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Title: Dinuclear manganese complexes for water oxidation: evaluation of electronic effects and catalytic activity

A library of dinuclear Mn$_{2}$$^{III}$ complexes for water oxidation has been synthesized and studied. Using the generated library, it was shown that incorporation of a distal group would promote water oxidation catalysis. The improved catalytic activity was suggested to stem from pre-orientation of the incoming water nucleophile, facilitating O–O bond formation, in connection with hydrogen bonding interactions between the distal group and high-valent Mn species. These results illustrate the importance of mastering both proton and electron transfer processes during water oxidation, and could be a general strategy for designing more efficient catalysts.
Dinuclear manganese complexes for water oxidation: evaluation of electronic effects and catalytic activity

Wael A. A. Arafah, Markus D. Kärkäs, Bao-Lin Lee, Torbjörn Åkermark, Rong-Zhen Liao, Hans-Martin Berends, Johannes Messinger, Per E. M. Siegbahn and Björn Åkermark*

During recent years significant progress has been made towards the realization of a sustainable and carbon-neutral energy economy. One promising approach is photochemical splitting of H2O into O2 and solar fuels, such as H2. However, the bottleneck in such artificial photosynthetic schemes is the H2O oxidation half reaction where more efficient catalysts are required that lower the kinetic barrier for this process. In particular catalysts based on earth-abundant metals are highly attractive compared to catalysts comprised of noble metals. We have now synthesized a library of dinuclear MnIIII catalysts for H2O oxidation and studied how the incorporation of different substituents affected the electronics and catalytic efficiency. It was found that the incorporation of a distal carboxyl group into the ligand scaffold resulted in a catalyst with increased catalytic activity, most likely because of the fact that the distal group is able to promote proton-coupled electron transfer (PCET) from the high-valent Mn species, thus facilitating O–O bond formation.

1. Introduction

The world's global energy consumption is primarily based on fossil fuels. However, the rapidly increasing global energy demand coupled to diminishing fossil fuel supplies call for the development of sustainable and carbon-neutral alternatives, with feedstock(s) that are environmentally friendly and abundant. There are already efficient solar cells for generation of electricity but in order for this to become truly useful, it is necessary that part of the energy can be stored. One attractive alternative would therefore be to store the energy in sunlight by the splitting of H2O into O2, protons and electrons (eqn (1)). The generated electrons can then be used to produce H2 or other solar fuels, thus allowing the energy to be stored in the form of chemical bonds. The production of solar fuels constitutes an attractive solution for a future sustainable energy economy.1–5

Nature has since a long time figured out how the process of splitting of H2O can be realized and the natural photosynthetic machinery involves the synchronization of several complicated features, such as light-harvesting, charge separation and electron transfer. Both in natural and artificial photosynthesis, catalytic H2O oxidation is an essential part of the intricate photochemical process of solar to chemical energy conversion. Although deceptively simple, H2O oxidation (eqn (1)) is associated with a high thermodynamic potential (eqn (2)), a high kinetic barrier and requires a multitude of bond breaking and bond making events. This makes the H2O oxidation half reaction the pitfall for the construction of artificial devices for the production of solar fuels.6–8

\[ 2\text{H}_2\text{O} \rightarrow \text{O}_2 + 4\text{H}^+ + 4\text{e}^- \]  

\[ E = 1.229 - (0.059\ \text{pH}) \text{ V vs. NHE} \]

In the natural system, H2O oxidation is mediated by the oxygen-evolving complex (OEC) comprised of a Mn4Ca oxo-bridged cluster.9–12 This has been a source of inspiration for scientists and has resulted in a plethora of structural and functional model systems of the OEC.13 Although substantial progress has been made during the latest years in producing highly active and rugged water oxidation catalysts (WOCs), the essential question still remains: how to construct WOCs that are capable of operating under the harsh reaction conditions required to oxidize H2O and how to reproduce the high activity and stability displayed by the natural system. The majority of the developed artificial molecular WOCs are based on the scarce transition metals Ru14–27 and Ir28–32. However, for large scale applications it is most likely necessary that the WOCs are
based on earth-abundant elements. This prompts the examination and development of molecular WOCs based on low-cost first-row transition metals. Although there have been a couple of reports on WOCs comprised of Mn, Co, Fe, and Cu, this research has not been as successful as with the rare metals Ru and Ir.

The incorporation of negatively charged groups into ligand frameworks has been shown to drastically reduce the redox potentials of the corresponding metal complexes. The electron-rich ligand environment helps to stabilize the metal center(s) in high oxidation states. We have previously reported the synthesis of Mn complexes housing ligands with benzylic amines. However, due to the labile benzylic positions, these complexes failed to mediate H₂O oxidation.

Our group recently reported on the synthesis and development of the bio-inspired ligand containing carboxylate, phenol and imidazole functionalities, which are all important elements in the natural system. Complexation of ligand 1 with Mn(OAc)₂ yielded the dinuclear Mn₂II,III complex 2, which in the solid state crystallized as the tetranuclear complex 2' with loss of the bridging acetate ligand, thus resembling the Mn₄Ca cluster in the OEC (Fig. 1). Maintaining a high redox flexibility of the active metal center(s) in the artificial WOCs constitutes a key parameter when designing synthetic WOCs, since the oxidation of H₂O requires the collective transfer of four electrons. It was therefore vital that the negatively charged functional groups in the dinuclear Mn₂II,III complex were shown to dramatically lower the redox potentials of the complex. By replacing the sensitive benzylic groups by imidazoles, along with the introduction of carboxylate moieties, it became possible for the dinuclear Mn₂II,III complex 2 to promote catalytic H₂O oxidation by use of the mild one-electron oxidant [Ru(bpy)₃]³⁺ (bpy = 2,2'-bipyridine). In addition to being capable of mediating chemical H₂O oxidation, photochemical H₂O oxidation was also realized when using [Ru(bpy)₃]²⁺-type complexes as photosensitizers. This has similarities to the natural photosynthetic system where light is employed to induce charge separation with subsequent H₂O oxidation.

Compared to heterogeneous WOCs, homogeneous molecular catalysts offer great advantages both in terms of kinetic studies and easy steric and electronic tuning of the catalysts. Thanks to the straightforward synthetic protocol associated with the catalysts 1 and complex 2, it was decided to study how the introduction of different functional groups into the ligand framework of 1 affected the catalytic activity and electronic properties of the corresponding dinuclear Mn complexes.

Here we report the synthesis and characterization of eight ligand scaffolds for dinuclear metal complexes, together with their corresponding dinuclear Mn₂II,III counterparts. The developed Mn complexes contained a wide variety of different substituents and their electronic and catalytic features were systematically examined. Moreover, this study has established the impact of having a non-innocent distal group in these complexes to promote proton transfer processes during H₂O oxidation catalysis. Within the library of the developed dinuclear Mn₂II,III complexes, complex 6 possessing the distal carboxyl group was found to evolve O₂ more efficiently than the other Mn complexes.

This effect could be due to both pre-orientation of the incoming H₂O nucleophile and to hydrogen bonding to the aqua and hydroxide coordinated to the Mn center(s). Hydrogen bonding could facilitate proton-coupled electron transfer (PCET) and pre-orientation of H₂O could facilitate nucleophilic attack on a high-valent Mn species, similar to the previously reported Co "Hangman" complexes. The incorporation of non-innocent distal groups into metal-based WOCs could thus be a general strategy for promoting catalytic H₂O oxidation activity. In addition, the constructed dinuclear Mn₂II,III WOCs have the advantage of being comprised of earth-abundant elements, which renders them especially attractive compared to noble metal-based WOCs.

2. Experimental section

2.1. Materials and methods

[Ru(bpy)₃]²⁺, 5-((1,3-dioxoisindolin-2-yl)methyl)-2-hydroxyisophthalaldehyde and 4'-methyl-2,2'-bipyridine-4-carboxylic acid were prepared according to a previously reported procedure. All other reagents including solvents were obtained from...
commercial suppliers and used directly without further purification. All solvents were dried by standard methods when needed. 

2.2. Electrochemistry

Electrochemical measurements were carried out using an Autolab potentiostat with a GPEs electrochemical interface (Eco Chemie), using a glassy carbon disk (diameter 3 mm) as the working electrode, and a platinum spiral as a counter-electrode. The reference electrode was an Ag/AgCl electrode (3 M KCl aqueous solution) and the electrolyte was a 0.1 M phosphate buffer (pH 7.2). All potentials were reported vs. NHE, using the [Ru(bpy)3]3+/[Ru(bpy)3]2+ couple as a reference.

2.3. Oxygen evolution measurements

Oxygen evolution was measured by mass spectrometry (MS), containing K2PO4 (3 mM). The catalyst solutions used in the experiments were then made by diluting the stock solutions with phosphate buffer (0.1 M, pH 7.2) to the desired concentrations. The solutions were then deoxygenated by bubbling with N2 for at least 15 min before being used in the experiments.

2.4. Chemical H2O oxidation with [Ru(bpy)]3+

In a typical run [Ru(bpy)]3+/PF6 (3.4 mg, 3.4 μmol) was placed in the reaction chamber and the reaction chamber was evacuated using a rough pump. ~ 35 mbar He was then introduced into the system. After a couple of minutes the catalyst solution (0.50 mL, 4.0 μmol) was injected into the reaction chamber. The generated oxygen gas was then measured and recorded versus time by MS.

2.5. Computational details

The geometry optimizations were performed at the B3LYP level of density functional theory as implemented in the Gaussian 09 package. The SDD pseudopotential was used to describe Mn, and the 6-31G(d,p) basis set was used for the C, N, O, H elements. Analytical frequency calculations were carried out at the same level of theory as the geometry optimization to obtain the Gibbs free energy corrections and to confirm the characteristics of the optimized structures. On the basis of these optimized geometries, single-point calculations were performed employing a larger basis set, where all elements, except Mn, were described by 6-311+G(2df,2p) at the B3LYP (15% exact exchange) level. Solution effects from the water solvent were calculated using the SMD continuum solvation model with the larger basis set at the B3LYP level. The B3LYP-D2 energies are reported, including dispersion corrections proposed by Grimme and Gibbs free energy corrections from B3LYP.

2.6. EPR measurements

X-band (9 GHz) electron paramagnetic resonance (ESR) measurements were performed at temperatures between 5 and 30 K with a Bruker E500 ELEXYS spectrometer using a 4122SHQE or a 4116DM resonator. For cooling with liquid helium an Oxford Instruments ESR 900 cryostat was installed. Frozen solutions of the complexes were prepared in 0.2 M phosphate buffer (pH 7) and for solid state spectra the complex and potassium bromide were ground together. All samples were frozen in liquid nitrogen prior to EPR measurements. The modulation amplitude was 5 G and the microwave power was varied between 200 μW and 6 mW.

2.7. Synthesis

Synthesis of 3,5-diformyl-4-hydroxybenzoic acid (4a). 4-Hydroxybenzoic acid (3a, 0.550 g, 4.00 mmol) and hexamethylenetetramine (4.50 g, 32.0 mmol) were dissolved in anhydrous CF3COOH (10 mL) and the reaction mixture was heated at 110 °C for 48 h. The reaction was cooled to room temperature before 4 M HCl (25 mL) was added to the reaction mixture. The precipitate was filtered, washed with H2O (3 × 10 mL) to afford the title compound (0.710 g, 86.2%). 1H NMR (400 MHz, [D6]acetone): δ = 12.21 ppm (br, 1H), 10.36 (s, 1H), 8.69 (s, 1H); 13C NMR (100 MHz, [D6]acetone): δ = 188.8, 160.0, 149.3, 122.0, 115.8, 113.1; HRMS (ESI) calcd for C8H1O3 [M+H]+: 193.0142; found: 193.0145.

Synthesis of 2,2′-(5-carboxy-2-hydroxy-1,3-phenylene)bis(1H-benzo[d][1,2,4]triazole-4-carboxylic acid) (5a). Compound 4a (1.10 g, 5.50 mmol) and 2-amino-3-nitrobenzoic acid (2.00 g, 11.0 mmol) were suspended in ETOH (50 mL). The suspension was heated until complete dissolution and then cooled to room temperature. A freshly prepared aqueous solution of Na2S2O4 (5.70 g, 32.8 mmol, 25 mL) was added and the reaction mixture was heated at 70 °C for 5 h. After the mixture had been allowed to cool to room temperature, a yellow precipitate was formed, which was filtered, washed with H2O (4 × 10 mL), acetone (4 × 10 mL) to afford the product.
(2.10 g, 83.9%). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 14.01 (br, 3H), 8.91 (s, 2H), 7.88 (d, $J$ = 7.91 Hz, 2H), 7.76 (d, $J$ = 7.47 Hz, 2H), 7.28 (t, $J$ = 7.93 Hz, 2H); $^{13}$C NMR (100 MHz, [D$_6$]DMSO): $\delta$ = 173.3, 167.7, 167.1, 154.7, 132.3, 123.0, 121.9, 121.0, 116.9, 114.3, 111.9; HRMS (ESI) calcd for C$_{23}$H$_{13}$N$_4$O$_7$ [M–H$^+$]/C$_0$: 457.0779; found: 457.0795.

Synthesis of dinuclear Mn$_{II,III}$ complex 6a. Mn(OAc)$_2$·4H$_2$O (0.153 g, 0.625 mmol) and NaOAc (0.209 g, 25.0 mmol) were added to a suspension of ligand 5a (0.114 g, 0.250 mmol) in MeOH (4 mL). The mixture was heated at 70 $^\circ$C for 20 h and then centrifuged and washed with MeOH (3 × 5 mL), and dried under vacuum to afford the complex (0.140 g, 76.6%). HRMS (ESI) calcd for C$_{26}$H$_{17}$Mn$_2$N$_4$NaO$_{10}$ [M+H$^+$]: 677.9598; found: 677.9593; anal. calcd for C$_{26}$H$_{22}$Mn$_2$N$_4$NaO$_{13}$ [6a·3H$_2$O]: C 42.70, H 3.03, N 7.66, Mn 15.02%; found: C 42.54, H 2.93, N 7.58, Mn 15.08%; IR (KBr): $\nu$ = 3428, 2926, 1563, 1497, 1474, 1403, 1385, 1320, 1260, 1020, 765, 676, 655, 619 cm$^{-1}$.

Synthesis of 5-bromo-2-hydroxyisophthalaldehyde (4b). 4-Bromo-phenol (3b, 1.47 g, 8.49 mmol) and hexamethylenetetramine (9.52 g, 67.9 mmol) were dissolved in anhydrous CF$_3$COOH (25 mL) and the yellow solution was heated at 110 $^\circ$C for 48 h. After cooling to room temperature, the mixture was added to aqueous HCl (4 M, 50 mL) and stirred for 3 h. The yellow precipitate was collected by filtration, washed with H$_2$O (35 mL) and dried under vacuum to yield the title compound as a yellow powder (1.67 g, 92.4%). $^1$H NMR (400 MHz, [D$_6$]acetone): $\delta$ = 11.71 (s, 1H), 10.28 (s, 2H), 8.19 (s, 2H); $^{13}$C NMR (100 MHz, [D$_6$]acetone): $\delta$ = 192.5, 140.5, 126.0, 112.5; HRMS (ESI) calcd for C$_8$H$_5$BrNaO$_3$ [M+Na$^+$]: 250.9314; found: 250.9310.

Synthesis of 2,2′-(5-bromo-2-hydroxy-1,3-phenylene)bis[1H-benzo[d]imidazole-4-carboxylic acid] (5b). An aqueous solution of Na$_2$S$_2$O$_4$ (6.09 g, 35.0 mmol, 6 mL) was added to an ethanolic solution containing 5-bromo-2-hydroxyisophthalaldehyde 4b (1.15 g, 5.00 mmol) and 2-amino-3-nitrobenzoic acid (1.82 g, 10.0 mmol). The resulting mixture was heated at 70 $^\circ$C for 5 h and the resulting yellow precipitate was filtered and washed with H$_2$O, acetone and finally dried under vacuum to give the product (2.00 g, 80.9%). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 13.64 (br, 2H), 8.58 (s, 2H), 7.97 (d, $J$ = 7.56 Hz, 2H), 7.85 (d, $J$ = 7.56 Hz, 2H), 7.33 (t, $J$ = 7.56 Hz, 2H); $^{13}$C NMR (100 MHz, [D$_6$]DMSO): $\delta$ = 166.8, 151.5, 141.1, 135.3, 133.9, 132.1, 124.6, 122.6, 122.1, 117.5, 115.4, 106.7; HRMS (ESI) calcd for C$_{22}$H$_{13}$BrN$_4$NaO$_5$ [M+Na$^+$]: 514.9962; found: 514.9967.

Synthesis of dinuclear Mn$_{II,III}$ complex 6b. Ligand 5b (0.100 g, 0.203 mmol) was added to a solution of NaOAc (0.166 g, 2.03 mmol) and Mn(OAc)$_2$·4H$_2$O (0.124 g, 0.506 mmol) in MeOH (3 mL). The resulting mixture was then refluxed for 20 h. The precipitate formed was isolated by centrifugation, washed with MeOH (5 × 5 mL) and dried under vacuum to give the desired product (0.120 g, 79.4%). HRMS (ESI) calcd for C$_{25}$H$_{16}$BrMn$_2$N$_4$NaO$_8$ [M+Na$^+$]: 711.8805; found: 711.8810; anal. calcd for C$_{25}$H$_{22}$Mn$_2$N$_4$O$_{11}$ [6b·3H$_2$O]: C 40.35, H 2.98, N 7.53, Mn 14.76%; found: C 40.43, H 3.05, N 7.49, Mn 14.71%; IR (KBr): $\nu$ = 3435, 2926, 1631, 1555, 1493, 1470, 1400, 1385, 1319, 1247, 1003, 765, 619 cm$^{-1}$.

Synthesis of 4-hydroxybiphenyl-3,5-dicarbaldehyde (4c). Biphenyl-4-ol (3c, 1.36 g, 8.00 mmol) and hexamethylenetetramine (22.4 g, 160 mmol) were placed in a round-bottomed flask which was purged with nitrogen. After addition of trifluoroacetic acid (48 mL), the mixture was refluxed at 110 $^\circ$C for 72 h. The reaction mixture was combined with 4 M HCl (240 mL) in a 500 mL round-bottomed flask and stirred for 3 h. The yellow precipitate was collected by filtration, washed with H$_2$O (35 mL) and dried under vacuum to yield the title compound as a yellow powder (1.67 g, 92.4%). $^1$H NMR...
(500 MHz, CDCl₃): δ = 11.62 (s, 1H), 10.32 (s, 2H), 8.20 (s, 2H), 7.61–7.57 (m, 2H), 7.51–7.45 (m, 2H), 7.43–7.37 (m, 1H); 13C NMR (100 MHz, [D₆]DMSO): δ = 192.2, 163.0, 138.2, 136.0, 133.7, 130.7, 129.3, 128.2, 126.8, 123.5; HRMS (ESI) calcd for C₁₄H₁₀NaO₃ [M+Na⁺]+: 249.0522; found: 249.0532.

Synthesis of 2,2'-((4-hydroxybiphenyl-3,5-diyl)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5c). A solution of 4-hydroxybiphenyl-3,5-dicarbaldehyde 4c (1.50 g, 6.63 mmol) and 2-amino-3-nitrobenzoic acid (2.42 g, 13.3 mmol) in EtOH (53 mL) was prepared by heating the mixture to 70 °C. This orange solution was brought to r.t. before addition of a freshly prepared solution of Na₂S₂O₄ (85 wt%, 8.15 g, 39.8 mmol) in H₂O (40 mL) after which the resulting red solution was heated for 5 h. Filtration and washing with EtOH (40 mL), H₂O (75 mL) and Et₂O (100 mL) with subsequent drying in a vacuum yielded the desired ligand as a red powder (2.58 g, 79.1%). 1H NMR (500 MHz, [D₆]DMSO): δ = 8.73 (s, 2H), 7.95 (d, J = 7.92 Hz, 2H), 7.81 (d, J = 7.58 Hz, 2H), 7.79 (d, J = 7.55 Hz, 2H), 7.51 (t, J = 7.55 Hz, 2H), 7.33 (t, J = 7.67 Hz, 3H); 13C NMR (100 MHz, [D₆]DMSO): δ = 166.9, 157.1, 153.0, 141.5, 139.5, 134.3, 128.9, 128.7, 128.4, 126.6, 126.0, 124.2, 121.8, 116.1, 115.3; HRMS (ESI) calcd for C₂₈H₁₉N₄O₅ [M+H⁺]: 491.1350; found: 491.1369.

Synthesis of dinuclear Mn²⁺,³⁺ complex 6c. Ligand 5c (1.00 g, 2.04 mmol), NaOAc (1.67 g, 20.4 mmol) and Mn(OAc)₂·C₄H₂O (1.25 g, 5.10 mmol) were suspended in MeOH (40 mL) and refluxed for 24 h. The product was retrieved by centrifugation, washed with MeOH (4×45 mL) and dried under vacuum to give the complex as a yellow powder (1.20 g, 77.5%). HRMS (ESI) calcd for C₃₀H₁₈Mn₂N₄O₈ [M–CH₃OH+]: 671.9891; found: 671.9892; anal. calcd for C₃₁H₂₉Mn₂N₄O₁₂ [6c·4H₂O]: C 49.03, H 3.85, N 7.38, Mn 14.47%; found: C 48.51, H 3.42, N 7.22, Mn 14.63%; IR (KBr): ν = 3412, 3174, 2955, 2925, 2854, 1602, 1555, 1506, 1491, 1468, 1384, 1315, 1252, 1211, 1155, 1023, 1003, 892, 803, 763, 700, 662, 602, 496 cm⁻¹.

Synthesis of 5-benzyl-2-hydroxyisophthalaldehyde (4d). 4-Benzylphenol 3d (1.47 g, 8.00 mmol) and hexamethylenetetramine (22.4 g, 160 mmol) were placed in a round-bottomed flask which was purged with nitrogen. After the addition of trifluoroacetic acid (48 mL), the mixture was refluxed at 110 °C for 72 h. The reaction mixture was then combined with 4 M HCl (240 mL) in a 500 mL round-bottomed flask and stirred for 3 h. The yellow precipitate was collected by filtration, washed with H₂O (500 mL) and dried under vacuum to yield the product as a yellow powder (1.49 g, 77.4%). 1H NMR (500 MHz, CDCl₃): δ = 11.50 (s, 1H), 10.20 (s, 2H), 7.78 (s, 2H), 7.36–7.12 (m, 5H), 4.00 (s, 2H); 13C NMR (100 MHz, [D₆]DMSO): δ = 192.2, 162.4, 139.7, 137.9, 133.4, 129.0, 128.9, 126.9, 124.5, 122.0, 115.6, 115.2, 40.7; HRMS (ESI) calcd for C₁₅H₁₂NaO₃ [M+Na⁺]: 263.0679; found: 263.0681.

Synthesis of 2,2'-((5-benzyl-2-hydroxy-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5d). A solution of 5-benzyl-2-hydroxyisophthalaldehyde 4d (1.20 g, 5.00 mmol) and 2-amino-3-nitrobenzoic acid (1.82 g, 9.99 mmol) in EtOH (40 mL) was prepared by heating the mixture to 70 °C. This orange solution was brought to r.t. before addition of a freshly prepared solution of Na₂S₂O₄ (85 wt%, 6.14 g, 30.0 mmol) in H₂O (30 mL) and the resulting red solution was heated for 5 h. Filtration, and washing with EtOH (16 mL), H₂O (25 mL) and Et₂O (150 mL) with subsequent drying in a vacuum yielded the desired ligand as a red powder (2.05 g, 82.0%). 1H NMR (500 MHz, [D₆]DMSO): δ = 8.38 (s, 2H), 7.95 (d, J = 7.89 Hz, 2H), 7.83 (d, J = 7.68 Hz, 2H), 7.43–7.29 (m, 6H), 7.25–7.18 (m, 1H); 13C NMR (100 MHz, [D₆]DMSO): δ = 192.2, 164.2, 139.7, 137.9, 133.4, 129.0, 128.9, 126.9, 123.3, 40.7; HRMS (ESI) calcd for C₂₉H₂₁N₄O₅ [M+H⁺]: 505.1506; found: 505.1501.
Synthesis of dinuclear Mn$_{II,III}$ complex 6d. Ligand 5d (1.00 g, 1.98 mmol), NaOAc (1.62 g, 19.8 mmol) and Mn(OAc)$_2$·4H$_2$O (1.21 g, 4.94 mmol) were suspended in MeOH (40 mL) and refluxed for 24 h. The product was retrieved by centrifugation, washed with MeOH (4×45 mL) and dried under vacuum to yield a yellow powder (1.19 g, 77.7%). HRMS (ESI) calcd for C$_{30}$H$_{18}$Mn$_2$N$_4$O$_6$ [M–CH$_3$COOH–H$^+$]: 639.9993; found: 639.9999; anal. calcd for C$_{32}$H$_{31}$Mn$_2$N$_4$O$_{12}$ [6d·4H$_2$O]: C 49.69, H 4.04, N 7.24, Mn 14.21%; found: C 49.25, H 3.92, N 7.02, Mn 14.45%; IR (KBr): $\nu$ = 3426, 3190, 2924, 2854, 1603, 1556, 1495, 1473, 1385, 1319, 1251, 1211, 1155, 1052, 1015, 892, 765, 700, 662, 536, 491 cm$^{-1}$.

Synthesis of 2,2′-(5-((1,3-dioxoisindolin-2-yl)methyl)-2-hydroxy-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5e). 2-Amino-3-nitrobenzoic acid (0.360 g, 2.00 mmol) and 5-((1,3-dioxoisindolin-2-yl)methyl)-2-hydroxyisophthalaldehyde (4e, 0.310 g, 1.00 mmol) were suspended in EtOH (30 mL). After addition of a freshly prepared aqueous solution of Na$_2$S$_2$O$_4$ (1.23 g, 6.00 mmol, 6.0 mL), the reaction mixture was gently heated at reflux for 24 h. The yellow precipitate was filtered, washed with H$_2$O, acetone and finally dried under vacuum to afford the title compound (0.450 g, 78.5%). HRMS (ESI) calcd for C$_{30}$H$_{18}$Mn$_2$N$_4$O$_6$ [M–CH$_3$COOH–H$^+$]: 639.9993; found: 639.9999; anal. calcd for C$_{32}$H$_{31}$Mn$_2$N$_4$O$_{12}$ [6d·4H$_2$O]: C 49.69, H 4.04, N 7.24, Mn 14.21%; found: C 49.25, H 3.92, N 7.02, Mn 14.45%; IR (KBr): $\nu$ = 3426, 3190, 2924, 2854, 1603, 1556, 1495, 1473, 1385, 1319, 1251, 1211, 1155, 1052, 1015, 892, 765, 700, 662, 536, 491 cm$^{-1}$.

Synthesis of 2,2′-(5-((5-carboxypentanamido)methyl)-2-hydroxy-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5f). Compound 5e (0.770 g, 1.73 mmol) was dissolved in pyridine (20 mL) and the resulting solution was cooled to 0 °C. Adipoyl chloride (0.330 g, 1.81 mmol) was added dropwise to this cold solution and the reaction mixture was then stirred overnight at room temperature. Evaporation of the solvent under reduced pressure gave a crude product that was washed with aqueous 2 M HCl (2×10 mL) and Et$_2$O (3×10 mL) to give the product.
(0.700 g, 85.9%). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 13.78 (br, 3H), 8.37 (s, 2H), 8.10–7.67 (m, 4H), 7.44 (s, 2H), 4.33 (s, 2H), 2.23 (s, 2H), 2.07 (s, 2H), 1.71–1.35 (m, 4H); $^{13}$C NMR (100 MHz, [D$_6$]DMSO): $\delta$ = 174.4, 172.1, 166.3, 150.8, 150.3, 132.3, 131.6, 125.7, 123.7, 120.0, 116.2, 41.7, 35.2, 33.5, 24.8, 24.3; HRMS (ESI) calcd for C$_{29}$H$_{25}$N$_5$NaO$_8$ [M+Na$^+$]: 594.1595; found: 594.1590.

Synthesis of dinuclear Mn$_{2}^{II,III}$ complex 6f. NaOAc (0.110 g, 1.32 mmol) and Mn(OAc)$_2$4H$_2$O (0.080 g, 0.330 mmol) were added to a suspension of ligand 5f (0.080 g, 0.140 mmol) in MeOH (3 mL). The reaction mixture was then heated at 70 °C for 20 h. The yellow precipitate was filtered, washed with MeOH (3 × 10 mL), and dried under vacuum to yield the title compound (0.700 g, 85.9%). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 13.29%; found: C 46.25, H 3.59, N 8.40, Mn 13.38%; IR (KBr): $\nu$ = 3426, 3191, 1603, 1564, 1502, 1400, 1383, 1263, 1003, 774, 751, 731, 672, 658, 618, 495 cm$^{-1}$.

Synthesis of 2,2′-(2-hydroxy-5-(4′-methyl-2,2′-bipyridine-4-carboxamido)methyl)-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5g). Pivaloyl chloride (1.22 mL, 10.0 mmol) was carefully added to a nitrogen purged solution of compound 5e (0.443 g, 10.0 mmol) in pyridine (15 mL) at 0 °C. After the reaction mixture was stirred at room temperature overnight. Upon pouring the reaction mixture into crushed ice, a yellow precipitate was formed, which was filtered, washed with H$_2$O (2 × 10 mL), Et$_3$N (3 × 10 mL), and dried under vacuum to yield the title compound (0.190 mmol). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 13.79%; found: C 46.77, H 4.50, N 8.47, Mn 13.29%; IR (KBr): $\nu$ = 3426, 3191, 1603, 1564, 1502, 1400, 1383, 1263, 1003, 791, 751, 672, 658, 618, 495 cm$^{-1}$.

Synthesis of 2,2′-(2-hydroxy-5-(pivalamidomethyl)-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5h). A solution of compound acid65 (0.150 g, 0.700 mmol) was refluxed in 10 mL of SOCl$_2$. The excess of SOCl$_2$ was removed under reduced pressure. The remaining residue (acid chloride) was dissolved (2 × 10 mL) and dried under vacuum to give the title compound (0.100 g, 0.700 mmol) was added to a solution containing Mn(OAc)$_2$:4H$_2$O (0.116 g, 0.474 mmol) and NaOAc (0.155 g, 1.90 mmol) in MeOH (4 mL). The reaction mixture was refluxed for 20 h and then centrifuged. The precipitate was washed with MeOH (5 × 5 mL) and dried under vacuum to afford complex 6g (0.123 g, 81.3%). HRMS (ESI) calcd for C$_{32}$H$_{31}$Mn$_2$N$_5$O$_{13}$ [M+Na$^+$]: 747.0540; found: 747.0576; anal. calcd for C$_{31}$H$_{28}$Mn$_2$N$_5$O$_9$: 767.0474; found: 767.0470; anal. calcd for C$_{32}$H$_{27}$Mn$_2$N$_5$O$_{11}$ [M+Na$^+$]: 781.0505; found: 781.0500; anal. calcd for C$_{31}$H$_{26}$Mn$_2$N$_5$O$_{13}$ [Mg$^+$H$_2$O]: C 46.74, H 4.56, N 8.79, Mn 13.79%; found: C 46.77, H 4.50, N 8.74, Mn 13.83%; IR (KBr): $\nu$ = 3435, 1564, 1498, 1474, 1400, 1385, 1316, 1263, 1003, 765, 662 cm$^{-1}$.

Synthesis of 2,2′-(2-hydroxy-5-((4′-methyl-2,2′-bipyridine-4-carboxamido)methyl)-1,3-phenylene)bis(1H-benzo[d]imidazole-4-carboxylic acid) (5i). 4′-Methyl-2,2′-bipyridine-4-carboxylic acid$^{5}$ (0.150 g, 0.700 mmol) was refluxed in 10 mL of SOCl$_2$ for 4 h. The excess of SOCl$_2$ was removed under reduced pressure. The remaining residue (acid chloride) was dissolved in 2.0 mL of dry DMF. The resulting reaction mixture was stirred at room temperature overnight. Upon pouring the reaction mixture into crushed ice and acidification with conc. HCl, a yellow precipitate was formed, which was filtered, washed with acetone (2 × 10 mL) and dried under vacuum to give the title compound (0.410 g, 89.1%). $^1$H NMR (400 MHz, [D$_6$]DMSO): $\delta$ = 13.29 (br, 2H), 8.84 (s, 2H), 8.56 (s, 2H), 8.40 (s, 1H), 8.25 (s, 1H), 8.00–7.89 (m, 5H), 7.61–7.40 (m, 3H), 4.45 (s, 2H), 2.41 (s, 3H); $^{13}$C NMR (100 MHz, [D$_6$]DMSO): $\delta$ = 166.8, 166.2, 156.4, 154.2, 151.6, 150.3, 150.0, 134.5, 139.3, 138.3, 131.1, 125.4, 125.0, 122.9, 122.4, 121.7, 121.3, 119.6, 118.2, 115.8, 42.5, 20.7; HRMS (ESI) calcd for C$_{32}$H$_{35}$N$_5$NaO$_6$ [M+Na$^+$]: 550.1697; found: 550.1703.
Synthesis of dinuclear MnII,III complex 6h. NaOAc (0.143 g, 1.75 mmol) and Mn(OAc)₂·4H₂O (0.107 g, 0.438 mmol) were added to a suspension of ligand 5h (0.112 g, 0.175 mmol) in MeOH (4 mL) and the reaction mixture was heated at 70 °C for 20 h. The resulting precipitate was centrifuged, washed with MeOH (3 × 5 mL) and dried under vacuum to afford the complex (0.125 g, 79.0%). HRMS (ESI) calcd for C₁₉H₂₆Mn₂N₇O₁₂ [M+Na⁺⁺]: 859.0602; found: 859.0608; anal. calcd for C₃₉H₃₆Mn₂N₇O₁₂: 662.1759; found: 662.1760.

3. Results and discussion

3.1. Synthesis

To investigate how the electronics affected the Mn centers in WOC 2, eight different dinuclear Mn complexes 6a–h, depicted in Scheme 1, were synthesized, where a variety of substituents were introduced. These alterations would hence result in tuning of the corresponding Mn complexes and thereby affect the properties and reactivities. The synthesis of the functionalized ligand scaffolds involved the reductive cyclization reaction between the appropriate phenolic dialdehyde 4 and 2-amino-3-nitrobenzoic acid, with Na₂S₂O₄ as the reducing agent. This allowed for easy access to a wide variety of ligands in good to excellent yields. The Mn complexes 6a–h were subsequently obtained by refluxing the ligands in the presence of Mn(OAc)₃ and NaOAc. It is believed that all of the synthesized complexes were isolated in their MnII,III state, in conformity with complex 2. This is based on elemental analyses and high-resolution mass spectrometry (HRMS). However, extensive attempts to prove this by EPR have failed so far since we have been unable to observe the characteristic EPR signal for MnII,III complexes. Instead, a broad signal at around g = 2 was observed both in solution and in the solid state, which is typical for uncoupled MnII and reminiscent of the signal recently observed for an octahedral MnII complex. This signal decreased in intensity upon the addition of the one-electron oxidant [Ru(bpy)₃]⁺. It therefore seems possible that the complexes are in fact polymeric or at least tetranuclear, as indicated by the generated crystal structure of 2′ from complex 2. The formation of the tetranuclear complex 2′ indicates that the acetate ligand in complex 2, and probably also in complexes 6, is somewhat labile and we assume that on dissolution in phosphate buffer, the acetate ligand is rapidly replaced by phosphate to yield the active catalyst. On dissolution of the dinuclear Mn complexes in aqueous solutions containing PO₄³⁻, in the presence of air, new signals appeared, as detected by HRMS. These signals corresponded to the formation of MnII,III complexes, which are expected to be EPR silent. However, this will be studied in more detail in future work.

3.2. Catalytic H₂O oxidation by use of the mild one-electron oxidant [Ru(bpy)₃]⁺⁺

A majority of the previously developed artificial WOCs require the use of the strong one-electron oxidant CeIV to drive H₂O oxidation. However, this approach is not useful in solar fuel devices since the oxidant cannot be regenerated photochemically, using sunlight as the terminal energy source. For this to be realized, CeIV and related oxidants have to be substituted by photosensitizer-based oxidants, such as the well-studied [Ru(bpy)₃]⁺⁺-type complexes (bpy = 2,2’-bipyridine). For this more preferable approach to be thermodynamically possible, the redox potential of the WOC has to be matched with that of the photosensitizer.

Since the previously developed Mn WOC 2 could promote catalytic H₂O oxidation with the mild one-electron oxidant [Ru(bpy)₃]⁺⁺ (E (RuIII/RuII) = 1.26 vs. NHE) as a pre-generated or photochemically generated, the catalytic experiments for the newly developed complexes 6 were conducted as previously established; the [Ru(bpy)₃]⁺⁺ oxidant was employed as the chemical oxidant in a buffered aqueous solution, under neutral conditions (0.1 M phosphate buffer, pH 7.2). The evolution of gaseous products was monitored and quantified by real-time mass spectrometry, a technique which we have previously used. In a typical reaction, [Ru(bpy)₃]⁺⁺ was added to an aqueous solution containing the Mn complex, triggering immediate O₂ evolution.

Several control experiments were conducted in order to verify that the observed O₂ evolution was mediated by the investigated Mn complexes; (1) no O₂ was generated when the Mn complex was omitted. The result was only degradation of the [Ru(bpy)₃]⁺⁺ oxidant, without any evolution of O₂. (2) To ensure that free Ru, possibly originating from the decomposition of the [Ru(bpy)₃]⁺⁺ oxidant, did not react with the free ligand to generate an active catalyst, an aqueous solution containing the free ligand was added to [Ru(bpy)₃]⁺⁺. However, this resulted in no observable O₂ evolution. Collectively, this supports that the detected O₂ formation was caused by the Mn complexes and not by any other unexpected catalytic process.

Fig. 2 and Table 1 show the O₂ evolution activity of Mn complexes 6. It is evident that Mn complexes 6a–e all have similar O₂ evolution activity, which highlights that the introduced substituents in 6a–e did not affect the catalytic activity to a large extent. The previously reported Mn complex 2 was also evaluated under these reaction conditions and was found to display comparable activity as catalysts 6a–e (not shown).
Scheme 1 Synthetic routes to ligands 5a–h and dinuclear Mn\textsuperscript{II,III} complexes 6a–h.
However, Mn complex 6f, housing the long aliphatic chain with a terminal carboxylate group, displayed higher catalytic activity than the other complexes. This effect could not be ascribed to significantly decreased redox potentials in complex 6f (vide infra), which would improve the O–O bond formation and subsequent O$_2$ formation. To additionally probe how big influence the substituent in complex 6f had on the electronics, it was decided to synthesize the related Mn complex 6g. Although the two complexes 6f and 6g have electronic resemblance, they do not at all display the same catalytic activity towards oxidizing H$_2$O. This highlights that there is a different reason as to why Mn complex 6f displays a higher catalytic activity and thus facilitates the oxidation of H$_2$O. To further establish the catalytic importance of the substituent in complex 6f, we turned our attention to electrochemistry in the hope that this would reveal the intrinsic reason for the better efficiency of this complex.

It should also be noted that the bpy functionalized Mn complex 6h was found to be catalytically active. This complex is a precursor of fundamental importance and could potentially be used for the construction of coupled photosensitizer-WOC systems for incorporation into devices for solar to chemical energy conversion (Fig. 3).

### 3.3. Electrochemical measurements

In order to assess the effect of the different substituents on the electronic properties of Mn complexes 6, it was decided to

| Table 1 | Catalytic data for dinuclear Mn$_{II,III}$ complexes 6a–h$^a$ |
|---|---|
| Complex | Initial rate of O$_2$ evolution$^b$ (nmol O$_2$ s$^{-1}$) | Turnover number (mol O$_2$ per mol catalyst) |
| 6a | 0.42 | 4.4 |
| 6b | 0.30 | 5 |
| 6c | 0.30 | 5 |
| 6d | 0.30 | 5 |
| 6e | 0.83 | 6.6 |
| 6f | 0.30 | 4.6 |
| 6g | 0.50 | 12 |
| 6h | 0.30 | 5 |
| 6i | 0.42 | 7 |

$^a$ Reactions were carried out in aqueous buffered solutions (phosphate buffer; 0.1 M, pH 7.2, 0.50 mL) containing the catalyst (20 μM, 10 nmol) and the oxidant [Ru(bpy)$_3$](PF$_6$)$_3$ (6.8 mM, 3.4 μmol). $^b$ Initial rates were obtained by linear fitting of the initial reaction (10–60 s).

Fig. 2 O$_2$ evolution kinetics by Mn complexes 6 as a function of time. Reaction conditions: an aqueous phosphate buffer solution (0.1 M, pH 7.2, 0.50 mL) containing the Mn complex (20 μM) was added to the oxidant [Ru(bpy)$_3$](PF$_6$)$_3$ (3.4 mg, 3.4 μmol) and the evolved O$_2$ was measured in real-time. 6a (--), 6b (--), 6c (--), 6d (--), 6e (--), 6f (--), 6g (--), 6h (--).

Fig. 3 Schematic pathway from the dinuclear Mn$_{II,III}$ complex 6h to a coupled Ru-Mn$_2$ assembly for use in a solar fuel device.

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examine the electrochemical properties of the synthesized Mn complexes 6a,b and 6e–h. Electrochemical measurements were carried out at neutral conditions (pH 7.2), under similar conditions to the catalytic H₂O oxidation experiments in order to obtain a deeper understanding of how the different substituents influenced the properties of the dinuclear Mn₂²⁺,³⁺ complexes. Cyclic voltammetry (CV) and differential pulse voltammetry (DPV) were used to analyze the redox properties. Cyclic voltammograms revealed catalytic currents, assigned to electrochemical oxidation of H₂O, for all of the studied Mn complexes 6 (Fig. 4). However, it was difficult to clearly assign the onset potentials due to overlapping redox processes in the onset potential region.

Therefore, to gain further insight into the redox chemistry of the newly synthesized Mn complexes 6, DPV was carried out on the complexes. The differential pulse voltammograms of the dinuclear Mn complexes 6 are depicted in Fig. 5 and displayed several peaks. Within the developed library, the differential pulse voltammograms of complexes 6f and 6h displayed five discernible peaks in the region 0.50 < E < 1.30 V vs. NHE (Table 2). These peaks were subsequently assigned to the formal oxidations of Mn₂²⁺,³⁺ → Mn₂³⁺,³⁺ → Mn₂³⁺,⁴⁺ → Mn₂⁴⁺,⁴⁺ → Mn₂⁵⁺,⁵⁺, in conformity with the previously reported result for a related dinuclear Mn complex.⁵¹ However, the other complexes displayed less than five redox peaks. This is probably related to the close potential gap, i.e. the narrow potential range for some of the redox couples, resulting in single observable two-electron redox peaks instead of two single-electron waves.

As is evident from these voltammograms and Table 2, the substituents in Mn complexes 6 did not cause any dramatic shift of the different redox processes in the complexes. This highlights that there must be another explanation for the increased efficiency of Mn complex 6f, containing the distal carboxyl group. Previous studies of so-called “Hangman” porphyrin and corrole complexes housing distal hydrogen bonding groups, such as carboxyl groups, have established the possibility of accessing metal complexes with increased affinity for promoting
PCET reactions.\textsuperscript{37,45,78–81} There even exists an example of an Fe complex comprising a Hangman platform, which in the solid state exhibits a hydrogen bonded H\textsubscript{2}O molecule bound between a distal xanthene carboxylic acid and an Fe porphyrin hydroxide by two hydrogen bonds (Fig. 6).\textsuperscript{78} This suggested that the carboxyl group in \textit{6f} could perhaps also interact with the Mn centers. To confirm this hypothesis, quantum chemical calculations were performed on complex \textit{6f}.

### 3.4. Quantum chemical insight

DFT calculations were performed on the dinuclear complex \textit{6f} in the relevant formal Mn\textsubscript{2}\textsuperscript{IV,IV} state. The Mn\textsubscript{2}\textsuperscript{IV,V} stage is believed to be capable of promoting O–O bond formation, as shown in the previous calculations on the dinuclear Mn complex [(H\textsubscript{2}O)(tpy)Mn(μ-O)\textsubscript{2}Mn(tpy)(OH\textsubscript{2})\textsuperscript{3+} (tpy = 2,2\textsubscript{0},6\textsubscript{0},2\textsubscript{00}-terpyridine).\textsuperscript{82} The presence of one high-valent Mn\textsuperscript{V} (or Mn\textsuperscript{IV–O}/C\textsubscript{15}) is also suggested in the OEC in the formation of O\textsubscript{2}.\textsuperscript{83–85} The optimized structures of the two low-energy isomers of complex \textit{6f} in its high-valent Mn\textsubscript{2}\textsuperscript{IV,V} state (A and B) are depicted in Fig. 7.

Both structures have a total charge of +0 and exist in the high spin octet state according to the calculations. The two Mn centers are bridged by an oxo group, an acetate, and the phenolate of the ligand. In addition, each Mn center also has a coordinated hydroxide ligand. Spin density analysis suggests

### Table 2  Electrochemical data for the dinuclear Mn complexes \textit{6}\textsuperscript{a}

| Complex | Mn\textsubscript{2}\textsuperscript{III,III}/Mn\textsubscript{2}\textsuperscript{II,III} (V vs. NHE) | Mn\textsubscript{2}\textsuperscript{III,IV}/Mn\textsubscript{2}\textsuperscript{III,III} (V vs. NHE) | Mn\textsubscript{2}\textsuperscript{IV,IV}/Mn\textsubscript{2}\textsuperscript{IV,IV} (V vs. NHE) | Mn\textsubscript{2}\textsuperscript{IV,V}/Mn\textsubscript{2}\textsuperscript{IV,IV} (V vs. NHE) |
|---------|-----------------------------------|-----------------------------------|-----------------------------------|-----------------------------------|
| \textit{6a} | 0.55                              | 0.78\textsuperscript{b}           | —                                 | 1.22\textsuperscript{b}           |
| \textit{6b} | 0.51                              | 0.86\textsuperscript{b}           | —                                 | 1.13                              |
| \textit{6c} | —                                 | 0.79\textsuperscript{b}           | 1.54                             | 1.21                              |
| \textit{6f} | 0.53                              | 0.79                              | 1.88                             | 1.17                              |
| \textit{6g} | 0.51                              | 0.85\textsuperscript{b}           | —                                 | 1.19\textsuperscript{b}           |
| \textit{6h} | 0.52                              | 0.78                              | 0.89                             | 1.18                              |

\textsuperscript{a} Potentials were obtained from differential pulse voltammetry. Conditions: measured in an aqueous phosphate buffer solution (0.1 M, pH 7.2), Scan rate 0.1 V s\textsuperscript{-1}, glassy carbon disk as a working electrode, a platinum spiral as a counter electrode and an Ag/AgCl electrode as a reference electrode. Potentials were converted to NHE by using the [Ru(bpy)]\textsubscript{3}\textsuperscript{3+}/[Ru(bpy)]\textsubscript{3}\textsuperscript{2+} couple as a standard (\textit{E}_{1/2} = 1.26 V vs. NHE). \textsuperscript{b} Possible two-electron redox process.

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**Fig. 6** (a) Structure of a metal “Hangman” porphyrin housing a distal carboxyl group and (b) representation of the hydrogen bonded H\textsubscript{2}O molecule bound between a distal xanthene carboxylic acid and a Fe porphyrin hydroxide by two hydrogen bonds.

**Fig. 7** Optimized structures of the dinuclear Mn complex \textit{6f} in its formal Mn\textsubscript{2}\textsuperscript{IV,V} state, showing the hydrogen bonding interaction between the distal carboxyl group and (a) the Mn bounded hydroxide(s), and (b) the imidazole moiety. Distances are given in Ångstrom, spin densities on Mn are given in italics, and relative energies are given in kcal mol\textsuperscript{-1}.
that both metals are in fact in their Mn$^\text{IV}$ state, with a coordinated ligand radical cation. This suggests that the ligand framework is redox-active, non-innocent, and highlights the intriguing properties of ligand scaffolds of this type. Redox-active ligands might thus be of importance in alleviating the metal centers from being heavily oxidized by storing redox equivalents at the ligand scaffolds during the oxidation of H$_2$O.

In structure A of the oxidized complex 6f, the distal carboxyl group generates two hydrogen bonds to the two Mn coordinated hydroxide ligands. These two hydrogen bonds might thus influence the subsequent O–O bond forming step(s). This supports that the distal carboxylate group in the ligand framework of Mn complex 6f can accommodate a bridging hydroxide/aqua molecule to yield a unique hydrogen-bonded scaffold that might facilitate H$_2$O oxidation and thus promote the crucial O–O bond formation through PCET. Multi-electron catalysis is at the heart of H$_2$O oxidation and the ability of synchronizing proton and electron transfer events via PCET is fundamental in a plethora of biological and chemical reactions. PCET allows the charge of a system to remain unchanged, whereas non-proton coupled single electron transfer processes result in charge accumulation and high-energy intermediates. Detailed studies and a fundamental understanding of how to couple proton and electron transfer are at the frontier and might be the key to realizing more efficient WOCs.

An additional interesting feature of the ligand platforms in Mn complexes 6 is the presence of the imidazole motifs, which have the ability to function as proton transfer mediators. In structure B, the carboxyl group has abstracted a proton from the imidazole unit. Structure B was found to be 2.0 kcal mol$^{-1}$ higher in energy than structure A. This hydrogen bonding capability of the distal carboxylate group renders it possible to also promote proton transfer reactions between the distal group and the protonated imidazole moiety. The alternative scenario for the enhanced activity of complex 6f is that the carboxyl group serves as a ligand to bridge the two metal centers. However, calculations revealed that the chain is not sufficient to accommodate this sort of coordination.

The possibility of including a non-innocent distal group within the ligand scaffolds in Mn complexes 6 is an interesting feature in these developed WOCs that has perhaps not received sufficient attention in H$_2$O oxidation schemes. The key feature of incorporating a distal group that acts as a general base to stimulate a H$_2$O molecule in undergoing nucleophilic attack on a high-valent metal-oxo species could be a general strategy for promoting catalytic H$_2$O oxidation activity. This idea of hydrogen bonding as a means to increase the nucleophilicity of H$_2$O is utilized in hydrolytic enzymes such as chymotrypsin. It has also been used by Nocera and ourselves to promote H$_2$O oxidation and also recently in two theoretical papers. In those previous studies, rather elaborate rigid systems have been used. However, our present results suggest that even a carboxyl attached to a highly flexible chain will participate efficiently. Using this concept we will therefore try to prepare analogues to complex 6f, and also analogues with a different central ligand, using the synthetic versatility available for modifying the ligand frameworks of the complexes 6.

4. Conclusions

In the present work we have synthesized eight new dinuclear Mn$^{II,III}$ complexes and examined the H$_2$O oxidation process for these complexes. It was demonstrated that all of the developed Mn complexes were catalytically active in H$_2$O oxidation. The complexes had sufficiently low redox potentials to allow for the use of the mild one-electron oxidant [Ru(bpy)$_3$]$^{2+}$ to drive H$_2$O oxidation at neutral pH conditions.

These studies have identified a potential general factor for obtaining WOCs that display increased catalytic efficiencies. Within the library of the synthesized Mn complexes, complex 6f containing the long aliphatic chain with a terminal carboxylate group as a distal group was found to exhibit a higher O$_2$ evolution rate than the other catalysts. This observation did not originate from decreased redox potentials, which would result in more favourable thermodynamics for carrying out H$_2$O oxidation.

Instead, the improved catalytic effect seems to result from pre-orientation of the incoming H$_2$O nucleophile and hydrogen bonding between the distal carboxyl group and a high-valent Mn-hydroxy group. This hydrogen bonding interaction can subsequently promote PCET processes, which decrease the energy of the catalytic H$_2$O oxidation process.

These findings highlight the importance of controlling both proton and electron transfer processes to promote O–O bond formation during H$_2$O oxidation. In the natural photosynthetic system, PCET is prevalent and provides the key to efficient H$_2$O oxidation catalysis. The present study provides a guide that might be helpful for designing more efficient WOCs in the future, which have the possibility of being incorporated into large scale systems for solar to chemical energy conversion.

Acknowledgements

Financial support from the Swedish Energy Agency (BÅ and JM), the Knut and Alice Wallenberg Foundation (Center of Molecular Catalysis at Stockholm University, Artificial Leaf Project Umeå), the Swedish Research Council (BÅ) and the Strong Research Environment Solar Fuels (Umeå University) is gratefully acknowledged. W. A. A. would also like to thank the Egyptian Government for a PAROWN research fellowship.

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