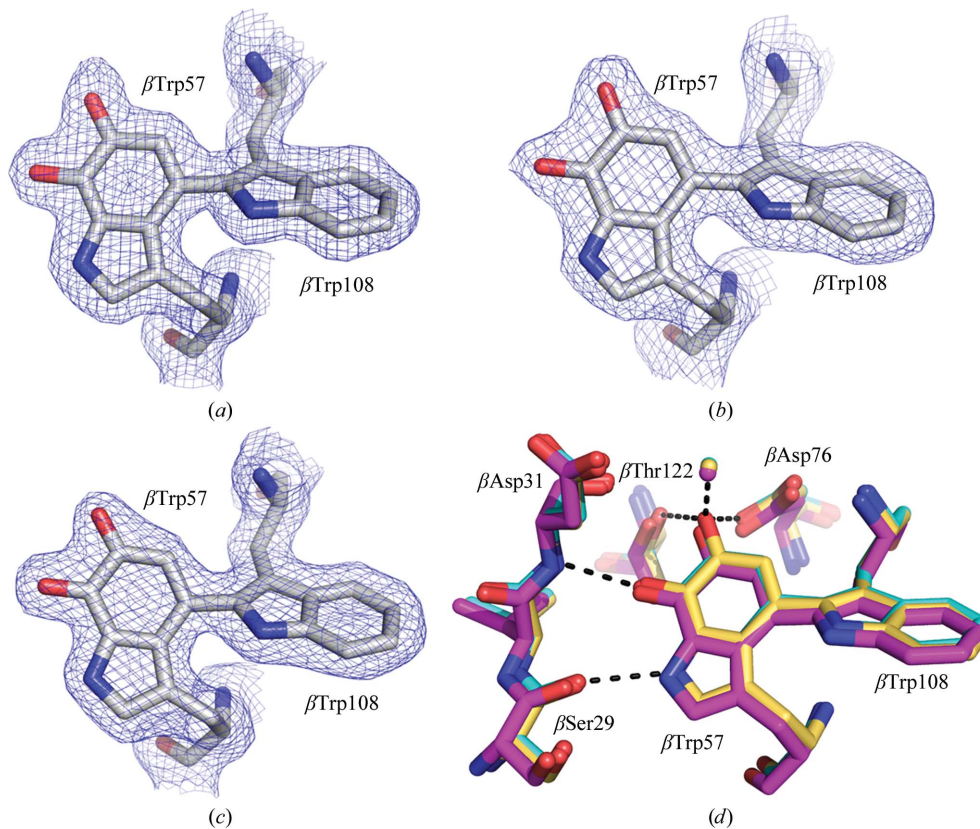
## 2. Methods

Native MADH with the mature TTQ<sub>OX</sub> cofactor was purified from *Paracoccus denitrificans* as described previously (Davidson, 1990). MauG was homologously expressed in *P. denitrificans* and purified by nickel-affinity chromatography as described previously (Wang *et al.*, 2003). The cocrystallization conditions were very similar to those for

the MauG–preMADH complex (Jensen *et al.*, 2010). Briefly, MauG and TTQ<sub>OX</sub> MADH were combined at 100 and 50  $\mu$ M (in terms of MADH tetramer), respectively, in 10 mM potassium phosphate pH 7.5. For the TTQ<sub>OO</sub> MADH complex, protein and reservoir solutions were made anaerobic by purging with argon gas and brought into an anaerobic glovebox (Belle Technologies, UK) maintained at ambient temperature and  $\leq 1.0$  p.p.m. O<sub>2</sub>. Sodium dithionite was added to both protein and reservoir solutions at a final concentration of 2 mM. MauG and MADH proteins were also purged and reduced separately in some experiments prior to crystallization. Upon the addition of dithionite to TTQ<sub>OX</sub> MADH alone, the solution went from green to colorless, consistent with reduction to the TTQ<sub>OO</sub> state. 1  $\mu$ l protein solution was combined with 3  $\mu$ l reservoir solution consisting of 0.1 M sodium acetate, 0.1 M MES pH 6.4, 22–26% PEG 8000. Plate-like



**Figure 1**  
TTQ cofactor maturation by MauG and reversible oxidation and reduction between the TTQ<sub>OX</sub> and TTQ<sub>OO</sub> forms.



**Figure 2**  
Electron density for the TTQ site of MauG–TTQ<sub>OO</sub> MADH in space groups  $P2_1$  (a) and  $C2$  (b) and MauG–TTQ<sub>OX</sub> MADH (c). Atoms are shown in stick form colored according to element and  $2F_o - F_c$  density contoured at  $1.0\sigma$  is shown as a blue mesh. (d) Comparison of hydrogen-bonding interactions to TTQ in MauG–TTQ<sub>OO</sub> MADH in space groups  $P2_1$  (yellow) and  $C2$  (cyan) and MauG–TTQ<sub>OX</sub> MADH (magenta). Water molecules are shown as spheres coloured as for the C atoms. This figure was produced using PyMOL (<http://www.pymol.org/>).

**Table 1**

X-ray crystallographic data-collection and refinement statistics.

Values in parentheses are for the highest resolution shell.

	MauG–TTQ <sub>OX</sub> MADH	MauG–TTQ <sub>OO</sub> MADH	MauG–TTQ <sub>OO</sub> MADH
PDB code	3sws	3sxt	4k3i
<b>Data collection</b>			
Space group	<i>P</i> 1	<i>P</i> 2 <sub>1</sub>	<i>C</i> 2
Unit-cell parameters			
<i>a</i> (Å)	55.22	62.64	346.36
<i>b</i> (Å)	99.42	135.66	55.56
<i>c</i> (Å)	102.83	111.56	112.55
$\alpha$ (°)	64.77	90	90
$\beta$ (°)	74.77	97.37	112.55
$\gamma$ (°)	75.14	90	90
Diffraction source			
Wavelength (Å)	APS 23-ID-D 1.03322	APS 23-ID-B 1.03320	APS 23-ID-B 1.03320
Detector	MAR Mosaic 300 mm CCD	MAR Mosaic 300 mm CCD	MAR Mosaic 300 mm CCD
Resolution range (Å)	50.00–1.86 (1.89–1.86)	50.00–1.81 (1.84–1.81)	50.00–2.00 (2.03–2.00)
Completeness (%)	97.5 (95.8)	98.3 (80.9)	99.8 (99.5)
Multiplicity	3.8 (3.7)	6.5 (2.9)	4.8 (4.1)
$\langle I/\sigma(I) \rangle$	13.9 (3.0)	16.7 (2.3)	13.6 (2.7)
$R_{\text{merge}}^{\dagger}$	0.097 (0.472)	0.093 (0.410)	0.104 (0.551)
<b>Refinement</b>			
Resolution range (Å)	49.12–1.86 (1.91–1.86)	35.01–1.81 (1.86–1.81)	43.06–2.00 (2.05–2.00)
No. of reflections (working/test set)	146343/7734	155734/8247	131712/6968
Final $R_{\text{cryst}}/R_{\text{free}}^{\ddagger}$	0.136/0.178	0.145/0.187	0.146/0.190
No. of non-H atoms			
Protein	13234	13241	13249
Ions	10	6	6
Other	225	192	192
Waters	2009	1824	1707
Total	15478	15263	15154
R.m.s. deviations			
Bonds (Å)	0.026	0.026	0.020
Angles (°)	2.142	2.092	2.065
Overall average <i>B</i> factor (Å <sup>2</sup> )	21.5	29.8	29.8
Ramachandran plot analysis $\S$			
Most favored regions (%)	95.98	96.45	95.99
Additionally allowed regions (%)	3.06	2.90	3.04
Disallowed regions (%)	0.96	0.65	0.97

$\dagger R_{\text{merge}} = \sum_{hkl} \sum_i |I_i(hkl) - \langle I(hkl) \rangle| / \sum_{hkl} \sum_i I_i(hkl)$ , where  $I_i(hkl)$  is the observed intensity and  $\langle I(hkl) \rangle$  is the average intensity of multiple measurements.  $\ddagger R_{\text{cryst}} = \sum_{hkl} ||F_{\text{obs}}| - |F_{\text{calc}}|| / \sum_{hkl} |F_{\text{obs}}|$ , where  $|F_{\text{obs}}|$  is the *i*th observed structure-factor amplitude and  $|F_{\text{calc}}|$  is the calculated structure-factor amplitude.  $R_{\text{free}}$  is the *R* factor based on 5% of the data excluded from refinement.  $\S$  Based on values obtained from the refinement validation options in *Coot*.

crystals were grown by hanging-drop vapor diffusion, appearing within several days and reaching full size within 3–4 weeks. The crystals were cryoprotected by soaking them in reservoir solution containing 10% PEG 400 and were cryocooled in liquid nitrogen. For the TTQ<sub>OO</sub> MADH crystals, cryoprotection and cryocooling were performed in the glovebox using a liquid-nitrogen port to prevent reaction with oxygen.

X-ray diffraction data were collected on GM/CA-CAT beamlines 23-ID-D and 23-ID-B of the Advanced Photon Source (APS), Argonne National Laboratory, Argonne, Illinois, USA. Data were collected at 100 K using a beam size matching the dimensions of the largest crystal face. The data were processed with *HKL-2000* (Otwinowski & Minor, 1997). All structures were solved by molecular replacement using *Phaser* (McCoy *et al.*, 2007) from the *CCP4* program suite (Winn *et al.*, 2011) with the entire MauG–preMADH complex (PDB entry 3l4m; Jensen *et al.*, 2010) as the search model. Restrained refinement with TLS was carried out using *REFMAC* (Murshudov *et al.*, 2011) in the *CCP4* program suite (Winn *et al.*, 2011) and model building was carried out in *Coot* (Emsley & Cowtan, 2004; Emsley *et al.*, 2010).

### 3. Results and discussion

Cocrystallization of TTQ<sub>OX</sub> MADH with MauG could be performed under ambient conditions, whereas the MauG–TTQ<sub>OO</sub> MADH

crystals had to be grown in an anaerobic glovebox in the presence of 2 mM dithionite to generate and preserve the reduced state of both the MauG hemes and the TTQ<sub>OO</sub> cofactor. Both treatments yielded diffraction-quality crystals under nearly identical conditions to those used for the crystallization of MauG–preMADH. Like MauG–preMADH, the MauG–TTQ<sub>OX</sub> MADH complex crystallized in space group *P*1 with two MauG monomers bound to the MADH tetramer in the asymmetric unit. However, the unit-cell parameters for these crystals were significantly different (Table 1). For MauG–TTQ<sub>OO</sub> MADH, monoclinic crystals were grown in either space group *P*2<sub>1</sub> or *C*2. Electron density at the TTQ site for all three crystal forms showed clear density for TTQ<sub>OO</sub> or TTQ<sub>OX</sub>, which are structurally indistinguishable within the resolution of the data (Figs. 2*a–c*). A single water molecule is within hydrogen-bonding distance (2.7–3.1 Å) of the O6 atom of TTQ in all three structures at an angle (91–107°) which is more consistent with the presence of TTQ<sub>OO</sub> than of TTQ<sub>OX</sub>. Furthermore, the arrangement of amino-acid side chains around the TTQ site is essentially identical in all three structures (Fig. 2*d*). It has previously been observed that TTQ<sub>OX</sub> is rapidly reduced to the TTQ<sub>OO</sub> form during X-ray data collection (Pearson *et al.*, 2007), suggesting that we may be observing a majority of TTQ<sub>OO</sub> in all three structures. Interference from the hemes, which are also reduced upon X-ray exposure to the iron(II) state (unpublished data), prevents direct spectroscopic assessment of the TTQ oxidation state in MauG–MADH crystals. Nevertheless, although there are ambiguities in the relative contributions of oxidation states to the

**Table 2**

Direct interactions between protein residues at the MauG–MADH interface.

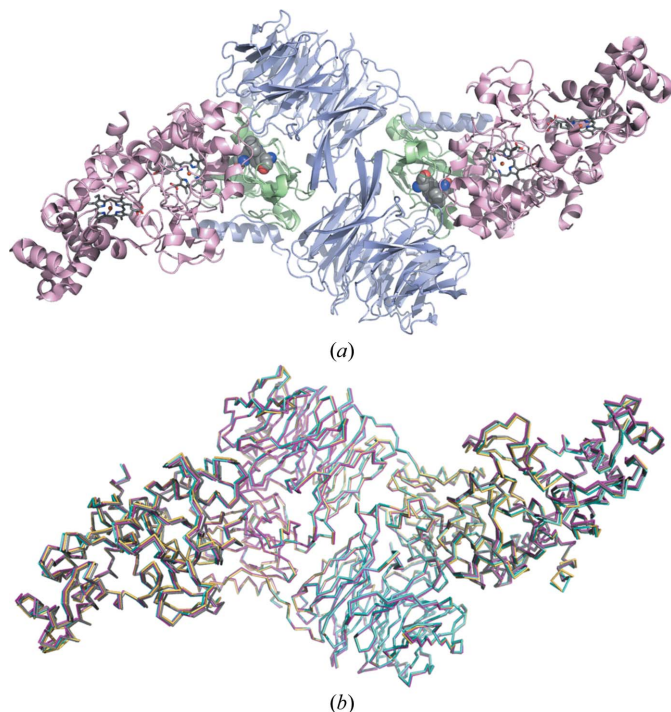
Only those interactions that are conserved between the independent copies of MauG and MADH at distances of less than 4 Å and for which there is strong electron density are listed, except where noted in the text. Specific values are for the *B* chain of MauG. The designation  $\alpha'$  indicates that the residue comes from the other  $\alpha$ -subunit.

Protein–protein interactions			Distance (Å)			
MADH residue	MauG residue	Interaction type	MauG–preMADH	MauG–TTQ <sub>OO</sub> MADH ( <i>C</i> <sub>2</sub> )	MauG–TTQ <sub>OO</sub> MADH ( <i>P</i> <sub>2</sub> <sub>1</sub> )	MauG–TTQ <sub>OX</sub> MADH
$\alpha$ Asp180	Arg338	Salt bridge	3.0	3.0	3.0	3.1
$\alpha$ Pro158	Met333	Hydrogen bond	2.9	2.8	2.9	2.9
$\alpha$ Ser157	Gly331	Hydrogen bond	2.6	2.4	2.5	2.5
$\alpha$ Arg197	Phe191	$\pi$ -Stacking	3.7	3.7	3.7	3.3
$\alpha'$ Gly29	Arg208	Hydrogen bond	3.1	3.0	2.7	2.9
$\alpha'$ Asp31	Lys209	Hydrogen bond	2.9	3.2	3.1	3.1
$\alpha'$ Asp31	Gln210	Hydrogen bond	3.0	3.0	3.0	3.1
$\beta$ Glu101	Arg338	Salt bridge	2.7	2.8	2.8	2.6
$\beta$ Glu101	Trp199	Hydrogen bond	3.4	3.5	3.2	3.1
$\beta$ Ile126	Gly211	Hydrogen bond	2.9	2.8	2.8	2.8
$\beta$ Thr54	Arg202	Hydrogen bond	2.8	4.1	4.8	2.7
$\beta$ Ser56	Thr198	Hydrogen bond	3.0	3.7	3.9	3.3
$\beta$ Thr44	Gln210	Hydrogen bond	3.4	3.2	3.2	3.3

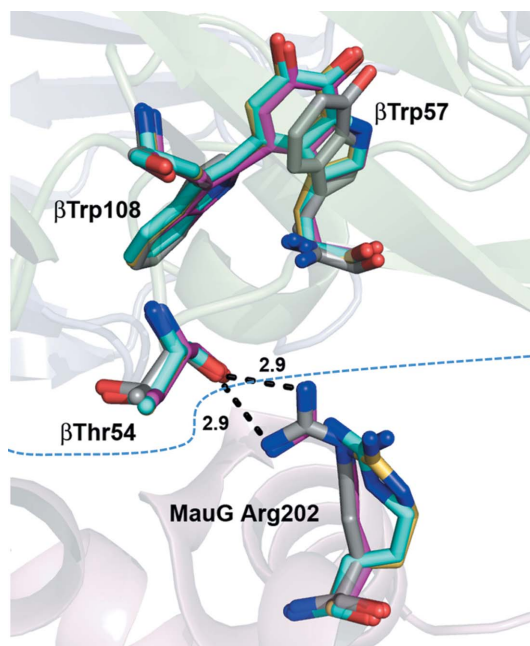
final electron-density maps, it is clear that the initial oxidation state of TTQ has an impact on packing during crystal growth.

Despite the differences in crystal form, all three structures are essentially identical to the MauG–preMADH substrate complex, with  $C^\alpha$  r.m.s.d. of  $\leq 0.52$  Å<sup>2</sup> over the entire complex (Fig. 3). The TTQ cofactor is buried at the interface between MauG and MADH, and so does not directly mediate crystal contacts. However, it could potentially affect the protein interface through changes in dynamics and the relative amounts of conformational states, or by structural changes that are within coordinate error. The observation that the binding affinities of MauG for TTQ<sub>OO</sub> and TTQ<sub>OX</sub> MADH are very similar and are approximately tenfold weaker than for preMADH

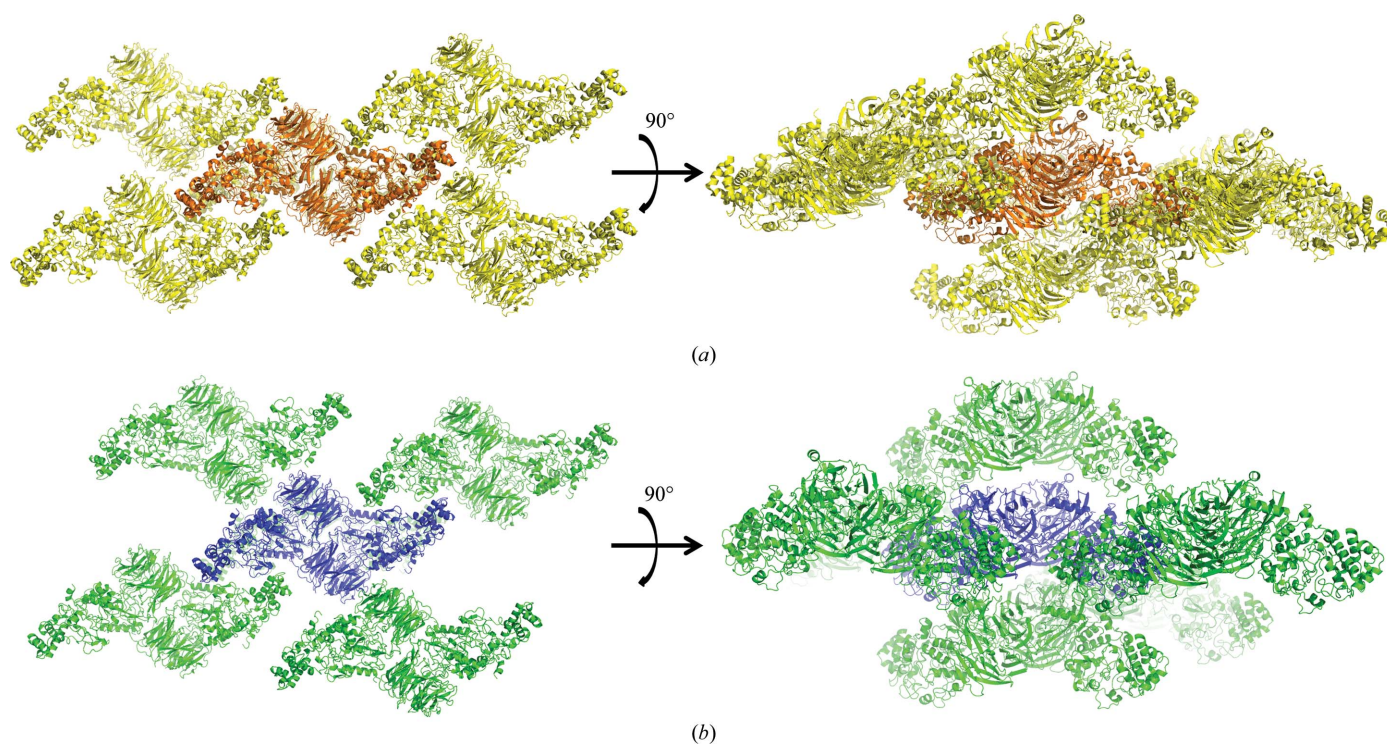
supports the transmission of a physical effect to the (pre)MADH surface (Lee *et al.*, 2009; Shin *et al.*, 2010). PreMADH has not been crystallized alone, and TTQ<sub>OX</sub> MADH only crystallizes alone when it has undergone limited proteolysis, so it is unknown whether there is a significant structural difference between these forms in the absence of a protein binding partner (Chen *et al.*, 1998). Interestingly, the packing differences between the MauG–MADH structures are not a consequence of a change in the relative angle between the two proteins within the complex, suggesting that this is stable (Fig. 3). In fact, the small amount of variability between the structures seems to stem from the MauG portion of the complex rather than the highly stable MADH core, suggesting some propagation of the MADH

**Figure 3**

(*a*) Structure of the MauG–preMADH complex (PDB entry 314m). MauG is shown in pink,  $\alpha$ -MADH in blue and  $\beta$ -MADH in green. The site of TTQ formation is shown as spheres and the MauG hemes are shown as sticks colored by element. (*b*) Backbone traces of MauG–TTQ<sub>OO</sub> MADH in space group *C*<sub>2</sub> (cyan), in space group *P*<sub>2</sub><sub>1</sub> (yellow) and MauG–TTQ<sub>OX</sub> MADH (magenta) overlaid on the MauG–preMADH structure (gray). This figure was produced using *PyMOL* (<http://www.pymol.org/>).

**Figure 4**

Comparison of an interface interaction in MauG–preMADH (gray), MauG–TTQ<sub>OX</sub> MADH (magenta) and MauG–TTQ<sub>OO</sub> MADH in space groups *P*<sub>2</sub><sub>1</sub> (yellow) and *C*<sub>2</sub> (cyan). The protein backbone cartoon is from the MauG–preMADH structure and is colored according to subunit as in Fig. 3(*a*). The MauG–preMADH interface is indicated by the blue dotted line and hydrogen-bond interactions are shown as black dotted lines with distances indicated. This figure was produced using *PyMOL* (<http://www.pymol.org/>).



**Figure 5**  
Crystal packing in MauG-preMADH (a) and MauG-TTQ<sub>OO</sub> MADH in space group C2 (b). This figure was produced using *PyMOL* (<http://www.pymol.org/>).

cofactor state through the interface to MauG which ultimately alters crystal packing.

It has previously been noted that the MauG-preMADH interface is mediated largely by water (Tarboush *et al.*, 2011) and that there are relatively few direct interactions between protein residues (Table 2). This is also true of the TTQ<sub>OO</sub> and TTQ<sub>OX</sub> structures, in which the positions and interactions of the interface waters are largely conserved. Nearly all of the direct interactions between protein residues are also highly conserved, with the exception of a hydrogen bond between MauG Arg202 and the backbone carbonyl of  $\beta$ -MADH Thr54, which is disrupted in both TTQ<sub>OO</sub> structures (Fig. 4) and in one of the monomers of the TTQ<sub>OX</sub> structure. The triclinic packing of MauG-preMADH is already very close to having a crystallographic twofold rotation axis owing to the noncrystallographic twofold symmetry of the complex (Fig. 5). Thus, although the loss of a single interface hydrogen bond is a relatively small change which does not affect the interface as a whole, it may be sufficient to shift the crystal packing sufficiently to allow the transition to a monoclinic space group. The observation of the Arg-Thr hydrogen bond in one subunit of the MauG-TTQ<sub>OX</sub> MADH structure and the fact that Arg202 is not conserved among MauG homologs argue against its relevance in directly modulating the binding affinity between MauG and different forms of MADH in solution, leaving how this is achieved an open and intriguing question. However, the ability to crystallize MauG-MADH in a higher symmetry space group has practical significance for structurally characterizing high-valent MauG heme intermediates before the onset of significant X-ray reduction. Experiments using these crystals for this purpose are currently under way in our laboratories.

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