Tuning the Electronic Properties of Prussian Blue Analogues for Efficient Water Oxidation Electrocatalysis: Experimental and Computational Studies

Elif Pınar Alsaç, Emine Ülker, Satya Vijaya Kumar Nune, Yavuz Dede, and Ferdi Karadas

Abstract: Although several Prussian Blue analogues (PBAs) have been investigated as water oxidation catalysts, the field lacks a comprehensive study that focuses on the design of the ideal PBA for this purpose. Here, members of a series of PBAs with different cyanide precursors have been investigated to study the effect of hexacyanometal groups on their electrocatalytic water oxidation activities. Cyclic voltammetric, chronoamperometric, and chronopotentiometric measurements have revealed a close relationship between the electron density of electroactive cobalt sites and electrocatalytic activity, which has also been confirmed by infrared and XPS studies. Furthermore, pH-dependent cyclic voltammetry and computational studies have been performed to gain insight into the catalytic mechanism and electronic structure of cyanide-based systems to identify possible intermediates and to assign the rate-determining step of the target process.

Introduction

Increase in energy demand has propelled the scientific community, particularly in the last two decades, to find alternative energy sources that will replace limited fossil-based fuels. Since solar energy that involves the production of H₂ from water has been one of the most promising candidates among sustainable sources of energy, much effort has recently been devoted to investigating efficient methods for splitting water. Despite several PBA families, including metal oxides, perovskites, amorphous materials, noble-metal-based materials, and metal-organic frameworks (MOFs), have been investigated as water oxidation catalysts (WOCs). Of these, cobalt oxides stand out due to their high catalytic activities, but are also associated with two main disadvantages: low stability and a high tendency to decompose in acidic media. Many inorganic systems, including metal oxides, perovskites, amorphous materials, noble-metal-based materials, and metal-organic frameworks (MOFs), have been investigated as WOCs due to their favorable characteristics, such as ease of preparation, stability over a wide pH range, and robustness during catalytic processes. Patzke et al. reported a carbodiimide-based material that could be used as a WOC, which is stable in acidic and neutral media. Members of a similar class of materials, metal dicyanamides, have also been shown to be promising candidates for water oxidation electrocatalysis. Cobalt hexacyanoferrates, members of the Prussian Blue analogue (PBA) family, are also exceptional candidates for electrocatalytic water oxidation due to their high catalytic activities, robustness, and stability at neutral pH. A further study by Patzke et al. showed that PBAs can also be used for a light-driven water oxidation process in the presence of [Ru(bpy)₃]⁺ as a chromophore. Despite their high turnover frequencies (TOFs), one of the main drawbacks of cyanide-based systems is their low concentration of electroactive cobalt sites. This is because of the relatively large distances between Co sites (ca. 10 Å) compared to oxide-based systems (ca. 3 Å). This problem has recently been overcome by our group with the use of a novel pentacyanoferrate-bound polymer as a precursor of Co-Fe PBAs, which resulted in a dramatic decrease in the crystallinities of PBAs, and thus a significant increase in the surface concentration of Co sites. Galán-Mascarós et al. approached the same problem by using a new synthetic method for the preparation of thin films of PBAs, which
involves chemical etching of cobalt oxides with a hexacyanoferrate solution to form an in situ PBA film. This novel method led to an impressive improvement in the stability of the electrode and electrocatalytic performance over a wide pH range. It required a much lower overpotential (510 mV) to obtain a current density of 1 mA cm⁻². In addition, Fukuzumi et al. investigated the photocatalytic water oxidation performances of a series of Co-Pt PBAs in the presence of the well-defined [Ru(bpy)₃]²⁺/S²O₈²⁻ couple. A systematic study performed with [Co(CN)₆]³⁻ and [Pt(CN)₆]⁴⁻ groups in different stoichiometries clearly showed the number of active sites to be highly dependent on the number of defects. Fukuzumi and co-workers also studied the effect of counter cations on the catalytic activity and quantum efficiency displayed by Co-Co PBAs in the photocatalytic water oxidation process, and showed that a quantum efficiency of 200% could be achieved with such systems incorporating calcium ions as counter cations.

The previous studies mentioned above have clearly shown that slight modifications in the structure of PBAs can lead to a significant increase in their catalytic activities. Although previous studies have taken advantage of the rich and well-established cyanide chemistry, to the best of our knowledge, no study has hitherto been performed to investigate the effect of hexacyanometal units on the electronic properties and catalytic performances of electroactive cobalt sites. In the present study, electrocatalytic measurements on a series of cobalt hexacyanometalates (CHCMs) incorporating different M(CN)₆ units (M = Co, Cr, and Fe) together with characterization studies, have been performed to investigate the effect of the type and oxidation state of the metal in the M(CN)₆ unit on the catalytic properties of PBAs. The effect of hexacyanometal groups on the electronic properties of electroactive cobalt sites has further been examined through electronic structure calculations employing density functional theory (DFT).

Results and Discussion

Electrochemistry

All of the electrochemical experiments were conducted with a PBA-modified fluorine-doped tin oxide (FTO) electrode. Cyclic voltammograms (CVs) of Co[M(CN)₆] (M = Co, Cr, and Fe) were measured in a phosphate buffer containing 1 M KNO₃ as the electrolyte in the potential range 0.2–1.7 V versus NHE (Figure 1). [Co⁵⁻Co⁷⁺] exhibits a quasi-reversible redox couple with an oxidation peak at 1.210 V and a reduction peak at 1.031 V versus NHE that can be assigned to Co²⁺/Co³⁺. Similar redox couples are also observed for the other PBAs. Another peak at a more positive potential of around 1.415 V versus NHE is observed for [Co⁵⁻Co⁷⁺], which can be assigned to the Co³⁺/Co⁴⁺ redox process. Tafel plots for each catalyst were performed to investigate the effect of the type and oxidation state of the metal in the Co(CN)₆ unit on the catalytic measurements on a series of cobalt hexacyanometalates (CHCMs) incorporating different M(CN)₆ units (M = Co, Cr, and Fe) together with characterization studies, have been performed to investigate the effect of the type and oxidation state of the metal in the M(CN)₆ unit on the catalytic performances of PBAs. The effect of hexacyanometal groups on the electronic properties of electroactive cobalt sites has further been examined through electronic structure calculations employing density functional theory (DFT).

A similarity of Tafel slopes indicates similar OER mechanisms. According to chronoamperometric measurements, onset overpotentials of 283, 303, 323, and 343 mV were obtained for [Co⁵⁻Co⁷⁺], [Co⁵⁻Co⁷⁺], [Co⁵⁻Co⁷⁺], and [Co⁵⁻Co⁷⁺], respectively, which are consistent with those of cyclic voltammetric studies (Figure S1). The surface coverage of electroactive Co²⁺ species on an FTO electrode, that is, the surface concentration, was determined by recording CVs at different scan rates (25 – 225 mV s⁻¹) in the range 0.8–1.6 V. Surface concentrations of the derivatives were estimated to be in the range 0–2.
Surface concentration was used to assess turnover frequencies (TOFs) of the PBAs. TOFs at an overpotential of 400 mV were evaluated as 5.0×10^{-2} s^{-1}, 3.0×10^{-3} s^{-1}, 4.4×10^{-3} s^{-1}, and 5.0×10^{-3} s^{-1} for [Co\textsuperscript{II}-Co\textsuperscript{III}], [Co\textsuperscript{III}-Fe\textsuperscript{II}], [Co\textsuperscript{III}-Fe\textsuperscript{III}], and [Co\textsuperscript{III}-Cr\textsuperscript{III}], respectively (Figure S3). Comparison of the TOFs shows that the available Co\textsuperscript{II} sites in [Co\textsuperscript{III}-Co\textsuperscript{III}] exhibit the highest catalytic activity. Chronopotentiometry (CP) was performed to determine the overpotential required to obtain a current density of 1 mA cm\textsuperscript{-2} during a 2 h experiment. The overpotential for [Co\textsuperscript{III}-Co\textsuperscript{III}] slightly decreased at first and then maintained a constant level, whereas those for the other PBAs gradually increased until stabilization. The overpotentials observed at 1 mA cm\textsuperscript{-2} are slightly higher than those extracted from the Tafel slopes due to the formation of O\textsubscript{2} bubbles on the electrode surface during the measurement. CP studies showed that [Co\textsuperscript{III}-Co\textsuperscript{III}] exhibited the lowest overpotential, and η\textsubscript{1mA} was determined as 531, 578, 661, and 692 mV for [Co\textsuperscript{III}-Co\textsuperscript{III}], [Co\textsuperscript{III}-Cr\textsuperscript{III}], [Co\textsuperscript{III}-Fe\textsuperscript{III}], and [Co\textsuperscript{III}-Fe\textsuperscript{II}], respectively (Figure 3, Table 1).

![Figure 3. Chronopotentiometry measurements of PB derivatives at 1 mA cm\textsuperscript{-2} in a 50 mM KPi buffer at pH 7.0.](image.png)

### Characterization studies

All samples were isostructural with the Prussian Blue crystal structure, adopting a face-centered cubic (fcc) form, with space group Fm\textit{3}m, as confirmed by powder XRD studies. The characteristic 2θ peaks for Prussian Blue were observed for all of the materials (Figure S6), and the lattice parameter was determined to be around 10 Å for each derivative (Table S1). XRD analysis in grazing incidence mode was also performed on the catalysts deposited on FTO before (pristine) and after (post-catalytic) the electrocatalytic studies, to investigate their structural stability during electrocatalysis. No additional peaks were observed in the XRD patterns of the post-catalytic samples, and the peaks corresponding to the Prussian Blue-type structure remained, confirming the stability of the catalysts (Figure 4).

The atomic ratio of metals in each compound was determined by EDX analysis (Table S2). The following molecular formulae were obtained, based on stoichiometric ratio of metals: K\textsubscript{0.76}Co\textsubscript{0.24}[Co(CN)\textsubscript{6}]\textsubscript{2+}, K\textsubscript{0.82}Co\textsubscript{0.18}[Cr(CN)\textsubscript{6}]\textsubscript{2+}, K\textsubscript{0.86}Co\textsubscript{0.14}[Fe\textsuperscript{II}(CN)\textsubscript{6}]\textsubscript{2+}, and K\textsubscript{0.91}Co\textsubscript{0.09}[Fe\textsuperscript{II}(CN)\textsubscript{6}]\textsubscript{2+} for [Co\textsuperscript{III}-Co\textsuperscript{III}], [Co\textsuperscript{III}-Cr\textsuperscript{III}], [Co\textsuperscript{III}-Fe\textsuperscript{III}], and [Co\textsuperscript{III}-Fe\textsuperscript{II}], respectively. Each compound has a similar potassium content in the range 0.6–0.8, which results in an average of about 4.5 CN groups per Co\textsuperscript{III} site. The coordination spheres of the Co\textsuperscript{III} sites are completed by water molecules, which play an active role in water oxidation (Figure S7).

Infrared studies showed that the PBAs exhibited the characteristic bands associated with Prussian Blue-type systems: a}

<table>
<thead>
<tr>
<th>Compound</th>
<th>Co\textsuperscript{II}</th>
<th>v(CN) \ [cm\textsuperscript{-1}]</th>
<th>TOF \ [(mA = 400 mV)]</th>
<th>Surface concentration \ [mmol cm\textsuperscript{-2}]</th>
<th>η\textsubscript{1mA} from Tafel plot \ [mV]</th>
<th>Tafel Slope \ [mV dec\textsuperscript{-1}]</th>
<th>η\textsubscript{1mA} from CP \ [mV]</th>
<th>η\textsubscript{max} \ [CV]</th>
</tr>
</thead>
<tbody>
<tr>
<td>[Co\textsuperscript{III}-Co\textsuperscript{III}]</td>
<td>1.010</td>
<td>2176</td>
<td>5.0×10\textsuperscript{-2}</td>
<td>4.11</td>
<td>531</td>
<td>99</td>
<td>565</td>
<td>283</td>
</tr>
<tr>
<td>[Co\textsuperscript{III}-Cr\textsuperscript{III}]</td>
<td>1.084</td>
<td>2173</td>
<td>5.0×10\textsuperscript{-3}</td>
<td>3.90</td>
<td>578</td>
<td>96</td>
<td>598</td>
<td>303</td>
</tr>
<tr>
<td>[Co\textsuperscript{III}-Fe\textsuperscript{III}]</td>
<td>1.084</td>
<td>2120</td>
<td>4.4×10\textsuperscript{-3}</td>
<td>5.48</td>
<td>661</td>
<td>127</td>
<td>717</td>
<td>323</td>
</tr>
<tr>
<td>[Co\textsuperscript{III}-Fe\textsuperscript{II}]</td>
<td>0.995</td>
<td>2072</td>
<td>3.0×10\textsuperscript{-3}</td>
<td>2.00</td>
<td>692</td>
<td>121</td>
<td>1079</td>
<td>343</td>
</tr>
</tbody>
</table>
sharp band at around 1610 cm$^{-1}$ and a broad feature at 3200–3500 cm$^{-1}$, which correspond to HOH bending and OH stretching, respectively; b) a sharp peak at around 490–590 cm$^{-1}$ due to M–C stretching; and c) a sharp peak at around 2120–2180 cm$^{-1}$ attributable to CN stretching (Table S3). PBAs exhibit higher CN stretching frequencies compared to their hexacyanometal precursors, which confirms the binding of nitrogen atoms of cyanide to Co$^3\text{II}$ sites$^{[28,40]}$ (Figure S8). Infrared analysis was also performed on the post-catalytic samples. The close similarity between the cyanide stretching IR bands of the pristine and post-catalytic samples suggests that the M-CN-Co$^3\text{II}$ type coordination mode of the catalysts is preserved during electrolysis (Figure S9). A slight shift to higher frequencies in the case of post-catalytic [Co$^3\text{II}$Fe$^{\text{III}}$], which was also observed in previous studies, can be attributed to partial oxidation of iron ions from $\pm 2$ to $\pm 3$ during the electrocatalysis.

XPS studies also confirmed the remarkable stability of the PBA electrocatalysts. In order to investigate the oxidation state of electroactive Co$^3\text{II}$ sites in the pristine and post-catalytic electrodes, the Co$^2\text{p}$ signal was examined in the binding energy region 810–775 eV. In previous studies, the binding energies of Co$^{2\text{p}_1/2}$ and Co$^{2\text{p}_3/2}$ signals for Co$^3\text{II}$ salts have been reported as 782.28 and 798.38 eV, respectively. For the pristine samples, the Co$^{2\text{p}_1/2}$ and Co$^{2\text{p}_3/2}$ signals were observed in the same range. The similarity between the binding energies of the Co$^2\text{p}$ signals obtained for pristine PBAs and previously reported Co$^3\text{II}$ salts suggests that the oxidation state of electroactive Co atoms is $\pm 2$ (Figure 5). No significant changes in the Co$^{2\text{p}_1/2}$ and Co$^{2\text{p}_3/2}$ signals were observed in the post-catalytic samples, indicating the stability of the Co$^3\text{II}$ sites.

In addition to Co$^2\text{p}$, the O$1\text{s}$ signals were also examined for both the pristine and post-catalytic samples (Figure S10). An O1$\text{s}$ signal with a binding energy higher than 530 eV indicated an absence of any cobalt oxide species before and after electrochemical experiments, even for [Co$^3\text{II}$Cr$^{\text{III}}$]. The observed values are displayed in Table S4. A moderate broadening of the O1$\text{s}$ signal is evident in the post-catalytic samples, indicating a partial and reversible oxidation of electroactive Co$^3\text{II}$ sites.

**Mechanism of catalytic water oxidation**

The CN stretching vibration may be considered as a fingerprint for cyanide-based coordination compounds. A comparison of shifts in the cyanide stretch can be used not only to confirm the bridging mode of the cyanide group, but also to evaluate the oxidation states, and hence the electron densities, of the metal ions. Considering that the cyanide stretching vibration shifts to higher frequencies as the oxidation state of the metal increases, a direct correlation can be established between the shift of the cyanide stretch and the electron deficiency of Co$^3\text{II}$ centers. Comparison of the cyanide stretches implies that the electron densities at the Co$^3\text{II}$ sites in our Prussian blue analogues can be ordered as: [Co$^3\text{II}$Co$^3\text{II}$] $\approx$ [Co$^3\text{II}$Cr$^{\text{III}}$] $<$ [Co$^3\text{II}$Fe$^{\text{III}}$] $<$ Co$^3\text{II}$ (Figure 6). This result is also in good agreement with the binding energies of the Co$^2\text{p}$ orbitals obtained by XPS studies. The ordering of the Co 2$p_{3/2}$ peaks is [Co$^3\text{II}$Co$^3\text{II}$] $>$ [Co$^3\text{II}$- Cr$^{\text{III}}$] $>$ [Co$^3\text{II}$Fe$^{\text{III}}$] $>$ [Co$^3\text{II}$Fe$^{\text{III}}$], implying that the Co$^3\text{II}$ sites in the [Co$^3\text{II}$Co$^3\text{II}$] analogue have the lowest electron densities in the series. The evaluation of electron densities can provide insight into the rate-determining step (r.d.s.) in water oxidation catalysis. Two steps have generally been reported as competing as the r.d.s. in water oxidation process: i) Co$^3\text{II}$–OH/OCo$^3\text{III}$–O (oxo) or Co$^3\text{II}$–OH/OCo$^3\text{III}$–O (oxyl) as the oxidation step and ii) the nucleophilic attack of water at the electrophilic oxygen atom of oxo/ oxyl species, resulting in O–O bond formation. An increase in the electron density at the Co$^3\text{II}$ site facilitates the former step,
while decreasing the electrophilic nature of the oxo intermediate and thus impeding the latter. The above discussion on the electron densities of Co\(^{3+}\) sites in PBAs and their electrocatalytic performances clearly shows that [Co\(^{4+}\)-Co\(^{3+}\)] stands out as the most efficient catalyst among the studied PBAs, which has Co\(^{3+}\) sites with the lowest electron density. This correlation implies that nucleophilic attack of water on the oxo/oxyl intermediate is the r.d.s. of the water oxidation process for PBAs. It should be noted that the electronic properties of the catalysts will differ when a potential is applied. Catalytically active cobalt ions will be in their higher oxidation states, particularly when the applied potential is above 1 V versus NHE. Nevertheless, the difference in the electron densities of the cobalt ions should be preserved, given that the structural integrity of the cyanide framework is preserved and that the metal ion in the M(CN)\(_{6}\) building block is not oxidized. While this assumption may be valid for hexacyanometal groups that contain metal ions in their 3+ oxidation state, the Fe\(^{2+}\) ion in the [Fe(CN)]\(_6\)\(^{3-}\) group would be expected to be oxidized when a potential above 1 V is applied.\(^{[41]}\) The oxidation of all Fe\(^{2+}\) ions is, however, a kinetically demanding process, since it requires more potassium ions to be transported from the framework to the electrolyte to maintain charge neutrality and, more importantly, there are insufficient potassium ions to produce a fully oxidized [Co\(^{6+}\)-Fe\(^{3+}\)] system. Therefore, the catalytically active species in [Co\(^{4+}\)-Fe\(^{3+}\)] contains a mixture of Fe ions with oxidation states of 2+ and 3+. The difference in the curvatures of the bands assigned to the Fe\(^{2+}/3+\) and Co\(^{2+}/3+\) redox processes for [Co\(^{4+}\)-Fe\(^{3+}\)] and [Co\(^{4+}\)-Fe\(^{2+}\)] also indicates different kinetics for these two analogues (Figure S11). The lower surface concentration and turnover frequency obtained for [Co\(^{4+}\)-Fe\(^{3+}\)] may thus be attributed to differences in the kinetics of its electron transfer and its electronic properties.

A further analysis of the mechanism was made based on the Pourbaix diagram (Figure 7), which was obtained by recording CVs for [Co\(^{8+}\)-Co\(^{3+}\)] at different pH values (Figure S12). The diagram shows that Co\(^{3+}/Co^{2+}\) redox process is pH-dependent in the range pH 4–10 with a slope of 64 mV log [H\(^+\)]\(^{-1}\), which refers to a 1H\(^++\)1e\(^-\) process.

Interestingly, the half-potential for the second redox step is preserved, regardless of pH (< 11), which indicates that a hydroxyl group is coordinated to the catalytically active Co\(^{6+}\)-oxo/Co\(^{5+}\)-oxyl intermediate under neutral conditions. An [N,Co\(^{6+}\)-OH] intermediate could be deprotonated in a subsequent step to form a cobalt oxo/oxyl complex, and then undergo nucleophilic attack of water to form a peroxy intermediate, which is one of the essential steps for O–O bond formation. A slightly different mechanism to that commonly accepted for oxides is thus proposed for PBAs. The presence of additional peaks in the CVs obtained at above pH 11 suggests that water oxidation proceeds with a different mechanism under basic conditions.

Electronic structure calculations

In order to gain insight into the different performances of the PB analogues studied in this work, electronic structure calculations were performed with DFT\(^{[37,38]}\) (see the SI for details). It is critical to understand the reason for the rate enhancement along the Fe\(^{6}\), Fe\(^{4}\), Cr\(^{6}\), and Co\(^{3+}\) cationic series used as the second metal separated from the catalytically active Co site by a cyanide bridge.

Oxidation of water would require the proton coupled electron transfer (PCET) steps to afford the formal Co\(^{6+}\)-oxo/Co\(^{4+}\)-O\(^-\) moiety from the substrate-bound aqua center with a +2 formal charge, that is, Co\(^{2+}\)(OH)\(^-\)→Co\(^{3+}\)(OH)→Co\(^{4+}\)(O)/Co\(^{4+}\)(O). Once Co\(^{4+}\)(O) is accessed, it is attacked by water to afford the O–O bond. This picture is consistent with the existing mechanistic data in the literature.\(^{[39,42]}\) Therefore, the structural and electronic properties of the Co\(^{5+}\)(O)/Co\(^{6+}\)(O) center are the main focus of our quantum chemical calculations. It is important to note that there are no restrictions on the distribution of electrons in our calculations, and thus formal assignments

**Figure 6.** FTIR spectra of PB derivatives showing cyanide stretches.

**Figure 7.** Pourbaix diagram of [Co\(^{4+}\)-Co\(^{3+}\)] in KPi buffer over the range pH 2–13. Cyclic voltammograms recorded at these pH values are shown in Figure S10.
of Co\textsuperscript{IV}(O) or Co\textsuperscript{III}(O) for the Co center are less comprehensive levels of describing the Co–O bond compared to the completely delocalized (canonical) orbital picture given by quantum chemical calculations.

DFT calculations suggest that the catalytically active Co–O site is a local quartet and hence hosts three unpaired electrons. Depending on the nature of the neighboring metal in a strong field environment, either one or zero electrons contribute to the total spin when a bimetallic model is considered, as given in Table S5. The local quartet spin arrangement of the Co\textsuperscript{IV} center is also verified by spin density analysis, as given in Table 2. Interestingly, the reactive Co\textsuperscript{IV} center has a high degree of radical character distributed over the Co–O bond. Such electronic structure fingerprints of high-valent Co moieties were recently shown to be related to reactivity,\textsuperscript{[43]} where an apparent Co\textsuperscript{IV}=oxo species having a mixed electronic structure of the oxyl/oxo type was reported. In that work, one of us (Y.D.) showed that, similarly to the case reported here, the oxo-alkaline\textsuperscript{[48]} was indeed not broken, and the local quartet spin arrangement, delocalized along the Co–O bond, showed substantial radical character on oxygen. Overall, a more appropriate electronic structure assignment for the in situ generated, catalytically competent, O–O bond-forming species is a mixed Co-oxo/oxyl.

CN stretching frequencies (\nu(CN)) and molecular orbitals were investigated to elucidate the molecular basis for the catalytic activity (Table 2).

The calculated trend in \nu(CN) is in good agreement with the experimental results, and shows the flow of electron density from the Co site. More importantly, the critical O–O bond-forming step can be readily understood by analyzing the attack of water on the Co–O center. The oxygen lone pairs borne by water are seeking vacant orbitals on the Co–O center, for which the best candidate is the LUMO. As shown in Table S5, the \sigma\textsuperscript{*} MO generated from Co\textsubscript{d} and O\textsubscript{p} contributions is obtained at lower energies (E\textsubscript{LUMO} in Table 2) through the Fe\textsuperscript{II}, Fe\textsuperscript{III}, Cr\textsuperscript{II}, Cr\textsuperscript{III}, and Co\textsuperscript{IV} sites. Thus, the electron affinity of the Co\textsuperscript{IV}–O center is increased; the attack of water becomes more facile, and this accounts for the lower overpotentials measured in our electrochemical experiments. Our quantum chemical calculations thus show that the reactive Co-oxo/oxyl center possesses a local quartet spin arrangement. Two of the three quartet spin electrons are distributed over Co and one over oxygen; however, all three electrons can be better described as sharing Co\textsubscript{d} and O\textsubscript{p} orbitals through the Co–O bond. The reactivity correlates with attaining the LUMO, to be attacked by incoming water, at lower energies, as summarized in Figure 8. Note that this truncated quantum chemical model may not capture all of the structural features of the PB surface, but it is a good compromise between accuracy and cost. Moreover, with the assistance of the experimental data, the electronic structure of the active species could be assigned as Co\textsuperscript{IV}(O)/Co\textsuperscript{III}(O), and hence the model chemistry is useful.

![Figure 8. O–O bond formation at the Co\textsuperscript{IV}–O center.](image)

| Table 2. Structural and electronic properties of the M/Co PBAs computed at the UM06L/cc-pVTZ level of theory.\textsuperscript{[44]} |
|-----------------|-------------|-------------|-------------|-------------|
|                  | R Co–O [Å]  | \nu CN [cm\textsuperscript{-1}] | \rho [O]    | \rho [Co\textsuperscript{IV}] | E\textsubscript{LUMO} [eV] |
| [Co\textsuperscript{IV}–Co\textsuperscript{II}] | 1.652       | 2195        | 1.22        | 1.63        | -14.34       |
| [Co\textsuperscript{IV}–Co\textsuperscript{III}] | 1.654       | 2158        | 1.24        | 1.62        | -13.85       |
| [Co\textsuperscript{IV}–Fe\textsuperscript{II}] | 1.653       | 2160        | 1.23        | 1.62        | -12.91       |
| [Co\textsuperscript{IV}–Fe\textsuperscript{III}] | 1.684       | 2114        | 1.17        | 2.10        | -11.06       |

[a] Surfaces were generated at 0.05 a.u.
Conclusions

Previous electrochemical studies on Prussian Blue analogues have shown that having a cobalt site coordinated to nitrogen atoms of a cyanide bridging group is essential to obtain efficient PBA electrolysis catalysts for water oxidation. Here, members of a series of cobalt hexacyanometalates with the general formula \( \text{K}_n\text{Co}_{x}\text{M} \) \( \text{[CN]}_y \) \( \text{[M} \) \( \text{CN]}_z \) \( \text{] } \) (\( \text{M} = \text{Co}^{II}, \text{Cr}^{III}, \text{and} \text{Fe}^{III} \)) have been prepared to investigate the effect of the hexacyanometal groups on the electrocatalytic activity of CHCMs. Tafel analysis and chronoamperometry experiments have revealed that \( \text{[Co}^{III}-\text{Co}^{II}] \) serves as the most efficient electrocatalyst for water oxidation among the studied CHCMs. Infrared and XPS studies have indicated that it has the Co\(^{II}\) center with the lowest electron density, which has also been confirmed by DFT studies.

Overall, experimental and computational studies have led to the following conclusions:

i) The electron density of Co\(^{II}\) is a decisive electronic criterion for achieving efficient water oxidation electrocatalysis, and this parameter may be tuned by changing the type of hexacyanometal group.

ii) The electron density of Co\(^{III}\) can be reduced by increasing the oxidation state of the metal ion of the \( \text{[M(CN)}_y \text{]}_z \) group.

iii) Electrophilicity of the Co\(^{III}\)-O center can be probed by molecular orbital analysis, and this might be a useful tool for the realization of new PBAs bearing reactive Co species as potent WOCs.

iv) Nucleophilic attack of water on the cobalt-oxo intermediate should be the rate-determining step for water oxidation catalysis with PBAs.

v) Theoretical calculations show that the O\( \rightarrow \)O bond-forming species has mixed Co-oxo/oxyl character.

In conclusion, the following mechanism may be proposed, based on the experimental and computational studies (Figure 9). This study has shown that electronic properties, and hence the electrocatalytic activity of the catalytically active cobalt site, can be easily tuned through the versatile chemistry of PBAs, and neighboring metal ions should also be considered as an important parameter to evaluate the catalytic activities of water oxidation electrocatalysts. Detailed electronic structure calculations, employing multi-reference techniques, on various Co–O systems are underway aimed at corroborating the nature of the Co\(^{III}\)-O bonding and reactivity.

Experimental Section

Chemicals and solutions

Potassium hexacyanocobaltate \( \text{K}_n\text{Co}_{x}\text{M} \) \( \text{[CN]}_y \) \( \text{[M} \) \( \text{CN]}_z \) \( \text{] } \) \( \text{] } \) \( \text{] } \) \( (> 97.0 \% ) \), cobalt(II) chloride hexahydrate \( \text{CoCl}_2 \cdot 6\text{H}_2\text{O} \) (98.0%), potassium hexacyanochromate \( \text{K}_2\text{[CrCN]}_6 \) (99.99%), potassium hexacyanoferrate \( \text{K}_4\text{[FeCN]}_6 \) (> 97.0%), and potassium hexacyanoferrate trihydrate \( \text{K}_3\text{[Fe(CN)}_6 \text{]}_3 \cdot 3\text{H}_2\text{O} \) (98.5–102%) were all obtained from Sigma–Aldrich. All the solutions were prepared with Millipore Milli-Q deionized water with a resistivity of 18.2 MΩ cm.
Physical measurements

XRD patterns were measured by means of a PanAnalytical X’Pert-Pro multipurpose X-ray diffractometer (MDP) employing CuKα radiation (λ = 1.5418 Å). GI-XRD patterns were recorded by using a PanAnalytical X’Pert 3 MRD material research diffractometer (MRD) with CuKα, X-ray radiation (λ = 1.5418 Å) at an incident angle of 0.58°. FTIR spectra were acquired with a Bruker Alpha Platinum-ATR spectrometer over the wavenumber range 4000–400 cm⁻¹. An FEI-Quanta 200 FEG ESEM was used for imaging and EDAX analysis, operated at 5 kV beam voltage for imaging and 30 kV for EDAX. XPS analysis was performed on a Thermo Scientific K-Alpha X-ray photoelectron spectrometer system with an AlKα microfocused monochromator source operated at 400-mm spot size and by = 1486.6 eV accompanied by a flow gun, 200 eV for survey scans and 30 eV for individual scans. Origin Pro 8.5 software was used to plot and analyze the results.

Acknowledgements

The authors thank the Science and Technology Council of Turkey, TUBITAK (Project No. 215Z2249) for financial support. E.U. thanks TUBITAK for support (Project No. 1929B011500059). Y.D. acknowledges ECOSTBio (CM 1305) for support and thanks the M.N. Parlar Foundation, BAGEP, and TÜBA-GEİBİ for young investigator awards. TUBITAK TRGRID infrastructure is gratefully acknowledged for HPC resources. We also thank Prof. Burak Ülgüt for helpful discussions on electrochemistry.

Conflict of interest

The authors declare no conflict of interest.

Keywords: cyanides · density functional calculations · electrocatalysis · Prussian Blue · water oxidation