

Identification of YdhV as first molybdoenzyme binding a bis-Mo-MPT cofactor in Escherichia coli

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ABSTRACT

The oxidoreductase YdhV in *Escherichia coli* has been predicted to belong to the family of molybdenum/tungsten cofactor (Moco/Wco) containing enzymes. In this study, we characterized the YdhV protein in detail, which shares amino acid sequence homologies to a tungsten-containing benzoyl-CoA reductase binding the bis-W-MPT (for metal-binding pterin) cofactor. Our studies showed that YdhV has a preference for molybdenum over tungsten as metal to be inserted into the MPT backbone. The cofactor was identified to be of a bis-Mo-MPT type, which represents a novel form of Moco that has not been found earlier in any molybdoenzyme. In-depth characterization of YdhV by X-ray absorption and EPR spectroscopy revealed that the bis-Mo-MPT cofactor in YdhV is redox active, while a bis-W-MPT cofactor is redox inactive. The bis-Mo-MPT and bis-W-MPT cofactors include metal sites with the metal centers binding the four sulfurs from the two dithiolene groups in addition to a cysteine and likely a sulfido ligand. The unexpected presence of a bis-Mo-MPT cofactor opens an additional route for cofactor biosynthesis in *E. coli* and expands the canon of the structurally highly versatile molybdenum and tungsten cofactors.

INTRODUCTION

Molybdenum and tungsten are important trace elements that are essential in many biological systems.¹ Both transition metals share comparable physicochemical properties that are utilized by numerous enzymes for catalytic reactions involving metal redox changes and group transfer reactions in concert with intramolecular electron and proton transfer.

The prototypical Mo or W cofactors include an unusual tricyclic pyranopterin moiety referred to as metal-binding pterin (MPT).² After metal insertion, the MPT moiety is further modified, resulting in the diverse molybdenum (Moco) or tungsten (Wco) cofactor types. Moco or Wco containing enzymes are divided into three families according to the metal coordination environment in the cofactors as well as distinct active site structures and reaction types (Figure 1), namely the xanthine oxidase (XO), the sulfite oxidase (SO), and the dimethyl sulfoxide (DMSO) reductase families.³ The XO family is characterized by a (MPT)Mo/W(OSO⁻) core in the oxidized state, with one MPT ligand bound to the metal. The sulfido-group (i.e. Mo=S) is cyanide labile.⁴ In enzymes of the XO family in prokaryotes, the cofactor can be modified by attachment of a CMP nucleotide to the phosphate group of Mo-MPT, forming the metal-binding-pterin cytosine dinucleotide (MCD) cofactor (Figure 1).^{5, 6}

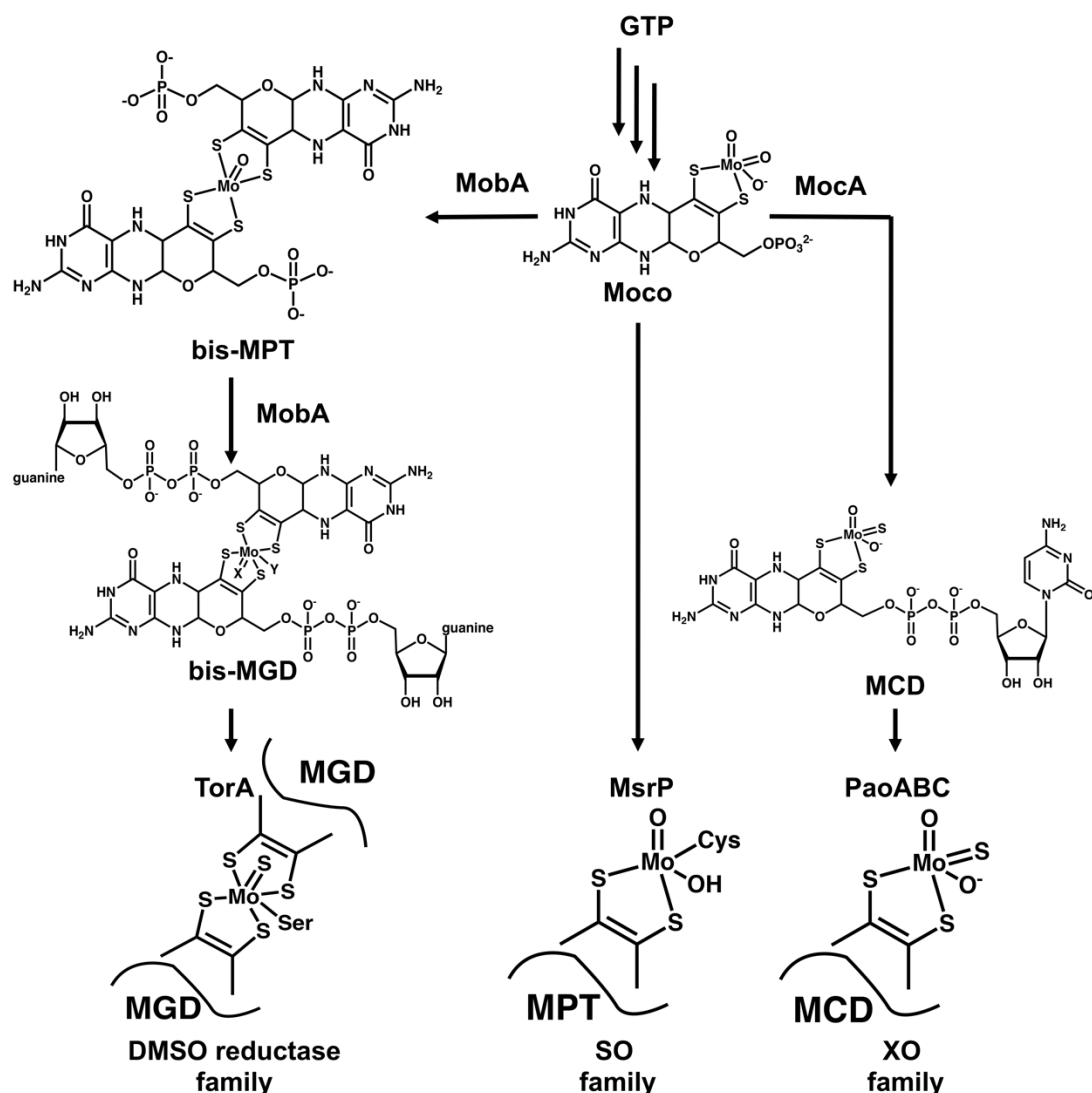


Figure 1: **The three families of molybdoenzymes in *E. coli*.**

The metal-free form of the cofactor is denoted metal-binding pterin (MPT), a 5,6,7,8-tetrahydropyranopterin. MPT is converted to Mo-MPT by insertion of the molybdenum atom. Mo-MPT (shown in the tri-oxo structure⁷) can be further modified and three different molybdenum-containing enzyme families are classified according to their coordination at the molybdenum atom: the XO, SO, and DMSO reductase families. The SO family is characterized by a MPT-Mo^{VI}O₂Cys ligand sphere and this cofactor is present in the *E. coli* MsrP protein. The XO family contains a MPT-Mo^{VI}OS(OH) core. Here, the MPT core can be modified by an additional CMP nucleotide at the phosphate group, forming MCD. The representative protein of this family in *E. coli* is the PaoABC protein. The DMSO reductase family contains a MGD₂-Mo^{VI}XY core with X being either a sulfur or an oxygen ligand and Y either being a hydroxo or amino acid ligand (Ser, Cys, Sec, and Asp ligands have been identified thus far). The well-characterized representative of this family in *E. coli* is the TorA protein, that solely contains bis-MGD with a sulfido ligand and a serine ligand from the protein backbone.

CMP attachment to Mo-MPT is catalyzed by the Moca protein in *Escherichia coli*.⁸ Enzymes of the SO family show a (MPT)Mo(O₂) core in the oxidized state and usually an additional

cysteine ligand at the metal. The DMSO reductase family is exclusively present in bacteria and archaea and all members bind two pterin ligands at the metal in a (MPT)₂Mo/W(XY) core. The ligand X can be a sulfido or oxo group. The Y ligand is typically an amino acid residue such as serine, cysteine, selenocysteine, or aspartate, but can be also a coordinated water species. In DMSO reductase family enzymes, further MPT modification occurs by addition of a GMP mononucleotide to the phosphate group, resulting in two MPT guanine dinucleotide (MGD) moieties at the bis-MGD cofactor.^{1, 9-12} During bis-MGD synthesis, binding of two MPT molecules to one metal ion precedes addition of the GMP nucleotides. The step of bis-MPT formation and GMP addition is catalyzed by the MobA protein. For bis-MPT formation, metal ligation to MPT is obligatory¹³, but no further proteins are involved in the reaction. In the next step, a GMP molecule is added to each MPT bound to MobA and the resulting bis-MGD cofactor is inserted into the target enzymes by bis-MGD binding chaperones.^{7, 14, 15} Tungsten-containing enzymes grouped into the DMSO reductase family, like the aldehyde-ferredoxin oxidoreductase AOR from *Pyrococcus furiosus* or the benzoyl-CoA reductase BamBC from *Geobacter metallireducens*,^{16, 17} were shown to bind a bis-W-MPT cofactor without guanine nucleotides attached to the terminal phosphate group of MPT. The synthesis of this cofactor is unexplored in these organisms.

In *E. coli*, numerous MPT-containing enzymes were identified, which all bind molybdenum at their active site.^{5, 18} A well-characterized typical member of the XO family is the periplasmic aldehyde oxidoreductase PaoABC, binding the sulfido-containing MCD cofactor.^{5, 18, 19} The periplasmic methionine sulfoxide reductase (MsrP) is a typical member of the SO family that binds the Mo-MPT cofactor.^{5, 20} However, the majority of molybdoenzymes in *E. coli* belong to the DMSO reductase family, binding the bis-MGD cofactor.³ Different ligands at the Mo-atom of members of this family were identified, including a recently characterized sulfido-ligand of *E. coli* TMAO reductase that is present at the Mo atom in

addition to a serine ligand.²¹ This sulfido-ligand was first identified to be essential for the activity of formate dehydrogenases and periplasmic nitrate reductases that bind a protein-derived cysteine ligand instead.^{22, 23} It has been suggested that the sulfido-group is sensitive to oxidative treatment, therefore this ligand might be present in more molybdoenzymes than originally expected.²¹

The so far uncharacterized proteins encoded by the *ydhYVWXUT* operon in *E. coli*²⁴ were also grouped in the DMSO reductase family of molybdenum/tungsten enzymes.⁵ The *ydhYVWXUT* operon encoding a predicted oxidoreductase in *E. coli* was first described by Partridge et al. in 2008.²⁴ These studies showed that the *ydhYVWXUT* genes are transcribed from an FNR-dependent promoter under anaerobic conditions. YdhV has been predicted to bind a MPT cofactor based on about 45% amino acid sequence homology to *P. furiosus* AOR and *G. metallireducens* BamBC.^{16, 17, 25} An YdhV mutant was reported to be impaired in anaerobic utilization of certain sulfur sources (e.g. L-cysteine, D-cysteine, L-cysteic acid, hypotaurine, butane sulphonic acid).²⁴ No further characterization of this enzyme was available so far.

In this study, we characterized the purified cofactor-binding protein YdhV from *E. coli* in detail. The cofactor in the native enzyme was identified to be of the bis-Mo-MPT type and also the respective bis-W-MPT form can be assembled in the enzyme. bis-Mo-MPT represents a novel form of Moco that has not been observed in any molybdoenzyme before. The bis-Mo/W-MPT cofactors in YdhV likely contain a sulfido and a cysteine ligand at the metal.

MATERIALS AND METHODS

Bacterial strains, plasmids, media and growth conditions

Escherichia coli BW25113 $\Delta mobA$ and $\Delta mocA$ cells were obtained from the Keio collection.²⁶ The $\Delta mobA/\Delta mocA$ strain was constructed by introducing the *mocA* gene deletion into the $\Delta mobA$ strain using P1 transduction.²⁶ Successful deletion of *mobA* and *mocA* was checked by PCR amplification and enzyme activities of proteins expressed in these strains. For the expression of proteins, cells were generally grown, unless otherwise indicated, at 30 °C under aerobic conditions in LB medium containing 1 mM molybdate, 20 μ M isopropyl β -D-1-thiogalactopyranoside (IPTG) and 150 μ g/mL ampicillin for 24 h.

Expression and purification of recombinant proteins

The gene encoding *ydhV* was amplified from the *E. coli* K12 genome.²⁷ The published gene sequence was used to design primers that permitted cloning into the *SacI* and *HindIII* sites of the expression vector pTrcHis.²⁸ The resulting plasmid, designated pSR153, contains the *ydhV* gene with a His₆-tag fused to the N-terminus of YdhV. For heterologous expression in *E. coli*, pSR153 was transformed into BW25113 cells. Additionally, two primer pairs were designed to clone the *ydhY* gene into the MCS-1 (multiple cloning site) and the *ydhWXUT* genes into MCS-2 of pACYCDuetTM-1 (Novagen), resulting in plasmid pSR242. The expression of *E. coli* TMAO reductase (TorA) from plasmid pJF119EH²⁹, MsrP from plasmid pSR8³⁰ and PaoABC from plasmid pMN11¹⁹ was performed according to the respective published procedures.

The *E. coli* K-12 strain BW25113 was employed for the expression of all proteins described herein, including those containing a deletion of the *mobA*, *mocA* and *moaA* genes. Expression pre-cultures were prepared in LB-medium containing 1 mM Na₂MoO₄, 20 μ M IPTG and 150 μ g/mL ampicillin. Expression cultures were started with 2 mL/L of an overnight culture and

incubated at 30 °C and 130 rpm for 24 h. YdhV additionally was expressed with different $\text{Na}_2\text{WO}_4^{2-}$ (0 - 100 μM) or $\text{Na}_2\text{MoO}_4^{2-}$ concentrations (0 - 100 μM) in presence of either 10 μM $\text{Na}_2\text{MoO}_4^{2-}$ or 10 μM $\text{Na}_2\text{WO}_4^{2-}$, respectively. Anaerobic expression of YdhV was performed similarly to aerobic expression, only in that the cultures incubated statically in closed flasks at 30 °C for 24 h.

Purification was performed either aerobically at 4 °C or anaerobically in a Coy chamber ($\text{O}_2 < 10$ ppm). The cells were harvested by centrifugation and the cell pellets were resuspended in phosphate buffer (50 mM NaH_2PO_4 , 300 mM NaCl, pH 8.0). 20 mL of cell pellet were lysed by sonification in a 50 mL Falcon tube for 6 min in an icebath. The cleared lysate was applied to 0.7 mL nickel-nitrilotriacetate (Ni-NTA) resin per liter of culture. The column was washed with 20 column volumes of phosphate buffer each, first with one containing 10 mM imidazole and then with one containing 20 mM imidazole. Proteins were eluted with phosphate buffer containing 250 mM imidazole. Subsequently buffer exchange was performed by using Sephadex G25 desalting columns with the following protein-specific buffers: MsrP, 50 mM NaH_2PO_4 , 300 mM NaCl, pH 8.0; PaoABC, 50 mM Tris-HCl, pH 7.5; YdhV and TorA, 100 mM Tris-HCl, pH 7.2.

YdhV samples for X-ray absorption spectroscopy were prepared in a similar fashion as described above. Mo-containing YdhV was expressed aerobically, whereas W-containing YdhV was expressed anaerobically. Slight modifications to the expression and purification procedures were applied. Briefly, expression cultures were supplemented with either MoO_4^{2-} or WO_4^{2-} concentrations of 1 mM or 10 μM and 1 mM, respectively. YdhV was purified as described above, however for Br^- containing samples, the 300 mM NaCl present in the imidazole buffers was replaced by 300 mM NaBr, and samples underwent buffer exchange into 100 mM Tris-HBr, pH 7.2. For chloride-free samples, YdhV purified with chloride underwent buffer exchange into 100 mM Tris-acetate, pH 7.2.

Activity assays

The activity of PaoABC was determined spectrophotometrically by monitoring the reduction of 1 mM ferricyanide at 420 nm using a UV-visible spectrophotometer in a 1 mL cuvette as described previously.¹⁹ The substrate vanillin was used in 100 mM Tris pH 6.8 with protein concentration of 5-20 μ M.

The activity of TorA and MsrP using an enzyme concentration of 5-20 μ M was measured in 1 mL cuvettes at 600 nm containing 4 μ L of 100 mM benzyl viologen, 5 μ L of 1.5 M TMAO, and sodium dithionite as described previously.²⁹⁻³¹ The activity of TorA was measured in 100 mM KH_2PO_4 (pH 6.5) and the activity of MsrP was measured in citrate phosphate buffer (pH 5).

To attempt identification of the substrate of YdhV, several substrates under different conditions were tested, i.e. in the context of the broad scope of reactions that so far characterized molybdoenzymes are able to perform. Briefly, the oxidation and reduction of formaldehyde, acetaldehyde, crotonaldehyde, butyraldehyde, hexanal, glyceraldehyde, benzaldehyde, glutaraldehyde and benzoyl-CoA in the presence of benzyl viologen and/or methyl viologen was tested at pH 5.0, 7.2 and 9.0. YdhV electron acceptor/donor activity was assessed using ferricyanide/ferrocyanide, methylene blue and DCPIP. In addition, redox reactions coupled to methyl viologen regeneration, as described above for TorA and MsrP, were adapted to YdhV with the following potential substrates: cystine, cysteine, methionine sulfoxide, sulfate, methionine, sodium dithionite, dithiothreitol (DTT), Na_2S , DMSO, TMAO, xanthine, vanillin, sodium nitrate, sodium chlorate, sodium perchlorate, O_2 , sodium molybdate, FeCl_3 , glucose and sucrose.

We further tried activity assays of YdhV co-expressed with the *ydhYWXUT* operon after aerobic or anaerobic expression. Redox cycling or the anaerobic reconstitution of the Fe-S cluster in YdhV were carried out in addition.

Quantification of metal and Moco saturations

Metal contents of proteins were determined for protein concentrations of about 10-1000 μM . The quantifications of Mo, W, Fe, or P were performed using inductively coupled plasma optical emission spectroscopy (ICP-OES) on a PerkinElmer Life Sciences Optima 2100DV instrument or total reflection X-ray fluorescence (TXRF) analysis on a PicoFox spectrometer (Bruker) as described previously.^{32, 33}

MPT contents were determined fluorometrically after conversion of the molecule to FormA, as described previously.³⁴ While an incubation for 30 min at 95°C in the presence of acidic iodine oxidizes all released Moco derivatives to the fluorescent FormA, an overnight incubation at room temperature results in FormA-NMP derivatives with the nucleotide retained. 0.5-30 μM of proteins were used for the oxidation by acidic I_2/KI . FormA and FormA-GMP or FormA-CMP were separated by a C18 reversed-phase high pressure liquid chromatography column (4.6 x 250 mm Hypersil ODS, 5 μm particle size) according to the method described previously.³⁴ The fluorescence of FormA, FormA-CMP and FormA-GMP was quantified by an Agilent 1260-series detector using excitation at 383 nm and emission detection at 450 nm.

ICP-MS

Aliquots of 100 μL of 16.4 μM YdhV were digested using 1 mL HNO_3 and 900 μL of H_2O (containing 5 $\mu\text{g}/\text{L}$ Rh as internal standard) in a MARS 6 closed vessel microwave system (CEM) at 200 °C for 20 min. The digestion solution was transferred to a sample vial and diluted 1:10 at least. Blank digestions were processed in the same way.

The concentrations of Mo and P were simultaneously quantified by ICP-MS (Agilent 8800 ICP-QQQ) in the MS/MS mode using oxygen as reaction gas. The nebulizer gas flow and parameters of lenses, Q1, collision cell and Q2 were tuned daily for maximum sensitivity

(oxide ratio <1.0% ($^{140}\text{Ce}^{16}\text{O}^+ / ^{140}\text{Ce}^+$), double charged ratio <1.5% ($^{140}\text{Ce}^{++} / ^{140}\text{Ce}^+$), background counts <0.1 cps). For quantification an element mix (100 mg/L, Merck - XVI) was diluted in 5% HNO_3 as an external calibration standard.

X-ray absorption spectroscopy

Protein samples for XAS were prepared by concentrating YdhV (Merck Amicon Ultra-0.5 Centrifugal Filter Units, 10 kDa cutoff) in the absence (YdhV^{ox}) or presence (YdhV^{red}) of a 10-fold excess of sodium dithionite, resulting in YdhV in the as-isolated (oxidized) or reduced states, respectively. Samples were loaded into acrylic-glass sample holders (50 μL), sealed with Kapton tape, and were flash-frozen and stored in liquid nitrogen until use. To follow redox changes of YdhV, the prepared XAS samples were subjected to additional treatments at the beamline. Following XAS (Mo) data collection for YdhV^{red} , the sample was thawed and exposed to air for 10 min reoxidation before freezing again in liquid nitrogen ($\text{YdhV}^{\text{reox}}$). Likewise, following XAS (Mo) data collection for $\text{YdhV}^{\text{reox}}$, the sample was thawed and incubated with a 10-fold excess of sodium dithionite for 10 min reduction before freezing again (YdhV^{red}).

Mo K-edge³⁵ XAS was carried out at the Samba beamline at Soleil synchrotron (Paris, France) using a standard set-up (double-crystal Si[220] monochromator, 36 element energy-resolving Ge detector for fluorescence data collection) as previously described.^{7, 35, 36} Fe K-edge and W L₃-edge XAS was carried out at beamline KMC-3 at BESSY-II (Helmholtz Center Berlin, Germany) using a standard set-up (double-crystal Si[111] monochromator, 13-element energy-resolving Ge detector). Samples were held in a liquid-helium cryostat at 20 K. The monochromator energy axes were calibrated using Mo, Fe, or W metal foils (reference energies: Mo, 20003.9 eV; Fe, 7112.0 eV; W, 10206.8 eV). Up to 10 XAS scans (~30 min duration, one scan per sample spot) were averaged for signal-to-noise ratio improvement.

XANES and EXAFS spectra were derived as described earlier.^{33, 35-37} EXAFS simulations were performed with in-house software³⁷ and phase functions calculated by FEFF9 ($S_0^2 = 1.0$, Mo and W, or 0.85, Fe).³⁸ EXAFS Fourier transforms (FTs) were calculated for k -ranges of 1.6-14.0 \AA^{-1} (Mo and W) or 1.6-12.0 \AA (Fe) (10 % cos windows).

EPR spectroscopy

EPR samples were prepared either aerobically in an ice bath or anaerobically in a Coy Chamber at O_2 levels < 10 ppm at 4 °C in a thermoblock, in 100 mM Tris-HCl, pH 7.2. Typical EPR sample preparation methods involved addition of 20 μL of freshly-prepared sodium dithionite or potassium ferricyanide (10 mM final concentration) to 180 μL of as-isolated YdhV residing in a quartz EPR capillary (3.9 mm O.D.), followed by brief mixing and immediate freezing (5-10 seconds aerobically, 15-20 seconds anaerobically) in a liquid N_2 -cooled ethanol bath before final freezing in liquid N_2 . Anaerobic samples were additionally gas-tight sealed prior to freezing outside of the chamber. EPR sample anaerobicity is noted in Table S1.

CW X-band EPR spectra were obtained using a laboratory-built spectrometer (microwave bridge, ER041MR, Bruker; lock-in amplifier, SR810, Stanford Research Systems; microwave counter, 53181A, Agilent Technologies) equipped with a Bruker SHQ resonator. An ESR 910 helium flow cryostat with an ITC503 temperature controller (Oxford Instruments) was used for temperature control. A Cu(II)/EDTA standard was used as a reference for spin quantitation of YdhV samples. Spin quantitation was performed using the utility ‘spincounting’ (<http://lcts.github.io/spincounting/>) in Matlab (Mathworks). Spectral simulations were performed using EasySpin in Matlab.³⁹ Field corrections were applied as needed through measuring a reference $N@C_{60}$ sample at ambient temperature to give the expected spin Hamiltonian parameters as reported elsewhere.⁴⁰ Typical parameters for EPR data acquisition were (unless otherwise noted): temperature, 80 K; modulation amplitude, 5 G; microwave

power, 4.0 mW; microwave frequency, 9.38 GHz. Microwave power saturation data were fitted as a double logarithmic plot using OriginPro (version 9.1.0) to the equation:

$$S = K \frac{\sqrt{P}}{\left(1 + \frac{P}{P_{1/2}}\right)^{0.5b}}$$

where S is the signal intensity, K is a proportionality constant, P is the microwave power, $P_{1/2}$ is the microwave power at half-saturation, and b is the inhomogeneity parameter.⁴¹

RESULTS

YdhV protein expression and purification

For characterization of the YdhV protein, a homologous expression system was established in *E. coli*. For separate YdhV expression, the coding sequence was amplified from *E. coli* K12 total DNA and cloned into the vector pTrcHis, which results in an N-terminal fusion of the protein with a His₆-tag. Additionally, the other genes of the operon were cloned into the pACYC-duet1 vector, *ydhY* into MCS1 and *ydhWXUT* into MCS2, which enabled coexpression with YdhV. Both vectors or vector pTrcHis-YdhV alone were transformed into BW25113 cells and expressed as described in Materials and Methods. In both cases, Ni-NTA chromatography yielded only the YdhV protein and no further proteins (data not shown), so that for all further characterizations the pTrcHis-YdhV construct was used for protein expression. YdhV was expressed with a yield of 17 mg protein per liter of *E. coli* culture and was purified by Ni-NTA chromatography and size exclusion chromatography, resulting in a monomeric protein with a molecular mass closely resembling the calculated value of 77.8 kDa (Figures 2A and B). The optical absorption spectrum shows the characteristic features for an enzyme containing a molybdenum cofactor⁴², which was reducible by the addition of sodium dithionite (Figure 2C). Characteristic features corresponding to the [4Fe-4S] cluster predicted to be bound to the protein were not clearly discernable in the absorption spectrum (Figure 2C).

Unfortunately, so far we were unable to establish an activity assay for YdhV, since no activity could be detected with any of the numerous substrates that were tested (see Materials and Methods for details).

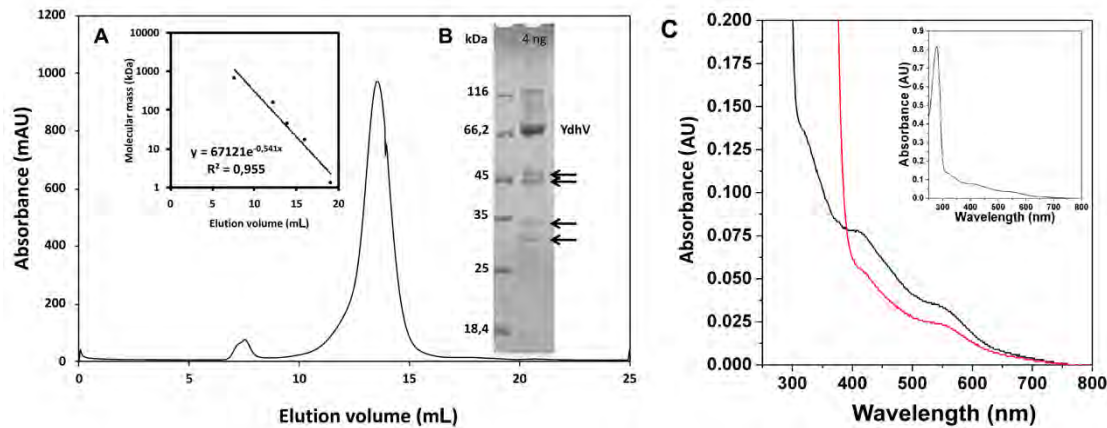


Figure 2: Purification of YdhV

(A) Size exclusion chromatography of YdhV. YdhV was analyzed by analytical size exclusion chromatography in 100 mM Tris and 200 mM NaCl (pH 7.2) using a Superdex 200 column. Inset: Relation of molecular weight and elution volume for standard proteins. Size exclusion chromatography markers (Bio-Rad): c-globulin (158 kDa), ovalbumin (44 kDa), myoglobin (17 kDa), and vitamin B12 (1.3 kDa). (B) 10% SDS polyacrylamide gel of YdhV after Ni-NTA chromatography. The bands marked by arrows were assigned to degradation products of YdhV by MALDI-TOF mass spectrometry. (C) UV-visible absorption spectra of 6.5 μ M YdhV in 100 mM Tris/HCl pH 7.2 (as-purified YdhV, black line, and reduced YdhV with 10-fold excess of sodium dithionite, red line.) The inset shows the overall spectrum of the as-purified protein.

Identification of the type of Moco present in YdhV

To analyze the type of Moco present in YdhV, we expressed the enzyme in different *E. coli* mutant strains. So far, only molybdoenzymes have been identified in *E. coli* and tungsten-containing enzymes were not expected to be produced. YdhV was expressed in the presence of 1 mM molybdate and the molybdenum and iron contents in the enzyme were determined. To identify whether YdhV binds a bis-MGD cofactor with additional guanine nucleotides as present in TMAO reductase (TorA), a MCD cofactor with a cytosine nucleotide as in PaoABC, or a Mo-MPT cofactor identified in MsrP, we quantified the metal, Moco and nucleotide content (as FormA-NMP) of YdhV in comparison to the mentioned enzymes serving as

controls (Table 1). The proteins were expressed in the BW25113 wild-type *E. coli* strain, $\Delta mobA$ and $\Delta mocA$ deletion strains impaired in bis-MGD or MCD biosynthesis, and in a $\Delta mobA/\Delta mocA$ double-deletion strain, respectively.

Table 1: Relative activity and metal/Moco loading of TorA, PaoABC, MsrP and YdhV.

TorA ^a	Mo ^b	Fe ^b	FormA ^c	FormA-NMP ^c	Relative activity ^d
BW25113	1.00 ± 0.03	n.d.	1.00 ± 0.11	1.00 ± 0.07	1.00 ± 0.04
$\Delta mobA$	0.60 ± 0.04	n.d.	0.08 ± 0.01	n.d.	n.d.
$\Delta mocA$	1.05 ± 0.02	n.d.	1.06 ± 0.08	1.00 ± 0.21	1.18 ± 0.06
$\Delta mobA\Delta mocA$	0.65 ± 0.03	n.d.	0.07 ± 0.01	n.d.	n.d.

PaoABC ^a	Mo ^b	Fe ^b	FormA ^c	FormA-NMP ^c	Relative activity ^d
BW25113	1.00 ± 0.03	1.00 ± 0.06	1.00 ± 0.07	1.00 ± 0.26	1.00 ± 0.07
$\Delta mobA$	0.55 ± 0.04	0.99 ± 0.04	0.56 ± 0.17	0.35 ± 0.16	0.47 ± 0.12
$\Delta mocA$	0.33 ± 0.04	1.01 ± 0.04	0.40 ± 0.20	n.d.	n.d.
$\Delta mobA\Delta mocA$	0.42 ± 0.03	0.97 ± 0.03	0.27 ± 0.16	n.d.	n.d.

MsrP ^a	Mo ^b	Fe ^b	FormA ^c	FormA-NMP ^c	Relative activity ^d
BW25113	1.00 ± 0.02	n.d.	1.00 ± 0.13	n.d.	1.00 ± 0.07
$\Delta mobA$	0.87 ± 0.03	n.d.	1.02 ± 0.07	n.d.	0.88 ± 0.04
$\Delta mocA$	0.91 ± 0.02	n.d.	0.91 ± 0.13	n.d.	1.19 ± 0.05
$\Delta mobA\Delta mocA$	0.98 ± 0.01	n.d.	1.51 ± 0.01	n.d.	1.34 ± 0.07

YdhV ^a	Mo ^b	Fe ^b	FormA ^c	FormA-NMP ^c	Relative activity ^d
BW25113	1.00 ± 0.02	1.00 ± 0.03	1.00 ± 0.09	n.d.	n.d.
$\Delta mobA$	0.71 ± 0.15	0.85 ± 0.04	2.23 ± 0.27	n.d.	n.d.
$\Delta mocA$	0.75 ± 0.02	0.88 ± 0.04	0.84 ± 0.07	n.d.	n.d.
$\Delta mobA\Delta mocA$	0.63 ± 0.02	0.84 ± 0.03	2.12 ± 0.23	n.d.	n.d.

^aThe enzymes were expressed in *E. coli* BW25113 and $\Delta mobA$, $\Delta mocA$ or $\Delta mobA/mocA$ deletion strains for comparison.

^bMetal contents (metal/enzyme) were determined by ICP-OES and normalized to the values for enzymes purified from the BW25113 wild-type strain set to 1.

^cThe Moco content of the enzymes was quantified after conversion of MPT or MPT-NMP to FormA or FormA-NMP and normalized to the values for enzymes purified from the BW25113 wild-type strain set to 1.

^dEnzyme activities were determined with various substrates in units/mg and normalized to the values for enzymes purified from the BW25113 wild-type strain set to 1.

n.d. = not detectable.

Data represent the mean values from at least three independent measurements (± standard deviations).

The controls showed the expected behavior, with TorA being inactive in the $\Delta mobA$ and $\Delta mobA/\Delta mocA$ strains impaired in bis-MGD synthesis (no FormA or FormA-GMP detectable),

while a *mocA* deletion had no effect on TorA. PaoABC was inactive in the $\Delta moca$ and $\Delta mobA \Delta moca$ strains (no FormA-CMP detectable). For MsrP no impairment in activity was detected in all mutant strains and the data were comparable to wild-type levels. For YdhV, we also did not detect the presence of FormA-NMP in any of the analysed strains. The molybdenum and iron contents of YdhV expressed in the deletion strains were 80-100% of the wild-type strain. While the FormA levels in the $\Delta moca$ strain were comparable to the wild-type strain, an up to two-fold higher FormA content was detected in $\Delta mobA$ and $\Delta mobA/\Delta moca$ mutant strains. This shows that the Moco, which is assembled and inserted in YdhV, does not depend on the activity of MobA or MocA. The absence of FormA-NMP indicates that likely no nucleotide is attached to the Moco of YdhV and consequently a Mo-MPT cofactor has to be bound to the enzyme.

After identification of a nucleotide-free Moco form in YdhV, we studied whether Mo-MPT, as present in MsrP, or a bis-MPT cofactor form containing two MPT moieties, as present in the homologous tungsten-containing enzymes AOR and BamBC, is bound to YdhV. In the past, we have identified a bis-Mo-MPT cofactor in MobA, e.g., by determination of the molybdenum to phosphorous ratio in the protein. The expected P:Mo ratios for the bis-MPT and MPT cofactors are 2:1 and 1:1, respectively.⁷ For comparison, here we applied two different methods to determine the P:Mo ratio in YdhV (Table 2). Quantification by ICP-MS revealed a P:Mo ratio close to 2:1. This ratio was confirmed by P quantification using TXRF (Table S2) in relation to the Mo content from ICP-OES, which gave a ratio of about 2.2:1. Based on these results a bis-Mo-MPT cofactor is most likely present in YdhV.

Table 2: Phosphorus to **molybdenum ratio in YdhV**.

Sample	P [per protein] : Mo [per protein]
YdhV ^a	2.07 (± 0.58) ^b : 1 (± 0.03) ^b
YdhV ^a	2.17 (± 0.22) ^c : 1 (± 0.07) ^d

^a1.64 μM YdhV was used for ICP-MS and 3.175 mM YdhV for TXRF and 50 μM YdhV for ICP-OES analyses. The molybdenum saturation in the proteins was normalized to 1.

^bMo and P were determined by ICP-MS

^cP contents were determined by TXRF (Table S2).⁷

^dMo was quantified by ICP-OES.⁷

Specificity of YdhV towards molybdenum or tungsten

Since a bis- MPT cofactor was so far identified only in enzymes that contain a tungsten ion, the ability of YdhV in *E. coli* to bind tungsten versus molybdenum was assessed. Competitive Mo vs. W binding to YdhV was studied by expression of YdhV in the presence of 10 μM molybdate and increasing tungstate concentrations or vice versa. After purification of YdhV, the Mo and W contents in the enzyme were quantified (Figure 3). With 10 μM molybdate, YdhV showed ~20 % Mo loading. The molybdenum saturation of YdhV increased up to 65 % when 100 μM tungstate were additionally present in the medium during expression, likely based on a better uptake of molybdate into the cells (Figure 3A). In contrast, the tungsten loading was rather small (~10 %) after expression with 10 μM molybdate and 100 μM tungstate, whereas with 10 μM tungstate in the absence of molybdate, the tungsten levels reached 20% (Figure 3B). However, with increasing molybdate concentrations during expression, the tungsten content decreased, meaning that W binding to the enzyme was outcompeted by Mo binding due to an apparently higher specificity for Mo insertion (Figure 3B). These findings show that when Mo is available even at low concentrations, the bis-Mo-MPT cofactor is preferentially inserted into YdhV.

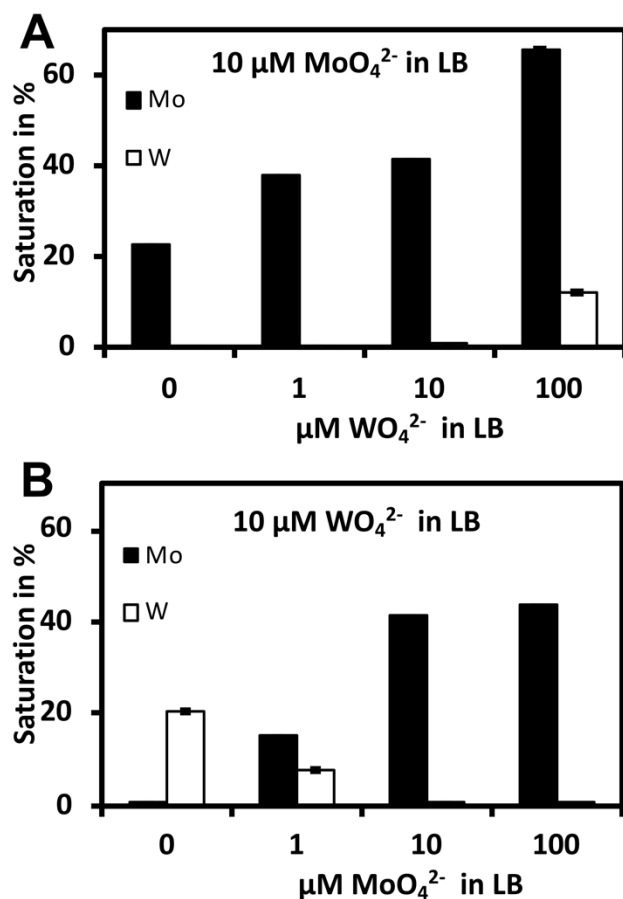


Figure 3: **Competition of molybdenum versus tungsten insertion into YdhV.** Expression of YdhV in the presence of 10 μM Mo in LB with increasing W concentrations (A). Expression of YdhV in the presence of 10 μM W in LB with increasing Mo concentrations (B).

Mo site structure in YdhV from XAS

XAS was employed to characterize the metal site structure in YdhV expressed with molybdenum. Figure 4 shows XAS spectra at the Mo K-edge. We compared the as-isolated (oxidized) enzyme purified under aerobic conditions (YdhV^{ox}) to the dithionite-reduced enzyme (YdhV^{red}). The XANES of YdhV^{ox} showed a resolved pre-edge feature due to resonant 1s \rightarrow 4d electron excitation, which almost disappeared in YdhV^{red} (Figure 4A). In addition, YdhV^{ox} showed a considerably higher K-edge energy (\sim 0.5 eV at edge half-height), which indicates significantly more oxidized Mo compared to YdhV^{red}. The K-edge energy of YdhV^{ox} (\sim 20012.8 eV) and the relatively small pre-edge feature are in good agreement with predominantly sulfur-coordinated Mo(VI).^{7, 30, 36} The edge energy downshift due to Mo

reduction presumably is superimposed by a counter-directed edge shift due to coordination changes at Mo, as indicated by the pre- and main-edge shape changes and the EXAFS analysis below, meaning that the metal was reduced at least to the Mo(V) if not to the Mo(VI) level in YdhV^{red}. Redox cycling of YdhV (air re-oxidation of YdhV^{red} leading to YdhV^{reox} and NaDT re-reduction of YdhV^{reox}, leading to YdhV^{rered}) caused respective increase and decrease of the pre-edge feature and similar K-edge energy changes in YdhV^{reox} or YdhV^{rered}, emphasizing the reversibility of the redox events at the Mo center.

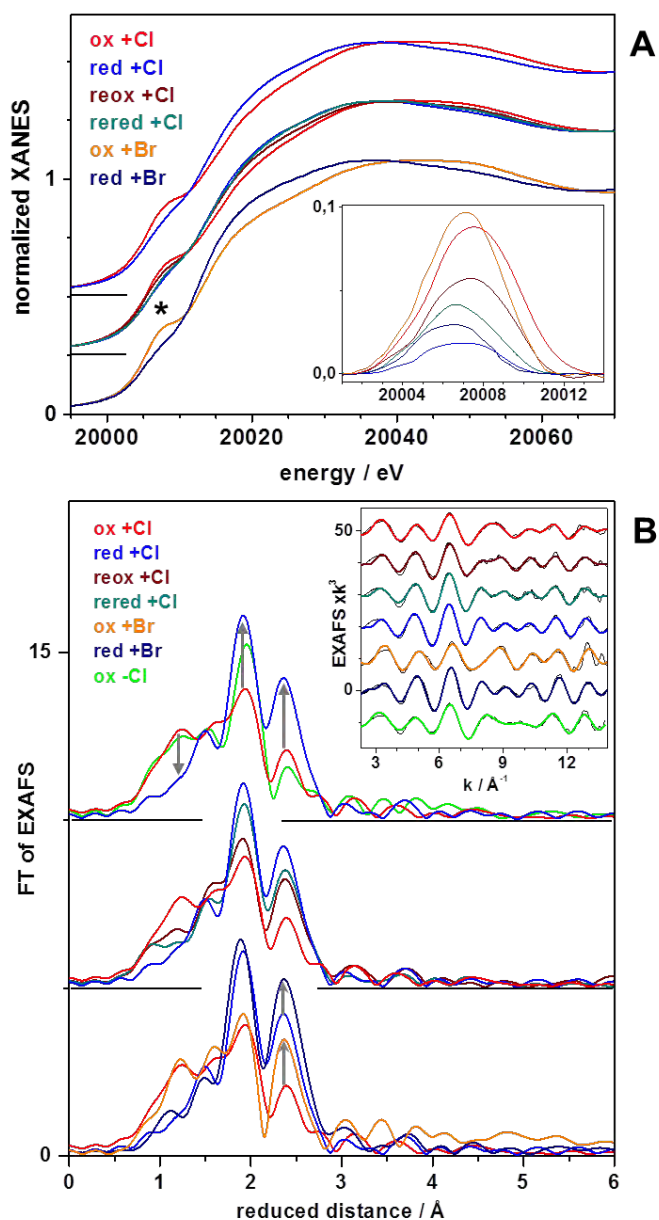


Figure 4: **XAS analysis of YdhV.** (A) Mo K-edge spectra (XANES) of (top) as-isolated, oxidized (ox) or sodium-dithionite reduced (red) enzyme in Cl⁻-containing buffer, (middle) of the latter enzyme samples as well as of enzyme after re-oxidation (reox) followed by re-reduction (rered) in Cl⁻-containing buffer, and (bottom) of oxidized or reduced enzyme in Br⁻-containing buffer. The inset shows the isolated pre-edge features (asterisk) after removal of the main edge rise. (B) Experimental Fourier-transforms (FTs) of EXAFS spectra in the inset of proteins as in (A) and of oxidized YdhV in Cl⁻-free buffer (inset: black lines, experimental data; colored lines, simulations with parameters in Table 3). Arrows emphasize spectral alterations due to the redox or anion changes.

EXAFS analysis was performed to determine the coordination of the Mo ion in the YdhV preparations. The Fourier transform (FT) of the EXAFS spectrum of YdhV^{ox} revealed

at least three peak features, suggesting a complex Mo ligation pattern. For YdhV^{red}, pronounced spectral changes occurred, such as vanishing of the lowest-distance FT peak due to Mo=O bonds and increasing of the two higher-distance FT peaks, which indicates significant coordination changes at the metal. The relative FT amplitude patterns and changes were well reproduced in the redox-cycled YdhV. EXAFS simulations provided coordination numbers and interatomic distances in the first and second Mo ligation spheres (Table 3).

Table 3: **Mo EXAFS fit parameters of YdhV.**^a

YdhV preparation	Mo=O,-OH _n	Mo=S,-SH	Mo-S _{MPT/Cys}	Mo---Cl,Br	R _F
	N [per Mo] / R [Å] / 2σ ² x10 ³ [Å ²]				[%]
ox +Cl	0.7 [#] / 1.68 / 2* 0.4 [#] / 1.80 / 2*	0.3 [#] / 2.17 / 2*	4.0* / 2.34 / 21 0.6 [#] / 2.56 / 2*	0.5 / 2.88 / 2*	12.4
red +Cl	0.3 [#] / 1.70 / 2* 0.8 [#] / 2.23 / 2*	0.4 [#] / 2.35 / 2*	4.0* / 2.37 / 14 0.5 [#] / 2.53 / 2*	1.2 / 2.86 / 2*	9.8
rered +Cl	0.5 [#] / 1.70 / 2* 0.6 [#] / 2.14 / 2*	0.4 [#] / 2.29 / 2*	4.0* / 2.39 / 19 0.5 [#] / 2.55 / 2*	1.0 / 2.87 / 2*	11.2
reox +Cl	0.6 [#] / 1.69 / 2* 0.3 [#] / 1.80 / 2*	0.3 [#] / 2.18 / 2*	4.0* / 2.35 / 17 0.8 [#] / 2.54 / 2*	0.9 / 2.87 / 2*	9.6
ox -Cl	0.8 [#] / 1.69 / 2* 0.2 [#] / 1.81 / 2*	0.4 [#] / 2.17 / 2*	4.0* / 2.36 / 16 0.6 [#] / 2.55 / 2*	0.2 / 2.84 / 2*	11.5
ox +Br	0.8 [#] / 1.69 / 2* 0.2 [#] / 1.78 / 2*	0.2 [#] / 2.17 / 2*	4.0* / 2.33 / 19 0.8 [#] / 2.54 / 2*	0.3 / 2.69 / 2*	12.6
red +Br	0.4 [#] / 1.70 / 2* 0.8 [#] / 2.19 / 2*	0.3 [#] / 2.27 / 2*	4.0* / 2.38 / 19 0.5 [#] / 2.49 / 2*	0.7 / 2.67 / 2*	13.4

^aN, coordination number; R, interatomic distance; 2σ², Debye-Waller factor; R_F, fit error sum (deviation between Fourier back-transform in a 1-3 Å reduced distance window and simulation curve). Fit restraints: *fixed parameter, #N-values coupled to yield a sum of 2. Parameters correspond to EXAFS simulation curves in the inset of Figure 4B.

First, we searched for a consistent simulation approach that described all spectra well and limited the number of free parameters to gain statistical significance. Unrestraint fits of all spectra provided Mo-S coordination numbers (*N*) in the range of 3.5-4.1 (not shown). Taking this result and a bis-Mo-MPT structure into account, the *N*-value of the Mo-S_{MPT} shell was restraint to 4. A presumably related cofactor structure in W-containing benzoyl-CoA reductase (BamBC) shows a cysteine bound at the metal, which is also present in the YdhV sequence

(Figure S1A), and an oxo ligand (Mo=O). In other DMSO-reductase family enzymes, a sulfido ligand (Mo=S) at the metal was found.^{21, 22, 36, 43-45} Accordingly, Mo=O/S distances as well as Mo-S_{Cys} and alternative water or oxygen species were included in the simulations. YdhV prepared in a Br⁻ instead of Cl⁻ containing buffer revealed pronounced EXAFS spectral changes, in particular increased highest-distance FT peaks (Figure 4), so that a halide in the second Mo coordination sphere was further included. We note that S and Cl⁻ cannot be discriminated in EXAFS simulations due to their comparable atomic weight, but the heavier backscatterer Br⁻ is well discernable from lighter Cl⁻ species. Table 3 summarizes the simulation results, which show good fit qualities for all spectra as judged by a mean error sum (R_F) around 10 %.

YdhV^{ox} revealed ~0.7 Mo=O (~1.7 Å) and ~0.3 Mo=S (~2.2 Å) interactions with typical bond lengths,³⁵ close to 0.5 Mo-OH (~1.8 Å) and Mo-S_{Cys} (~2.6 Å) bonds, and ≤0.5 Cl⁻ in the 2nd coordination sphere (~2.9 Å), besides of the 4 Mo-S_{MPT} (~2.35 Å) bonds (Table 3). In YdhV^{red}, the number of Mo=O bonds decreased in favor of longer Mo-OH_n (~2.2 Å) bonds, short Mo=S bonds were replaced by longer Mo-S bonds (~2.3 Å), the number of Mo-S_{Cys} bonds was diminished, and close to 1 Cl⁻ was detectable. YdhV^{reox} and YdhV^{rered} revealed similar, but less pronounced coordination number and bond lengths changes, including gain of longer Mo-S/O bonds at the expense of shorter Mo=O/S bonds and of the Cl⁻ coordination number in the more reduced enzyme (Table 3). YdhV^{ox} prepared in the absence of halides showed a diminished Cl⁻ coordination number and an otherwise similar Mo coordination (Table 3). TXRF on YdhV prepared with Br⁻ instead of Cl⁻ revealed ~0.5 Br⁻ per protein (Figure S2, Table S2). The EXAFS simulations of the spectra of oxidized or reduced YdhV prepared with Br⁻ showed similar first-sphere coordination and ligation changes as the Cl⁻ containing proteins, including higher numbers of Mo=S and Mo-S_{Cys} bonds in the more oxidized enzyme (Table 3). However, about 0.3 or 0.7 Br⁻ ions at closer distance to Mo (~2.7 Å) vs. Cl⁻ were discernable

in the oxidized or reduced proteins, in agreement with (partial) replacement of Cl⁻ by Br⁻ in the second Mo coordination sphere.

A bis-Mo-MPT cofactor in YdhV is further supported by the Mo-S_{MPT} bond numbers and the bond lengths range similar to the highly distorted cofactors in (W) BamBC and AOX crystal structures.^{16, 46} The YdhV preparations likely contain a mixture of Mo site configurations due to varying amounts of oxidized or reduced and of native or modified cofactors. Detection of minor, but reproducible Mo=S bond numbers in oxidized YdhV suggests that a sulfido ligand is present in the native YdhV. The sulfido group is largely replaced by an oxo group in aerobically prepared protein, which further may show cysteine to hydroxo ligand exchange. The sulfido vs. oxo exchange resembles the situation for another DMSOR family enzyme, formate dehydrogenase.³⁶ A cysteine ligand seems to be bound to Mo in oxidized YdhV, which may be detached from the metal and replaced, for example, by a water or another amino acid species (O_x) in reduced enzyme. Mo site reduction further causes elongation of sulfur and oxygen bonds, which can be explained by ligand protonation to yield Mo-OH (1.9 Å³⁵) or Mo-SH (~2.4 Å^{47, 48}) instead of Mo=O or Mo=S bonds. In addition, Mo site reduction seems to cause appearance of a halide ion in the 2nd coordination sphere related to cysteine detachment, oxygen species binding, and ligand protonation, which may provide hydrogen-bonding interactions to the halide. Notably, a halide near Mo may be incorporated only in the absence of a substrate. In summary, our results suggest that native, oxidized YdhV contains a S=Mo(S_{cys})(S_{MPT})₄ site with Mo^{VI}, which is converted to a HS-Mo(O_x)(S_{MPT})₄ site presumably with Mo^{IV} in reduced enzyme (Figure 5). In modified YdhV, a O=Mo(OH)(S_{MPT})₄ site in oxidized enzyme seems to be converted to a HO-Mo(OH₂)(S_{MPT})₄ site in reduced enzyme.

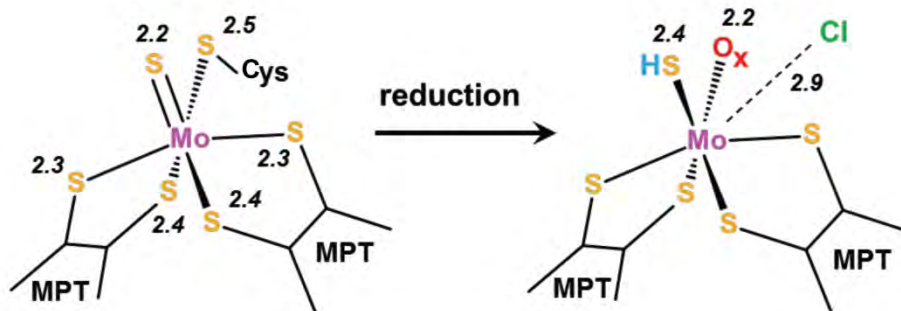


Figure 5: **EXAFS-derived molybdenum site structures in YdhV.** Schematic drawings tentatively adopt an overall cofactor geometry as in crystal structures of bis-W-MPT-Mg cofactors in BamBC and AOX. Selected (rounded) interatomic distances (in Å) from XAS are indicated. The Mo site in native, oxidized YdhV is proposed to bind sulfido and cysteine (Cys336 in *E. coli* YdhV) ligands. Sulfur protonation, oxygen against cysteine replacement, and remote halide (Cl⁻ or Br⁻) binding (at an unknown position around the metal) occur upon enzyme reduction; the oxygen species (O_x) may stem from water (H₂O) or an amino acid residue (e.g. Thr458 in *E. coli* YdhV). Oxidative modification (i.e. after aerobic enzyme purification) leads to sulfido to oxo and cysteine to hydroxo group exchange at Mo and to further oxygen protonation and/or O_x as well as halide binding in reduced vs. oxidized enzyme.

XAS on tungsten-substituted YdhV

XAS spectra of tungsten-substituted, anaerobically purified YdhV at the W L₃-edge are shown in Figure 6. The W XANES depended strongly on the W concentration during protein expression. A ~1 eV lower edge energy for YdhV expressed with 10 μM W vs. 1 mM W and edge shape changes suggested pronounced coordination changes at the metal. The edge shape for YdhV expressed with 1 mM W indeed was more similar to tungstate in solution (Figure 6A, see below). The EXAFS spectrum of YdhV expressed with 10 μM W revealed a major FT peak due to W-S bonds and minor W=O contributions (Figure 6B). Simulation of the EXAFS spectrum using a similar approach as for the Mo data revealed ~0.5 W=S (~2.2 Å), 4 Mo-S_{MPT} (~2.4 Å) and close to 1 Mo-S_{cys} (~2.5 Å) bonds, besides of W=O/-OH bonds. Cl⁻ in the second coordination sphere was barely detectable (Table S3). A more homogenous metal ligation in W vs. Mo YdhV was supported by the smaller Debye-Waller factor ($2\sigma^2$) of the W-S_{MPT} vs. Mo-S_{MPT} bonds. Notably, the W EXAFS of YdhV was reproducible in several protein preparations, for cell growth in media with varying tungstate concentrations ≤250 μM, or

increased salt content in the purification buffer (data not shown). These results suggest that native YdhV both in the W and Mo forms shows a fully sulfurated metal site. Oxidative modification was less pronounced for bis-W-MPT samples, facilitating more clear detection of sulfido and cysteine ligands at the tungsten center.

Interestingly, the EXAFS spectrum of YdhV at pH 7.2 expressed with 1 mM W closely resembled the spectrum of tungstate in aqueous solution at pH 4.0 (Figure 6B). EXAFS analysis of Na_2WO_4 solutions showed 4 W=O bonds at pH 10.6, protonation of one ligand at pH 7.1, but spontaneous tungsten oxide (polytungstate) formation, as evident from W-W and long W-O distances in the EXAFS, only at pH 4.0 (Table S3). For YdhV with 1 mM W, no tungsten-sulfur bonds were detectable, which showed the absence of MPT ligands at the metal, but comparable W=O/-OH_n and W-W distances as for polytungstate were indicated by the EXAFS simulation. These findings implied that polytungstate formation occurs in the YdhV protein even at neutral pH. Presumably, in the presence of super-stoichiometric W concentrations during protein expression, the cofactor binding site is occupied by several tungstate molecules, which promotes tungsten-oxide formation in YdhV.

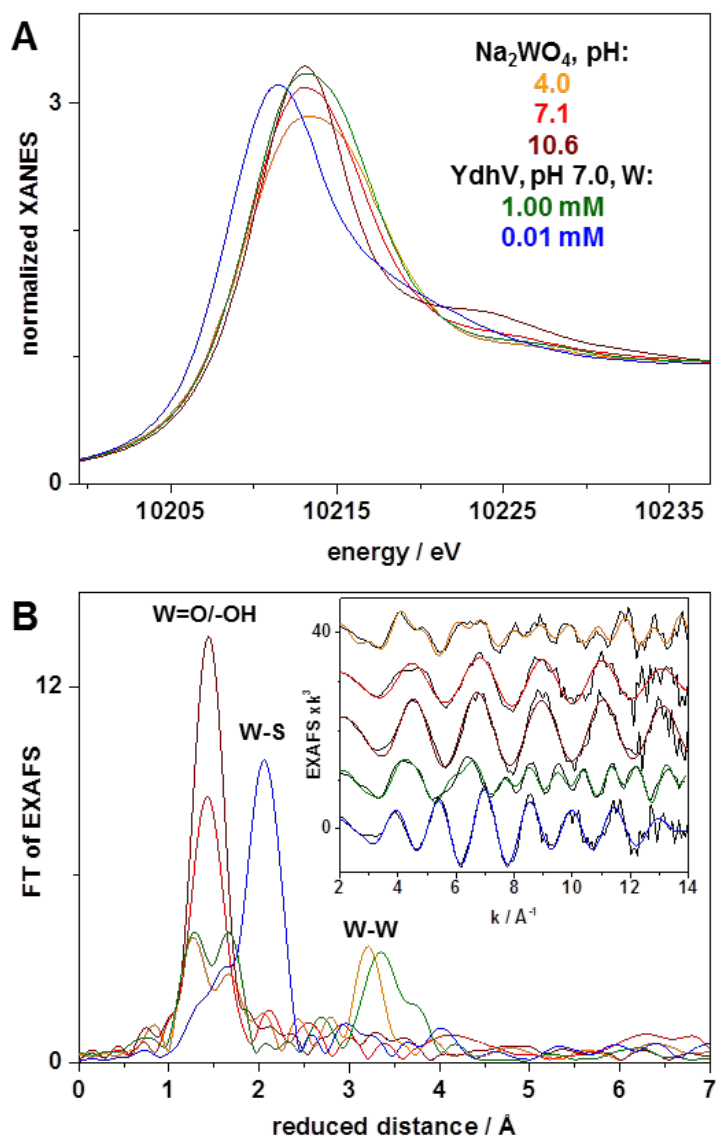


Figure 6: **W XAS of YdhV.** (A) W L₃-edge spectra of YdhV (pH 7.2) expressed with 10 μ M or 1 mM tungsten (anaerobic purification) compared to spectra of aqueous Na₂WO₄ solutions at indicated pH values. (B) Fourier-transforms (FT) of EXAFS spectra in the inset of samples as in (A) (black lines, experimental data; blue lines, fits with parameters in Table S3; dominant contributions of interatomic distances to the FT peaks are marked).

EPR characterization of YdhV

To characterize the YdhV protein further, paramagnetic cofactor species were studied by EPR spectroscopy. EPR spectra of Mo(V) in bis-Mo-MPT loaded YdhV are shown in

Figure 7A. As-isolated and dithionite-reduced YdhV showed an isotropic signal representing less than 0.05 spins/YdhV. This signal was more prominent for purified YdhV expressed in the $\Delta moaA$ deletion strain (apo-YdhV) in the presence of molybdate, accounting for about 0.1 spins/YdhV or 0.08 spins/Mo in the as-isolated state (Figure S3A; Table S1). Since binding of a bis-Mo-MPT can be excluded in the apo-YdhV variant, the isotropic signal is reminiscent to paramagnetic polymolybdate species reported for the tungstate transporter WtpA.⁴⁹ Therefore, the isotropic signal represents Mo(V) bound to YdhV in a non-bis-Mo-MPT form (Figure S3A). Spin quantification of the isotropic signal in the as-isolated, bis-Mo-MPT loaded YdhV revealed approximately 56 μM Mo(V), corresponding to $\sim 15\%$ of the total molybdenum (Figure 7A, Table S1). Interestingly, upon oxidative treatment with ferricyanide, a more prominent and different signal could be detected, representing a spin concentration of ~ 0.08 spins/YdhV at 80 K and of ~ 0.20 spins/YdhV at 150 K (Figure 7A, Table S1; Figure S4). We assign this signal to Mo(V) in the bis-Mo-MPT cofactor (see below).

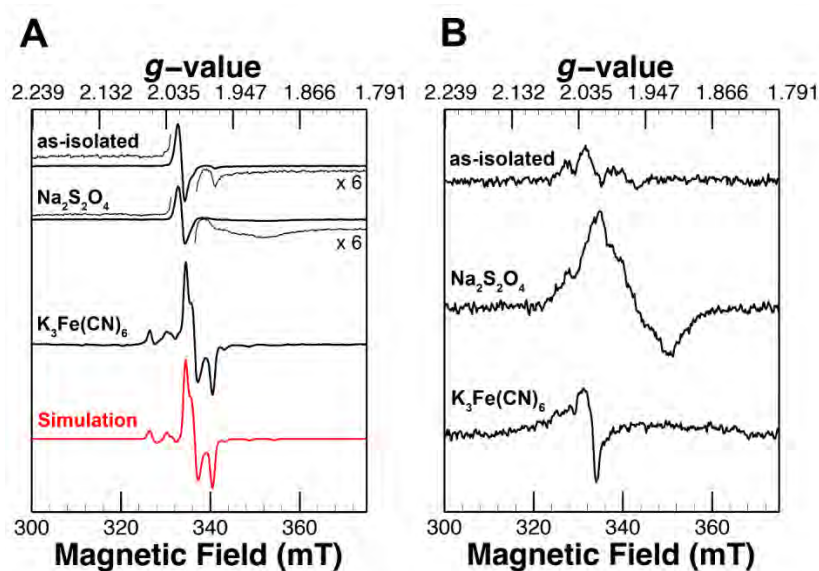


Figure 7: **X-band EPR spectra of YdhV.** Anaerobically expressed and purified YdhV was employed. (A) YdhV expressed in the presence of 1 mM Na₂MoO₄, (B) YdhV expressed in the presence of 10 μM Na₂WO₄. Spectral simulation of Mo(V) detected in ferricyanide-oxidized YdhV. The black spectra depict YdhV as-isolated (top), after treatment with 10 mM sodium dithionite (middle), and after treatment with K₃Fe(CN)₆ (bottom); the red spectrum in panel A represents a spectral simulation of the ferricyanide-oxidized spectrum. Features present in the baseline are enlarged as noted (thin lines). Spectra were recorded at 80 K (4 mW microwave power, 5 G modulation amplitude, 100 kHz modulation frequency). Spectra in

panel A represent a YdhV concentration of 202 μM ; spectra in panel B were scaled to comparable amplitudes for comparison. For associated concentrations of Mo and Fe, see Table S1:

By contrast, in EPR spectra of bis-W-MPT loaded YdhV no W(V) was detected (Figure 7B). These results possibly reflect a less redox-active bis-W-MPT as opposed to a more redox-active bis-Mo-MPT cofactor.

A spectral simulation of the Mo(V) signal in ferricyanide-treated YdhV is shown in Figure 7A (see also Figure S4). The simulation parameters ($g = 2.0082, 1.9957, 1.9724$; $g_{av} = 1.9915$) yielded a tensor reminiscent in anisotropy and rhombicity to parameters reported for the glycerol-inhibited and “high-g unsplit” Mo(V) states detected in *R. sphaeroides* DMSO reductase (Table S4).^{50, 51} Inclusion of a Mo(V) component due to unspecific molybdenum was not necessary to achieve a good simulation and resolved ^1H hyperfine couplings were not observed.^{35, 37} Inclusion of chloride hyperfine coupling also did not improve the simulation. The $^{95, 97}\text{Mo}$ hyperfine tensor ($A^{\text{Mo}} = [75.7, 152.5, 6.1]$ MHz) exhibits rhombic character. Rhombic $^{95, 97}\text{Mo}$ hyperfine tensors typically show a lowered point group symmetry due to trigonal-prismatic metal coordination through distortion of the Mo-S_{MPT} from a formally trigonal-prismatic metal coordination geometry.⁵¹ In turn, the simulated hyperfine parameters for the Mo(V) bis-Mo-MPT center are reflective of a low-symmetry structure.

The power saturation behavior of the Mo(V) species was studied (Figure 8A). The Mo(V) signal (at 12 K) of ferricyanide-treated YdhV exhibited a half-saturation power ($P_{1/2}$) of 90 mW, meaning that the signal was very difficult to saturate. While an interacting Fe-S cluster can be ruled out as a reason for relaxation enhancement of the Mo(V) signal since the cluster in YdhV (see below) was oxidized, apparent coupling to ferricyanide ions may cause the large $P_{1/2}$ (Figure S5, panel A). Indeed, removal of ferricyanide from the YdhV sample resulted in a dramatically slowed relaxation of the Mo(V) signal at 12 K without significant change in shape of signal, which now showed an about 135-fold smaller $P_{1/2}$ value (670 μW).

The latter value is similar to the Mo(V) signal of *E. coli* DMSO reductase in the presence of an oxidized [4Fe-4S] cluster.⁵² These observations are consistent with the binding of ferricyanide ions in proximity to molybdenum in YdhV, resulting in an apparent Fe(III)-Mo(V) electron spin-spin interaction.

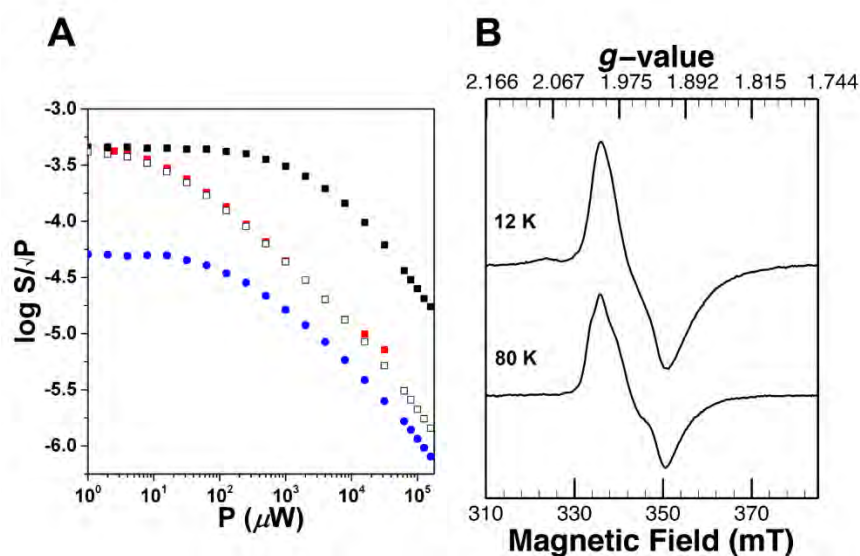


Figure 8: **Relaxation behavior of paramagnetic species in YdhV.** Panel A depicts the power saturation properties of Mo(V)-associated (squares) and Fe-S cluster-associated (circles) EPR signals recorded at 12 K (10 G modulation amplitude, 100 kHz modulation frequency). Solid squares represent the intensity of the bis-Mo-MPT Mo(V) signals (Figure 7, panel A) recorded in the presence (black squares) or absence (red squares) of $\text{K}_3\text{Fe}(\text{CN})_6$ (measured at 334.5 mT). Open squares represent the intensity of the signal of unspecific Mo(V) in apo-YdhV expressed in the presence of molybdate (measured at 335.0 mT; Figure S3, panel A). Solid circles represent the intensity of the EPR signal at 12 K attributable to an Fe-S cluster in YdhV (measured at 336.0 mT; Figure S3, panel B). Panel B shows EPR signals at two temperatures of apo-YdhV aerobically expressed without Na_2MoO_4 and reduced with 10 mM sodium dithionite (see also Figure S3, panel B) (4 mW microwave power, 10 G modulation amplitude, 100 kHz modulation frequency). Shown spectra are baseline-corrected. For concentrations of Mo and Fe in the EPR samples, see Table S1.

Fe XAS analysis

In addition to molybdenum, YdhV was found to also bind about 2-3 Fe ions per protein (Table S1). Sequence alignment revealed a 46% amino acid sequence homology of YdhV with the benzoyl-CoA reductase BamBC from *G. metallireducens* (Figure S1A) particularly in that the YdhV residues Cys307, Cys310, Cys314, and Cys558 are similar to those shown to

coordinate a [4Fe-4S] cluster in proximity to the bis-W-MPT-Mg cofactor in the crystal structure of BamBC (Figure S1B).¹⁷ To identify the Fe-S cluster type of YdhV, XAS data at the Fe K-edge were collected (Figure 9). The XANES spectrum shows the typical shape of predominately sulfur-coordinated, tetrahedral iron.^{53, 54} The EXAFS Fourier-transform displays two main peak features mainly due to Fe-S and Fe-Fe distances. Simulation of the EXAFS spectrum revealed less than 4 Fe-S bonds (~ 2.2 - 2.4 Å), about 1 Fe-O bond (~ 2.5 Å), and ~ 1.6 Fe-Fe distances (~ 2.7 Å), as well as longer Fe-C distances per iron ion (Table S5). The interatomic distances are typical for iron-sulfur clusters, but the coordination numbers deviate from the values expected for a [4Fe-4S] cluster (4 Fe-S bonds, 3 Fe-Fe distances) or a [2Fe-2S] cluster (4 Fe-S bonds, 1 Fe-Fe distance). Taking into account the sub-stoichiometric Fe content, the detectable Fe-O bonds, and the presence of four cysteines in YdhV, it seems likely that the Fe EXAFS represents an overlay of partially assembled and (oxidatively) degraded [4Fe-4S] clusters, explaining the non-integer Fe-Fe coordination number and fractional oxygen-ligation of iron.

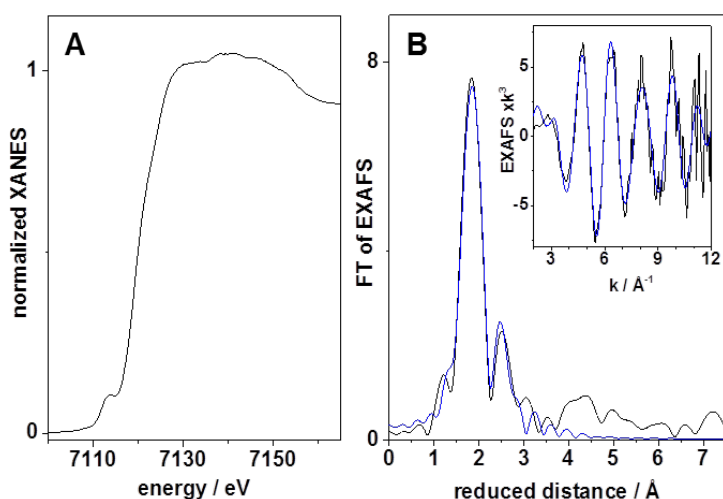


Figure 9: **Fe XAS of YdhV.** (A) Fe K-edge spectrum. (B) Fourier-transform (FT) of EXAFS spectrum in the inset (black lines, experimental data; blue lines, simulation with parameters in Table S5; the first FT peak reflects Fe-S,O bonds and the second peak Fe-Fe,C distances). The FT was calculated for a k-range of 2-12 Å⁻¹ (cos windows of 10 % at both k-range ends).

EPR spectroscopy of the FeS cluster present in YdhV

EPR spectroscopy was employed to further characterize [4Fe-4S] cluster binding to YdhV. To unambiguously differentiate spectral features associated with an Fe-S cluster, the apo-YdhV variant expressed in the absence of molybdate was characterized, that exhibited comparable Fe loading and substantially decreased Mo loading relative to bis-Mo-MPT-loaded YdhV (Table S1). Dithionite-reduced apo-YdhV revealed a broad EPR signal at 12 K and 4 mW microwave power, but did not show the isotropic signal observed for bis-Mo-MPT-loaded YdhV measured under similar conditions (Figure 8B, see also Figure S3B). To identify the Fe-S cluster type bound to apo-YdhV, EPR power saturation and temperature dependence studies were performed. EPR power saturation measurements at 12 K yielded a $P_{1/2}$ of 1.5 mW, a value qualitatively similar to [2Fe-2S] clusters (Figure 8A); by comparison, the 12 K and 80 K spectra appeared similar with modest changes in the signal lineshape and similar spin concentrations (~ 0.08 - 0.09 spins/YdhV) were observed under power non-saturating conditions (Table S1). These observations, in conjunction with the lowered Fe content (Table S1), XAS results, and featureless optical absorption spectra (Figure 2C) are consistent with an apparent [2Fe-2S] cluster of low occupation in YdhV. Likely, such a cluster reflects a species that has resulted from partial degradation of a [4Fe-4S] cluster as favored by the amino acid sequence of YdhV.

DISCUSSION

The variety of molybdoenzymes seems to be more versatile than originally expected, since we identified a novel bis-Mo-MPT cofactor without attached nucleotides in *E. coli* YdhV. A bis-Mo-MPT cofactor was consistently indicated by elemental analysis, nucleotide detection methods, and XAS. So far, the insertion of tungsten into the MPT cofactor of various molybdoenzymes was assumed to occur only unspecifically in *E. coli* (e.g. under high tungstate concentrations of 1 – 5 mM during expression). Our results reveal assembly of a bis-W-MPT cofactor in YdhV even for low tungstate concentrations (10 μ M). However, when both Mo and W were present, molybdenum is preferred over tungsten binding in YdhV. We therefore conclude that YdhV binds preferentially molybdenum so that under typical *E. coli* growth conditions, a bis-Mo-MPT cofactor is bound to the enzyme. The bis-Mo-MPT cofactor revealed to be redox active both in XAS and EPR spectroscopy, featuring (likely predominant) Mo(IV) and (minor) Mo(V) species, whereas EPR showed no indication for W(V) species of the bis-W-MPT cofactor. These findings suggest that a Mo(V) species may be involved as a semi-stable state in the two-electron reduction of the bis-Mo-MPT cofactor whereas the bis-W-MPT cofactor is either redox-inert or a W(V) species is less stable than the Mo(V) species.

In previous studies by our group, a bis-Mo-MPT cofactor was identified as an intermediate in the synthesis of the bis-MGD cofactor in *E. coli*, a reaction that is catalyzed by the MobA protein. In bis-MGD synthesis, two Mo-MPT molecules are fused on MobA to form the bis-Mo-MPT structure before nucleotide addition. For this reaction the ligation of molybdenum to MPT is essential.¹³ After formation of bis-Mo-MPT, two equivalents of GMP (from GTP) are added to the C4' phosphate of each MPT and the bis-MGD cofactor is formed. Our present results, however, reveal that formation of the bis-Mo-MPT cofactor in YdhV is independent of MobA. Also, the homologous protein MocA, which is involved in MCD cofactor formation in enzymes of the XO family, is likewise not involved in bis-Mo-MPT

synthesis in YdhV. This situation might indicate that either other proteins in *E. coli* are involved in bis-Mo-MPT formation or that this cofactor is directly formed on YdhV. So far, no function has been assigned for the two proteins YdhW and YdhT also encoded by the *ydhYVWXUT* operon. Amino acid sequence alignments show limited sequence identities of YdhW and YdhT with MobA. Thus, it is possible that YdhW and YdhT represent two proteins that act in conjunction in the formation of the bis-Mo-MPT cofactor in YdhV.

Despite extensive experimental efforts, we were unable to assign any substrate to YdhV. Previous studies by our group showed that the *ydhYVWXUT* operon is induced under anaerobic conditions, pointing to a role in anaerobic respiration.⁵⁵ Using XAS and EPR we demonstrated that the molybdenum atom in the bis-Mo-MPT cofactor of YdhV shows reversible redox activity. Our XAS results suggest that the Mo coordination sphere in the bis-Mo-MPT cofactor comprises a similar cysteine ligand as present in the homologous benzoyl-CoA reductase BamBC, in addition to a sulfido ligand. However, the sulfido ligand is easily replaced by an oxo ligand when YdhV is aerobically purified. In tungsten-containing YdhV an enhanced sulfido ligand content was observed. These findings suggest that the native YdhV active site contains a Mo-ligation sphere resembling that of formate dehydrogenases, which also bind either tungsten or molybdenum as well as cysteine and sulfido ligands at the metal in the bis-MGD cofactor (Figure 5). We note that the nature of the sixth ligand at the metal in the homologous enzyme BamBC remains to be established while cysteine ligation is visible in the crystal structure. A recent theoretical investigation disfavored a sulfido ligand in BamBC and favored an oxo ligand.⁵⁶ However, such a coordination sphere might be unique to BamBC since the particular Birch reaction catalysed by this enzyme so far is unique among molybdenum and tungsten enzymes.⁵⁷ The bis-W-MPT cofactor in BamBC contains a magnesium atom bridging the phosphate groups of the two pterin moieties. The methionine and serine residues binding Mg²⁺ in BamBC are conserved also in YdhV (Figures S1A and B), suggesting a similar

cofactor structure. Interestingly, we have obtained evidence that a halide ion (Cl^- or Br^-) may be present in the second metal coordination sphere in reduced YdhV. The presence of at least spurious chloride is difficult to exclude under the usual cell growth and protein expression/purification conditions. It should be noted, however, that the detected Mo(V) for YdhV displays no (large) hyperfine coupling to a chloride ion directly detectable in the EPR spectrum, at least in comparison to the coupling situation shown for Mo(V) in sulfite oxidase.⁵⁸⁻
⁶² Presumably, chloride may occupy a putative binding site close to the metal center only when it is in the Mo(IV) state and when a catalytically competent substrate is absent. Further studies are required to clarify a potential role of halides in YdhV.

EPR characterization of YdhV has provided distinct insights on paramagnetic molybdenum and iron species in comparison to other molybdenum enzymes. The signal of the Mo(V) species in the bis-Mo-MPT cofactor of YdhV resembles the high-pH signal of *E. coli* nitrate reductase NarGH⁶³ and the “high-*g*-unsplit” signal of DMSO reductase from *E. coli*, *R. sphaeroides*, or *R. capsulatus*,^{50-52, 64, 65} likely reflecting a similar Mo(V) coordination geometry in these enzymes. It should be noted that the coordination sphere of the aforementioned enzymes are thought to have a mixed sulfur/oxygen coordination geometry, with respect to binding of an oxo ligand and either a coordinated serine or aspartate residue. The *g*-matrix and hyperfine coupling parameters of the YdhV Mo(V) are reminiscent of the Mo(V) species of *R. sphaeroides* DMSO reductase generated in the presence of TMAO, albeit without observable ¹H hyperfine coupling. By comparison, the slightly elevated g_{av} value and altered *g*-tensor anisotropy in YdhV relative to DMSO reductase likely is due to coordination of the cysteine thiolate.⁵¹ A similarly rhombic ^{95,97}Mo hyperfine tensor likely reflects low coordination symmetry at the metal. As has been shown for the high- and low-pH Mo(V) EPR-signals of *E. coli* nitrate reductase, the dihedral angle between the oxo and aspartate ligands can modulate the Mo(V) *g*-values and the isotropic proton hyperfine interaction.⁶⁶ The

geometry of the two apical Mo(V) ligands may therefore also affect the EPR parameters of Mo(V) in YdhV.

Based on EPR spin Hamiltonian parameters for bis-MGD enzymes in literature, two tentative Mo(V) coordination structures are conceivable for YdhV (Figure 10C,D).

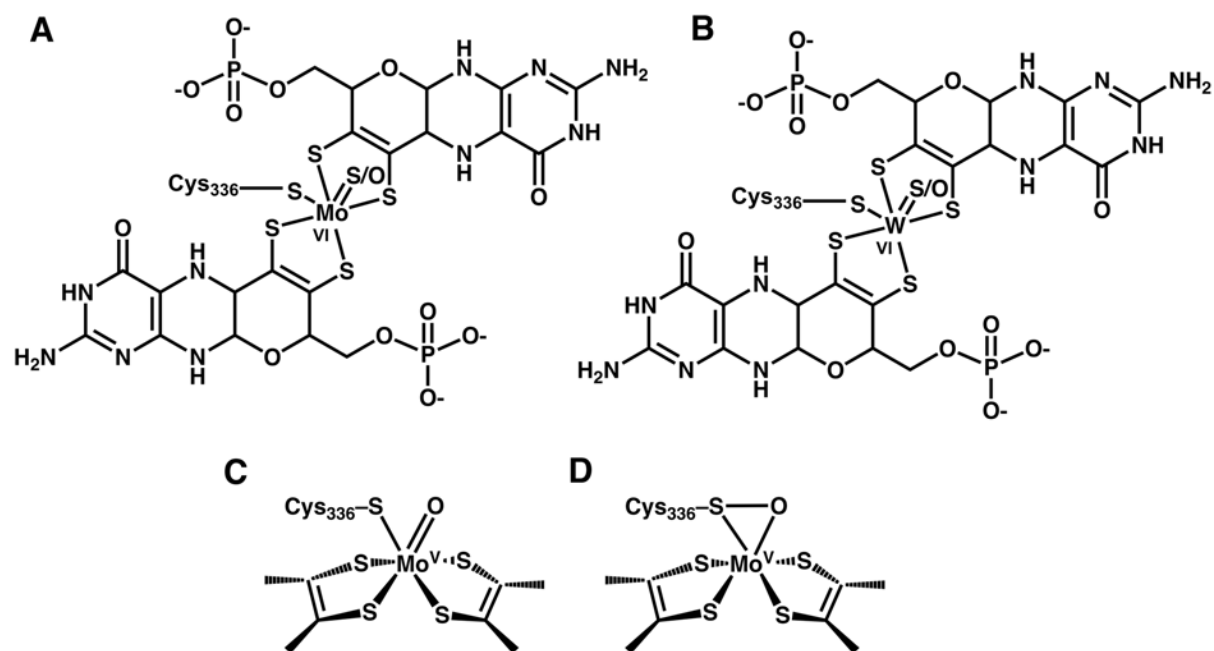


Figure 10. **Proposed bis-Mo-MPT and bis-W-MPT structures of YdhV.** Panels A and B depict (schematic) structures of bis-Mo/W-MPT cofactors (Mo^{VI}) suggested by our XAS and biochemical results while panels C and D depict possible cofactor structures in agreement with our EPR results (paramagnetic Mo^V; the EPR data reflect a mixed O/S coordination with respect to binding of the Cys₃₃₆ thiolate and an oxo ligand; the apparent lack of ¹H hyperfine coupling may occur for example via C) oxo ligand binding, or via D) sulfoxxygenated Cys₃₃₆ binding).

Given that YdhV coordinates a bis-Mo-MPT cofactor, a coordination geometry with four Mo–S_{MPT} bonds is most likely. The determined spin Hamiltonian parameters of the Mo(V) appear consistent with a mixed O/S coordination geometry with respect to binding of an oxo or a sulfido ligand, in addition to binding of the Cys₃₃₆ ligand, in agreement with the XAS data. Binding of the Cys thiolate to the bis-Mo-MPT is further supported by a relatively elevated g_{av} reminiscent to bis-MGD enzymes such as for formate dehydrogenases that coordinate a Cys thiolate ligand and a sulfido ligand.³⁶ Similarly, the arsenate reductase Mo(V) in complex with

arsenite shows a very similar Mo(V) *g*-tensor as YdhV, with a coordinated cysteine thiolate and coupling to a ⁷⁵As center.⁶⁷ Completing the coordination sphere, an oxo ligand is considered here, since the sulfido ligand was detected as a minor species. The apparent absence of substantial ¹H hyperfine coupling may be explained by a structure with an oxo ligand, as opposed to a hydroxo or water ligand. Alternatively, a sulfoxxygenated cysteine thiolate, as found, e.g., in [NiFeSe] hydrogenases⁶⁸ and NAD⁺-reducing [NiFe] hydrogenases,⁶⁹ might be possible. In conclusion, although the detected Mo(V) species represents a minor cofactor population, the properties of this Mo-MPT cofactor state in YdhV are comparable to Mo(V) species of bis-MGD cofactors in other enzymes.⁷⁰

Our apparent failure to identify any substrate reactions of YdhV may be explained by the lack of a functional [4Fe-4S] cluster in proximity to the bis-Mo-MPT cofactor. The amino acid sequence, in homology to BamBC and other enzymes, shows four cysteine residues that could bind a [4Fe-4S]-cluster in YdhV. However, even in anaerobically expressed and purified YdhV, the Fe-S cluster may be partially disassembled even by very minor residual O₂ contaminations. Characterization of the iron-sulfur cluster in YdhV by XAS and EPR revealed an apparent [2Fe-2S] species of low occupation, that likely has resulted from degradation of a [4Fe-4S] cluster. The apparently extreme sensitivity of the [4Fe-4S] cluster towards degradation may suggest that YdhV belongs to the group of enzymes that contain a so-called “hyper-active” cysteine residue according to a bioinformatics analysis.⁷¹ Such a cysteine may enhance [4Fe-4S]-cluster degradation also in YdhV.

In conclusion, our results reveal that YdhV belongs to a novel class of molybdenum oxidoreductases in *E. coli* that comprises a bis-Mo-MPT cofactor with a cysteine and a sulfido ligand at the metal (Figure 10) as well as likely a [4Fe-4S]-cluster, which is particularly sensitive to partial degradation. The enzyme binds preferentially molybdenum in the cofactor, but a similar cofactor with a tungsten center is formed when only tungsten is available (Figure

10). Accordingly, *E. coli* is able to assemble not only molybdenum-, but also tungsten-containing cofactors. The presence of a Mo/W containing bis-MPT cofactor in *E. coli* opens a new route of Moco biosynthesis and expands the canon of the versatile molybdenum and tungsten enzymes.

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Notes

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ABBREVIATIONS

bis-MPT guanine dinucleotide (bis-MGD)

cyclic pyranopterin monophosphate (cPMP)

cytosine dinucleotide (MCD)

electron paramagnetic resonance spectroscopy (EPR)

extended X-ray absorption fine structure (EXAFS)

iron-sulfur (Fe-S)

metal-binding pterin (MPT)

molybdenum cofactor (Moco)

trimethyl-N-oxide (TMAO)

X-ray absorption spectroscopy (XAS)

X-ray absorption near edge structure (XANES)

Supporting Information Available

TABLES

Table S1: **YdhV metal quantification and spin concentrations from EPR.**

Table S2: **Element contents in YdhV from TXRF.**

Table S3: **W EXAFS simulation parameters.**

Table S4: **Comparison of spin Hamiltonian parameters for selected molybdoenzymes.**

Table S5: **Fe EXAFS simulation parameters of YdhV.**

FIGURES

Figure S1: **Sequence alignment of YdhV and BamBC (A) and crystal structure of BamBC (B).**

Figure S2: **Elemental analysis with TXRF.**

Figure S3. **X-band EPR spectra of apo-YdhV.**

Figure S4. **EPR spectral simulation of the Mo(V) signal of ferricyanide-oxidized YdhV at 150 K.**

Figure S5: **Wide-sweep X-band EPR spectra of YdhV at low temperature.**

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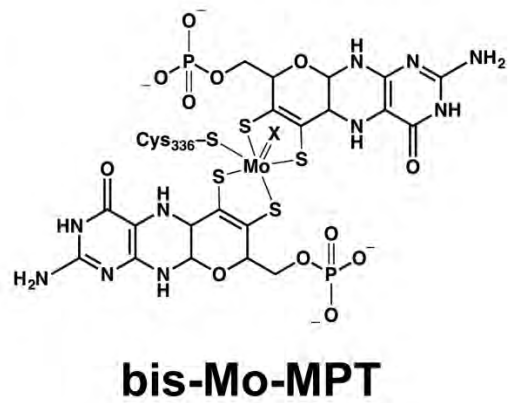
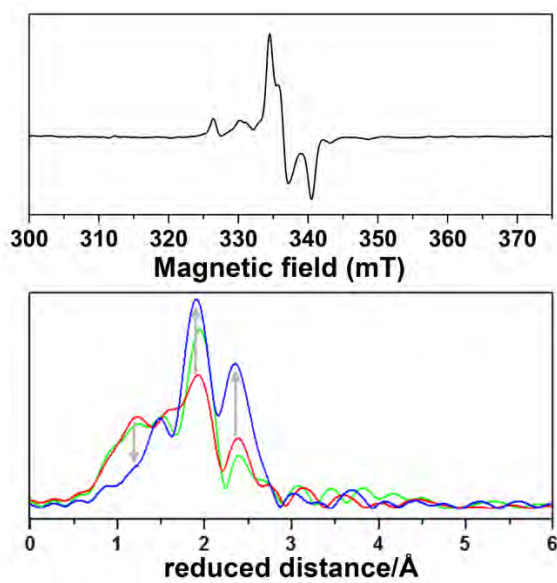
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*Identification of YdhV as first molybdoenzyme binding a
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Supporting Information

Identification of YdhV as first molybdoenzyme binding a bis-Mo-MPT cofactor in Escherichia coli

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Table S1: YdhV metal quantification and spin concentrations from EPR.

The concentrations of YdhV were determined by $A_{280\text{nm}}$ absorbance, while the concentrations of Mo, W, and Fe were determined by ICP-OES. The spin concentrations were determined through use of the Matlab toolbox ‘spincounting’ from the obtained spectra and associated tune picture of the resonator. The reported spin/YdhV relates the amount of spin to YdhV present. Quantitated spin, unless noted otherwise, detected from 80 K spectra measured with 4 mW microwave power, 5 G modulation amplitude, and 100 kHz frequency.

YdhV; Treatment	YdhV (μM)	Mo (μM)	W (μM)	Fe (μM)	Spin (μM)	Spin/YdhV
apo-YdhV (+MoO ₄ ²⁻) ^{1,2}	1094	1328	N/D	2100		
as-iso					104	0.095
dithionite-reduced					147	0.134
ferricyanide-oxidized					72	0.066
apo-YdhV (-MoO ₄ ²⁻) ¹	662	5	N/D	927		
as-isolated					2 ⁷	0.003
dithionite-reduced (12 K) ³					60	0.091
					57	0.086
ferricyanide-oxidized					4 ⁷	0.006
YdhV-Mo ^{2,5}	202	380	N/D	288		
as-isolated					5	0.040
dithionite-reduced					10	0.080
ferricyanide-oxidized (150 K) ⁴					17	0.084
					41	0.203
YdhV-W ^{5,6}	221	N/D	977	N/D		
as-isolated					1 ⁷	0.005
dithionite-reduced					28	0.126
ferricyanide-oxidized					4 ⁷	0.018

N/D not detected.

¹YdhV variant was purified and expressed aerobically. The bound Fe-S cluster was highly similar between aerobic and anaerobic expressions and purifications, respectively.

²YdhV was expressed in the presence of 1 mM MoO₄²⁻.

³Acquisition parameters at 8 μW microwave power, 10 G modulation amplitude, and 100 kHz modulation frequency.

⁴Acquisition parameters at 4 mW microwave power, 5 G modulation amplitude, and 100 kHz modulation frequency.

⁵YdhV variant with optimal loading of bis-Mo-MPT, as detected by EPR spectroscopy. YdhV was expressed in the $\Delta\text{mobA}/\Delta\text{mocA}$ deletion strain and was expressed and purified anaerobically.

⁶YdhV was expressed in the presence of 10 μM WO₄²⁻.

⁷It should be noted that the reported quantity of spin is at the limit of detection, therefore representing a negligible spin quantity.

Table S2: Element contents in YdhV from TXRF.^a

preparation	element per protein		
	P	Fe	Br
YdhV +Cl	1.7(5)	2.3(9)	0.1(1)
YdhV +Br	1.5(2)	2.9(5)	0.5(1)

^aData stem from two protein samples for each preparation and 3 repetitions per sample (standard deviation in parenthesis) and were determined relative to a gallium standard added to the protein solutions. Ni, Cu, and Zn contents were ≤ 0.4 metals per protein on average. From a series of YdhV samples to which increasing concentrations of a phosphorus standard were added (Supporting Figure S2), a P per protein ratio of 1.8(2) was determined.

Table S3: **W EXAFS simulation parameters.**^a

	W=O,-OH,-O	W=S	W-S _{MPT,cys}	W-Cl ^b ,W ^c	R _F
	N [per Mo] / R [Å] / 2σ ² x10 ³ [Å ²]				[%]
YdhV 10 μM W	0.5 [#] / 1.72 / 2 [*] 0.4 [#] / 1.81 / 2 [*]	0.5 [#] / 2.24 / 2 [*]	4.0 [*] / 2.40 / 5 0.6 [#] / 2.53 / 2 [*]	^b 0.1 / 2.79 / 2	19.2
YdhV 1 mM W	1.6 / 1.74 / 4 [§] 2.3 / 1.93 / 4 [§] 4.6 / 4.00 / 12 [§]	-	-	^c 4.4 / 3.45 / 12 ^{§*}	21.4
Na ₂ WO ₄ pH 4.0	2.0 / 1.74 / 8 [§] 1.9 / 1.91 / 8 [§] 6.5 / 3.57 / 2 [§]	-	-	^c 0.8 / 3.51 / 2	28.1
Na ₂ WO ₄ pH 7.1	3.0 / 1.77 / 5 [§] 1.0 / 2.10 / 5 [§]	-	-	-	23.0
Na ₂ WO ₄ pH 10.6	3.9 / 1.78 / 2	-	-	-	16.4

^aN, coordination number; R, interatomic distance; 2σ², Debye-Waller factor; R_F, fit error sum (deviation between Fourier back-transform in a 1.0-3.5 Å reduced distance window and simulation curve). Fit restraints: ^{*}fixed parameter, [#]N-values coupled to yield a sum of 2, ^{§,§}coupled to yield the same value. Parameters correspond to EXAFS simulations in Figure 6. ^bParameters correspond to a W-Cl distance, ^cparameters correspond to a W-W distance.

Table S4: Comparison of spin Hamiltonian parameters for selected molybdoenzymes. Table denotes signal type for the specified organism and enzyme, and associated g -tensor parameters reported, either by simulation or where applicable, by assignment. Where available, the associated $^{95,97}\text{Mo}$ hyperfine values, in addition to assigned Euler angles, are provided. Where applicable, associated nuclear quadrupole values and hyperfine values to nuclei other than Mo are excluded.

Organism, Enzyme	Signal Type	g -values	g_{av}	g_1-g_3	$(g_1-g_2)/(g_1-g_3)$	$^{95,97}\text{Mo}$ Hyperfine (Mo)	Euler Angles	Reference
<i>EcYdhV</i>	ferricyanide-oxidized	2.008, 1.996, 1.973	1.992	0.035	0.350	79, 149, 11	90, 104, 90	This work
<i>RcDMSOR</i>	high- g unsplit type 1	1.991, 1.983, 1.961	1.978	0.030	0.250	N/A	N/A	¹
	high- g unsplit type 2	1.987, 1.983, 1.963	1.978	0.024	0.160	N/A	N/A	¹
	high- g unsplit (glycol)	1.989, 1.983, 1.964	1.979	0.025	0.243	N/A	N/A	¹
<i>RsDMSOR</i>	high- g unsplit (glycerol)	1.99, 1.98, 1.96	1.98	0.030	0.333	N/A	N/A	²
	high- g split (TMAO)	1.999, 1.989, 1.972	1.987	0.027	0.370	40, 112, 156	N/A	³
<i>EcDMSOR</i>	poised at $E = -51$ mV	1.987, 1.976, 1.960	1.974	0.027	0.407	N/A	N/A	⁴⁻⁶
<i>EcNarGH</i>	high pH	1.987, 1.980, 1.960	1.976	0.027	0.258	97, 126, 98	N/A	⁷⁻⁹
	low pH	2.001, 1.985, 1.964	1.983	0.037	0.437	N/A	N/A	⁷⁻⁹
<i>EcFDH(Se)-H</i>	$g = 2.094$ signal	2.094, 2.001, 1.990	2.029	0.104	0.894	118, 71, 71	N/A	¹⁰
<i>CnFFDH</i>	dithionite-reduced	2.009, 2.001, 1.992	2.001	0.017	0.471	138, 82, 45	11, 0, 153	¹¹
<i>WsPsrABC</i>	very-high g (polysulfide)	2.017, 2.003, 1.987	2.002	0.029	0.481	56,56, 140	N/A	¹²
<i>S.sp. ArrAB</i>	turnover-detected	1.990, 1.980, 1.957	1.976	0.033	0.303	122, 68, 143	N/A	¹³
Chicken SO	high pH	1.990, 1.966, 1.954	1.970	0.036	0.666	164, 63, 34	0, 14, 22	¹⁴
<i>PpSoxCD1</i>	high pH	1.985, 1.961, 1.950	1.965	0.035	0.686	162, 64, 30	0, 21, 41	¹⁵
Bovine XO	very rapid	2.025, 1.955, 1.949	1.977	0.076	0.921	133, 55, 57	8, 36, 0	¹⁶

	rapid type 1 formamide	1.990, 1.971, 1.967	1.976	0.023	0.826	184, 74, 77	0, 18, 0	¹⁶
	desulfo	1.971, 1.966, 1.954	1.963	0.016	0.3110	196, 79, 81	81, 79, 196	¹⁷
	formaldehyde -inhibited	1.991, 1.977, 1.951	1.973	0.040	0.349	159, 69, 156	178, 69, 67	^{18, 19}

N/A not applicable

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Table S5: Fe EXAFS simulation parameters of YdhV.^a

Fe-O,C	Fe-S	Fe-Fe
N [per Fe] / R [Å]		
1.2 / 2.50	2.1 / 2.23	1.6 / 2.76
3.5 / 2.92	1.5 / 2.38	

^aParameters correspond to the EXAFS spectrum in Fig. 9 (N, coordination number; R, interatomic distance). The fit error sum, R_F , for reduced distances of 1-3 Å was 10.9 %. A Debye-Waller factor ($2\sigma^2$) of 0.002 \AA^2 was employed for all shells except the Fe-Fe distance ($2\sigma^2 = 0.005 \text{ \AA}^2$) in the fitting.

YdhV 1 MANGW T G N I L R V N L T T G N I T L -- E D S S K F K S F V G G M G F G Y K I M Y D E V P P G T K P F D E A N K L
BamB 1 M R Y A E T G Y V L E V D L T K G S I E R V A T D P R D T E L Y L G G L G T N A K I L W D R V P P E V E F F S P E N L L

YdhV 59 V F A T G P L T G S G A P C S S R V N I T S L S T F T K G N L V V D A H M G G F F A A Q M K F A G Y D V I I I E G K A K
BamB 61 I F A A G L L C G T P A T G C N R T I V S T V S P -- Q T K L M A F S M M G G F W A P E L K Y A G Y D K I I F R G K S P

YdhV 119 S P V W L N I K D D K V S L E K A D F L W G K G T R A T T E E T C - R L T S P E T C V A A I G O A G E N L V P L S G M L
BamB 119 E L V Y L Y I N N D K V E I R D A S H L K G K G A I E T A E I I K K E L N E P R A Q V A A I G K A G E N R V F Y A S I E

YdhV 178 N S R N H S G G A C T G A I M G S K N L K A I A I E C T K G V N I A D R O E M K S L N D Y M M T E L I G A N N H V V P
BamB 179 Q G R S S A S R G G I G A V M G D K G L K A V V V R G T K D L C V A K P E E Y I G L C N E V L D Y I K H R E E N - P I P

YdhV 238 S T P Q S W A E Y S ----- D P K S R W T A R K G L F W G A A E G G P I E T G E I P P G N O N T V G F R T Y K S V F
BamB 238 D V M P I L A G L G S P Q E M K V H D E K W H T - E N F N W G N A R T R R K ----- D F W T D E V S H

YdhV 292 D L G P A A E K Y T V K M S G C H S C P I F C M T O M N I P R V K E F G V P G T G G N T C V A N F V H T T I F P N G P K
BamB 284 A W E K T M D K A R T R L I S C Y N C P M K C G A T I S M E G L P T Y M M K ----- C F T K L T Y T ----- M A

YdhV 352 D F E D K D D G R V I G N L V G L N L F D D Y G L W C N Y G Q L H R D F T Y C - Y S K G V F K R V L P A E E Y A E I R W
BamB 332 A Y S D L D F G L R I A O K ----- A T E Y G L D G F S A P Q V M A F A F E L L E K G I L K ----- D ----- S D F

YdhV 411 D O L E A G D V N F I K D F Y Y R L A H C V --- G E L S H L A D G S Y A I A E R W N L G E E Y W G Y A K N K L W S P F
BamB 378 P G L P E C N --- E E R F F Y L L D K I V N R D G I G D I L A N G T Y W A A Q E I G N G A E D -- Y A H N N I K K H E

YdhV 468 G Y P V H H A N E A S A Q V G S I V N C M F N R D C M T H T H I N F I G S G L P L K L O R E V A -----
BamB 433 Q L P L K L S M -- L N P I Y Y L M Y C T G E K I N I T Q I E G Q F P Q A P Y P K L E Q R E A F V E D W I Q V P D E K F

YdhV 516 K E L F G S - E D A Y D E T K N Y T P I N D A K I K Y A K W S L L R V C L H N A V T I C N W V W P M T V S E - L K S R N
BamB 491 K K I F L E W E P R G E K S M P N F P T V D M C C D I V D W Q E M M H Y I D D A L G C C A G L S S F P L K P P Y H I H -

YdhV 574 Y R G D L A L E A K F F K A I T G E D M T O E K L D L A A E R I F T L H R A Y T V K L M O T K D M R N E H D L I C S W V
BamB 550 ----- N Y P K F I A A G A G I E M D T E K L K K A A K R Y R T L V R A F N I R R G M ----- R R -----

YdhV 634 F D K D F Q I P V F T E G T D K M D R D D M H A S L T M F Y K E M G W D P O L G C P T R E T L O R L G L E D I A A D L A
BamB 591 V D E Q P P A N H W K N R F P E L ---- E K E L L D S Y Y K L K G W N D - D G I P T K E T L D D L G L G Y V G D E F I

YdhV 694 A H N L L P A -
BamB 646 K R G I L S A G

Figure S1A: Sequence alignment of YdhV and BamBC.

Alignment was carried out with Clustal (1.2.4, <http://www.uniprot.org/align/>) using amino acid sequences for YdhV from *E. coli* (AKK48365.1) and benzoyl-CoA reductase BamBC (PDB ID 4Z3Y) from *Geobacter metallireducens*. Colors denote: magenta = Cys322 (C) ligand of W in BamBC, green = Thr close to W in BamBC, red = cysteine (C) ligands to the [4Fe-4S] cluster next to the W-cofactor in BamBC, orange = amino acid ligands at the magnesium ion bridging the two MPT units of the W-cofactor in BamBC.

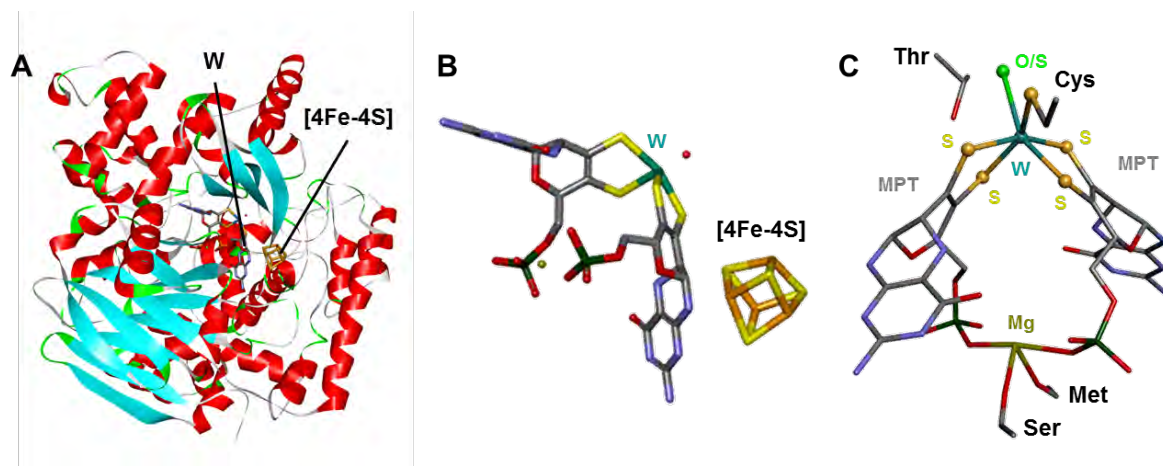


Figure S1B: Crystal structure of BamBC.

(A) Structure (truncated) according to PDB entry 4Z3Y (2.36 Å resolution) with positions of W-cofactor and [4Fe-4S] cluster marked. (B) W-bis-MPT-Mg cofactor with neighboring [4Fe-4S] cluster at ~ 9 Å distance. (C) W-cofactor with bound or neighboring amino acids marked (an O atom at W was assigned in the crystal structure).

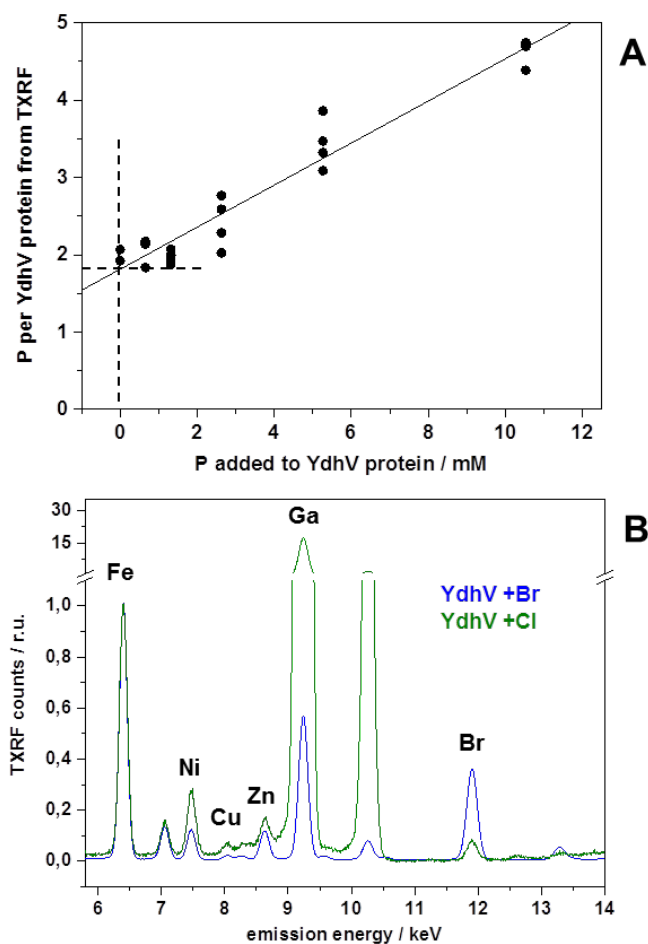


Figure S2: Elemental analysis with TXRF. (A) Phosphorus determination in YdhV protein. To YdhV samples (1.0 ± 0.1 mM protein), a phosphorus standard (Na_3PO_4) was added at increasing concentrations (in addition to a gallium concentration standard), the P contents were determined from TXRF spectra (P $\text{K}\alpha$ emission at ~ 2.0 keV, not shown), and P per protein ratios were determined. A linear fit (i.e. extrapolation to zero added P) provided a P per protein content in YdhV of 1.8 ± 2 . (B) Quantification of bromine in YdhV samples (green line, YdhV prepared with Cl⁻; blue line, YdhV prepared with Br⁻; spectra are normalized at the Fe $\text{K}\alpha$ emission). $\text{K}\alpha$ emission lines of selected elements are marked (unmarked features = $\text{K}\beta$ emission lines). A Ga standard was added to the samples at different concentrations. Element per protein ratios from the TXRF analysis are summarized in Table S2.

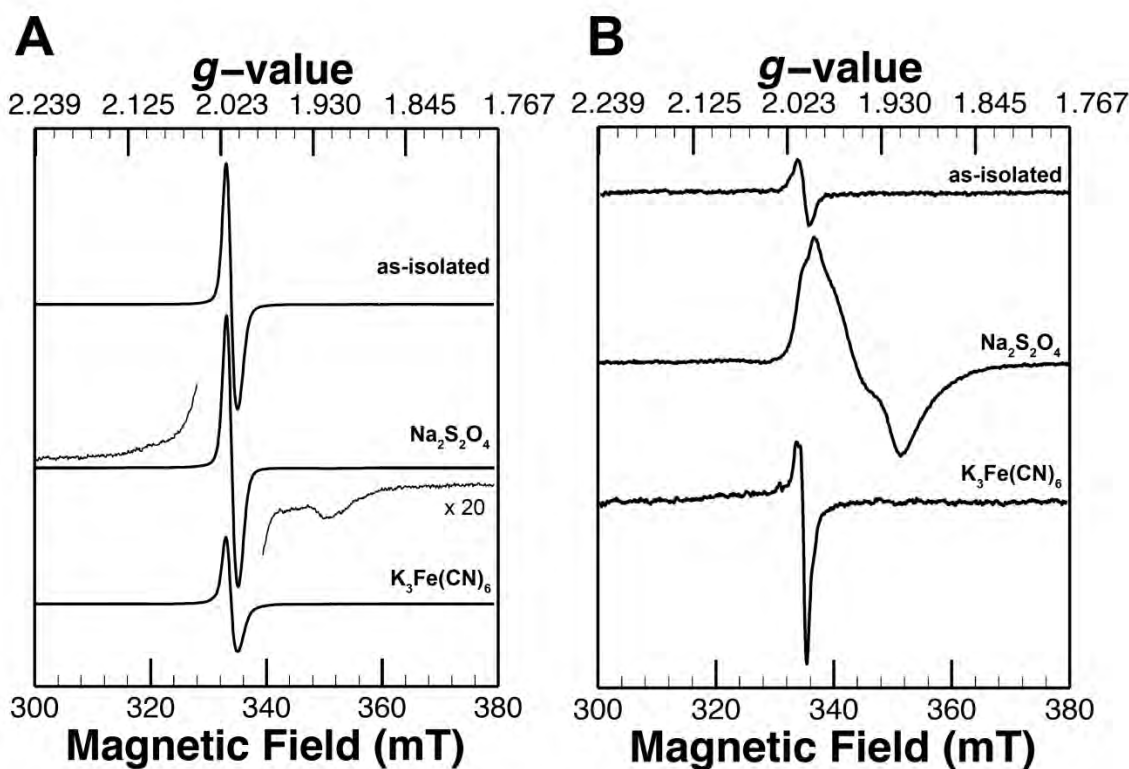


Figure S3: X-band EPR spectra of apo-YdhV. (A) apo-YdhV aerobically expressed in the presence of 1 mM Na_2MoO_4 , (B) apo-YdhV aerobically expressed without Na_2MoO_4 . For each panel, the top spectrum represents the as-isolated protein, the middle spectrum represents sodium dithionite-reduced protein, and the bottom spectrum represents $\text{K}_3\text{Fe}(\text{CN})_6$ -oxidized protein. Features present in the baseline are enlarged as noted (thin lines). Spectra were recorded at 80 K at 4 mW microwave power, 5 G modulation amplitude and 100 kHz modulation frequency. Spectra have been scaled to represent a YdhV concentration of 1094 μM . For associated concentrations of Mo and Fe, see Table S1.

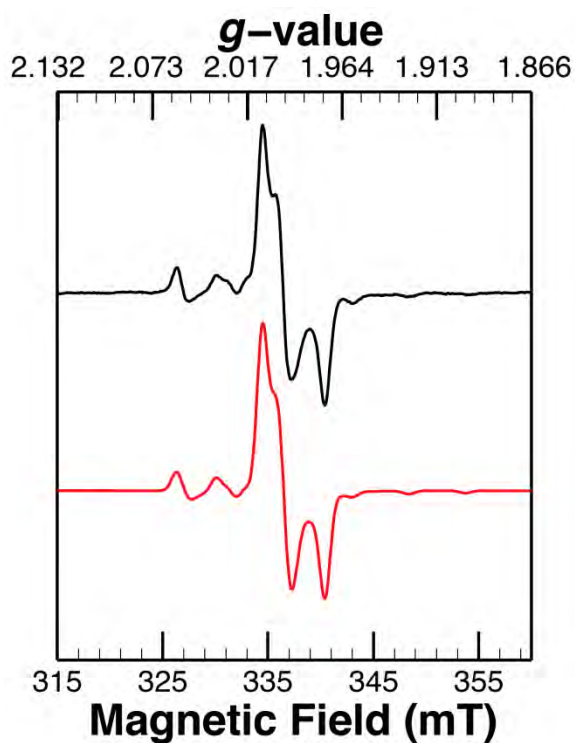


Figure S4: EPR spectral simulation of the Mo(V) signal of ferricyanide-oxidized YdhV at 150 K.

The experimental spectrum (black line) represents ferricyanide-oxidized YdhV as depicted in Figure 7 (panel A, bottom black spectrum), measured at 150 K at 4 mW microwave power, 5 G modulation amplitude and 100 kHz modulation frequency. The spectral simulation (red line) includes two components whereby the Mo isotopes that possess a nuclear spin were treated separately using the simulation toolbox EasySpin. Simulation parameters: g : 2.008, 1.996, 1.973 ($g_{av} = 1.992$, anisotropy = 0.0354, rhombicity = 0.3503), g -strain: 6.7, 8.2, 6.7 ($\times 10^{-3}$), A: ($^{95,97}\text{Mo}$) [78.8, 148.7, 11.4] MHz, Euler angles: 89.8, 103.5, 89.9 (degrees).

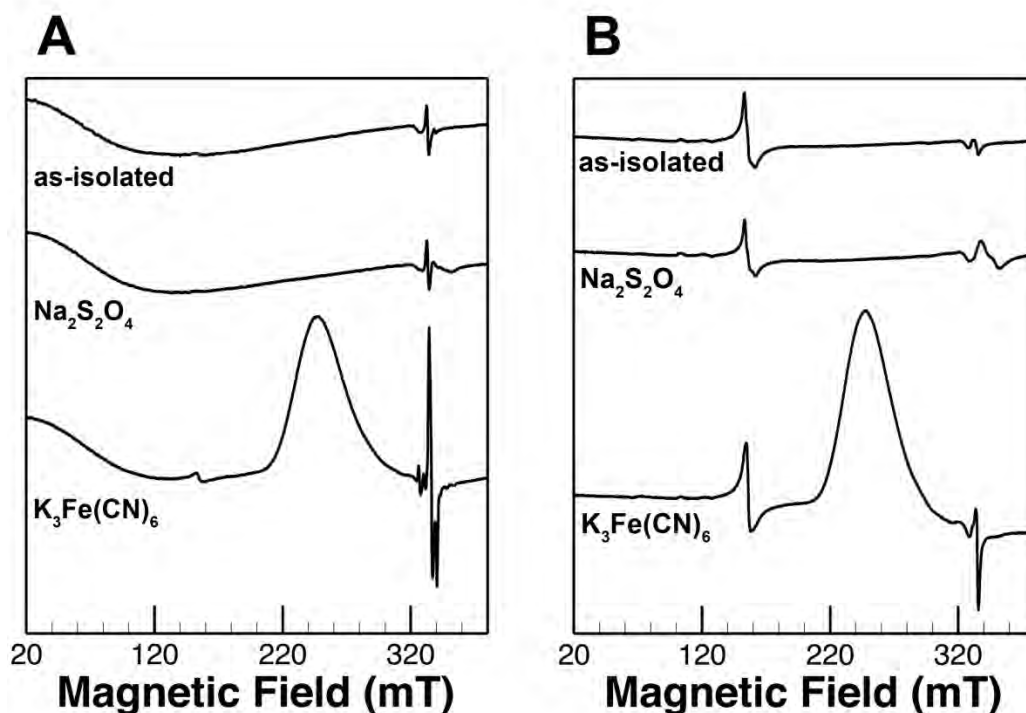


Figure S5: Wide-sweep X-band EPR spectra of YdhV at low temperature.

(A) bis-Mo-MPT-loaded YdhV, (B) apo-YdhV aerobically expressed without Na_2MoO_4 . Spectra refer to YdhV as-isolated (top), after treatment with 10 mM sodium dithionite (middle), and after treatment with $\text{K}_3\text{Fe}(\text{CN})_6$ (bottom), and correspond to the spectra of samples in Figure 1 (panel A) and Figure S1 (panel B). Spectra were recorded at 12 K at 4 mW microwave power, 5 G modulation amplitude and 100 kHz modulation frequency. Spectra in panel A represent a YdhV concentration of 202 μM , while spectra in panel B represent a YdhV concentration of 662 μM . The large feature at 250 mT in panel A represents the signal of excess paramagnetic ferricyanide in the samples that is readily observed at 12 K. For associated concentrations of Mo and Fe, see Table S1.