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Ruthenium–rhenium and ruthenium–palladium supramolecular photocatalysts for photoelectrocatalytic CO₂ and H⁺ reduction†‡

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Photoelectrocatalysis offers the opportunity to close the carbon loop and convert captured CO₂ back into useful fuels and feedstocks, mitigating against anthropogenic climate change. However, since CO₂ is inherently stable and sunlight is a diffuse and intermittent energy source, there are considerable scientific challenges to overcome. In this paper we present the integration of two new metal–organic photocatalysts into photocathodes for the reduction of CO₂ using ambient light. The two molecular dyads contained a rhenium carbonyl or palladium-based catalytic centre bridged to a ruthenium bipyridyl photosensitizer functionalised with carboxylic acid groups to enable adsorption onto the surface of mesoporous NiO cathodes. The photocathodes were evaluated for photoelectrochemical reduction of CO₂ to CO or H⁺ to H₂ and the performances were compared directly with a control compound lacking the catalytic site. A suite of electrochemical, UV-visible steady-state/time-resolved spectroscopy, X-ray photoelectron spectroscopy and gas chromatography measurements were employed to gain kinetic and mechanistic insight to primary electron transfer processes and relate the structure to the photoelectrocatalytic performance under various conditions in aqueous media. A change in behaviour when the photocatalysts were immobilized on NiO was observed. Importantly, the transfer of electron density towards the Re–CO catalytic centre was observed, using time resolved infrared spectroscopy, only when the photocatalyst was immobilized on NiO and not in MeCN solution. We observed that photocurrent and gaseous photoproduct yields are limited by a relatively low yield of the required charge-separated state across the NiO|Photocatalyst interface. Nonetheless, the high faradaic efficiency (94%) and selectivity (99%) of the Re system towards CO evolution are very promising.

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Introduction

For many decades, chemists have been trying to create molecular systems that mimic photosynthesis in order to capture the

energy available in sunlight in chemical bonds. There are now an abundance of chromophores and catalysts,^{1–3} and various colloidal semiconductors that can drive the water splitting reaction or reduce CO₂ to produce energy dense fuels such as H₂, CO, HCO₂H and hydrocarbons.^{4–8} These systems offer opportunities to tune the products with high selectivity and atom efficiency. Pioneering work into artificial photosynthesis and molecular photophysics from the 1970s and 1980s provided a wealth of information on the photophysics of such systems.^{9–10} The challenges of light-driven water-splitting and CO₂ reduction using molecular systems in solution are well documented.^{11–14} These include the difficulty of driving multi-electron chemistry when each photon absorbed only provides a single pair of an electron and hole. Storing charges on the appropriate timescale to drive the chemistry, while avoiding decomposition pathways, is a major focus of current research. As more elaborate chromophores have been developed to resolve these challenges, detailed mechanistic insight has been gained from investigations into primary photo-induced electron transfer and energy transfer. These processes are highly sensitive to minor

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structural changes and modification of the local environment, adding another layer of complexity.¹⁵ At this point, for systems in solution, the thermodynamic and kinetic requirements for developing effective assemblies to separate redox pairs from light harvesting arrays are well-understood. Some headway has also been made towards developing catalysts from earth-abundant metals.^{16–20} A key issue still remains for molecular and colloidal artificial photosynthetic systems. That is, that the charge required to drive the oxidation or reduction reactions, which are usually performed with two separate photocatalysts, is usually provided by a sacrificial reagent. This requirement limits the sustainability of the process as well as generating reactive waste products.

An alternative approach is to use a photoelectrochemical system where electrons liberated by the water oxidation reaction at the photoanode can then be consumed at the fuel forming reaction at the photocathode.^{21–26} Such systems perform with promising light-to-fuel efficiency but in aqueous solutions, CO₂ reduction competes with the kinetically more favourable hydrogen evolution reaction and selectivity is difficult to control.^{27–30} A challenge is finding a p-type semiconductor that is stable to corrosion and has an appropriate electronic band alignment with the desired electrochemical reactions to deliver a high photovoltage and photocurrent density.³¹ Photocathodes have been reported based on metal phosphides, chalcogenides, oxides, III–V, p-Si and metal free semiconductors.^{7,23,28,32} For example, Kang *et al.* demonstrated that photocathodes based on CuFeO₂ and CuO binary films selectively produced formate from aqueous CO₂ at ~1% efficiency over 24 h, under AM 1.5G; 100 mW cm⁻² simulated sunlight. However, the photocathode gradually degraded over 17 days.³¹ Zhang *et al.* reported a Cu₂O-SnO_x nanowire-based photocathode that produced syngas (CO and H₂ with a ratio that could be adjusted with composition and applied bias) with a 11.61 mA cm⁻² at -0.55 V *versus* RHE, and a total faradaic efficiency of 90.32% at -0.35 V *versus* RHE.³³ p-Si decorated with catalytic metal nanoparticles (*e.g.* Cu, Ag, Au) has been used to convert CO₂ to CO with faradaic efficiencies >80%, but parasitic light absorption can limit the overall solar to fuel efficiency. Gong *et al.* reported an amorphous silicon photocathode, protected by a thin layer of TiO₂ and decorated with Au of variable thickness that enabled photocurrent densities of up to 6 mA m⁻² and CO/H₂ ratios that could be tuned between 1:2 and 1:3.1 by changing the applied bias between -0.1 V_{RHE} to 0.4 V_{RHE}.³⁴ Lewis *et al.* optimized a microstructured p-Si photocathode decorated with Cu and attained a photocurrent density of 30 mA cm⁻² in 0.1 M KHCO₃(aq) at -0.62 V *vs.* RHE under AM 1.5 simulated solar illumination. CO, CH₄, and C₂H₄, were detected but the major product was hydrogen, with only 10% faradaic efficiency for carbon products.²⁸

A higher selectivity and lower parasitic absorption can be reached with molecular catalysts immobilized on semiconductor photocathodes.⁷ For example, Kumar *et al.* reported a p-type silicon photocathode coated with [Re(bipy⁴Bu)(CO)₃Cl] that reduced CO₂ to CO under non-aqueous conditions with a faradaic efficiency of 97%.³⁵ Re(bpy)(CO)₃Cl was immobilised *via* phosphonate linkers on the surface of a photocathode

assembled from a nanostructured TiO scaffold deposited on TiO₂-protected Cu₂O photocathodes.³⁶ Under non-aqueous conditions, the photocathode produced an initial photocurrent density of 2.5 mA cm⁻² and faradaic yields for CO between 80 and 95%. The catalyst molecules exceed 70 turnovers during a 1.5 h test at -2.05 V *vs.* Fc, but there was a loss of activity. There are few reports of such high selectivity for CO under aqueous conditions, however.

In recent years, the wealth of knowledge gained from decades of research into dye-sensitized solar cells (DSSC) has been applied to optimizing photoelectrochemical (PEC) systems for both water-splitting and CO₂ reduction.^{37–41} Photoelectrochemical systems based on dye-sensitized electrodes (DS-PEC)^{42–44} can be used to generate solar fuels, circumventing one of the main obstacles encountered by molecular photocatalytic systems in solution, namely the frequent need for a sacrificial electron and proton source other than water.^{45,46} Instead, the molecular chromophore and catalyst are immobilized on the surface of a transparent electrode and excitation of the chromophore drives a cascade of electron transfer reactions between the electrode and the catalyst.^{21,22,47,48} TiO₂ is the optimum material for the photoanode. While a number of mesoporous materials were used for the photocathode, NiO is the most common choice due to its good transparency, p-type character, and stability under the conditions of the experiments.^{49–57} NiO is a well-understood p-type material with a valence band edge that lies sufficiently positive in energy to donate an electron to an excited molecular photosensitizer. It is typically chosen as the cathode material in photoelectrochemical systems due to its relatively good stability under a moderate pH range used and its optical band gap (3.6 eV) is sufficiently large that it does not compete for light absorption with the dyes and photocatalysts.^{58–61}

While these systems are fairly straightforward to assemble, and they maintain the high atom efficiency of the catalyst while benefitting from the robust semiconductor scaffold, the mechanism is complicated and optimization of the system is not so simple. The dynamics of the light-driven processes determined *via* studies of analogous chromophores in dilute solution do not directly transfer to the equivalent DS-PEC device.^{62,63} Electron transfer from a dye/catalyst anchored to a semiconductor surface by the traditional strategy of an acid or phosphonate anchoring group leads to multiple competing recombination pathways, many similar to those seen with DSSCs.^{57,64–67} It is also now known that the chromophore's anchoring group itself can have a significant impact on charge-recombination.⁶⁸ Recent work has begun to recognise the challenges in engineering suitable catalysts and sensitizers for DS-PEC, which involves a combination of device engineering, optimizing reaction conditions and dye/catalyst design.^{48,69–77} A key design feature in DS-PEC systems is the spatial separation of the catalytic centre from the semiconductor surface to slow down charge-recombination and maximise catalytic turnover. Mastering all these facets simultaneously and scaling up devices presents a major challenge.

CO₂ reduction with DS-PEC has been investigated previously with devices incorporating a Ru(II)-Re(I) molecular





Fig. 1 Molecular structures of the photocatalysts used in the present study. $[\text{Ru}(\text{bpy})(\text{dcb})(\text{bpt})\text{Re}(\text{CO})_3\text{Cl}](\text{PF}_6)[\text{PC1}]$, $[\text{Ru}(\text{bpy})(\text{dcb})(\text{bpt})\text{PdCl}_2](\text{PF}_6)_2[\text{PC2}]$, $[\text{Ru}(\text{bpy})(\text{dcb})(\text{bpt})](\text{PF}_6)_2[\text{PC3}]$, bpy = 2,2'-bipyridine, bpt = 3,5-di(pyridin-2-yl)-triazole, dcb = 2,2'-bipyridine-4,4'-dicarboxylic acid. PC1 is designed for CO_2 reduction, PC2 is designed for H_2 evolution and PC3 is a catalyst-free control.

photocatalyst. Ishitani and co-workers successfully demonstrated such a model system utilizing a phosphonic acid anchor for the RuRe catalyst on NiO where CO_2 was reduced predominantly to CO with a MeCN electrolyte at an applied potential of -1.2 V vs. Ag/AgCl.⁷⁸ To function effectively over 15 hours in an aqueous environment, a polymer layer was added on top of the catalyst sensitized NiO to prevent desorption.^{43,44} Optimization of this system led to remarkable stability over 100 hours and the total TON of the photocatalytic CO_2 reduction exceeded 1200 ($\text{TON}_{\text{CO}} = 576$, $\text{TON}_{\text{HCOOH}} = 695$, $\text{TON}_{\text{H}_2} = 172$).⁷⁷

Previously we have shown that supramolecular photocatalysts containing ruthenium bipyridyl photosensitizers linked through a conjugated bridging ligand to a rhenium carbonyl or Pd catalytic centre drive photoelectrocatalytic reduction of CO_2 to CO or H^+ to H_2 when integrated in NiO-based photocathodes. The Ru component enables the photocatalyst to absorb further into the red (towards 600 nm or beyond) than the Re complex alone (~ 450 nm).⁷⁹ A second advantage is that the absorption coefficient of the bimetallic system is higher than the Ru system alone, which means the NiO film can be quite thin to overcome the slow hole-transport.⁸⁰ Co-immobilization of the photosensitizer and catalyst would lead to competition for binding sites at the surface and so the thickness of the NiO would need to be even thicker to immobilize the equivalent number of chromophores as the supramolecular system. The third advantage of the structure is to improve the photophysics. The supramolecular structure is intended to provide conjugation from the NiO through to the catalytic centre so that excitation of the photocatalyst leads directly to reduction. In other systems the catalyst has been electronically decoupled from the photosensitizer and several sequential electron transfer steps are required to reduce the catalyst. These processes compete with charge-recombination, which is prominent in NiO photoelectrodes, and can limit the overall efficiency.⁶³

In our previous work, the photocatalysts were adsorbed on the NiO surface through ester groups, but only trace amounts of CO were produced and it was noted that carboxylic acids may be preferable for anchoring to metal oxide surfaces.^{63,80,81} In this

paper, we consider two photoelectrocatalysts, one for CO_2 reduction (PC1), one for H_2 evolution (PC2), with carboxylic acid groups for NiO photocathodes. PC3 was included as a catalyst-free compound to benchmark the activity (Fig. 1). We also tested a Pt analogue of PC2, but this gave negligible photocurrent, so we have not included it here. This paper describes the optimization of reaction conditions, analysis of the electrochemical and photophysical properties in addition to chromatography measurements of gaseous products. Some of the major bottlenecks that we found in designing DS-PEC systems where the chromophore and catalyst are integrated in one molecule are discussed.

Results and discussion

Optical, electronic and electrocatalytic properties of the photocatalysts

The photocatalysts were synthesized according to the procedures provided in the ESI† and were characterized by NMR, mass spectrometry and FTIR spectroscopy. Steady-state absorption and emission properties of the photocatalysts PC1–PC3 in solution are summarized in Fig. 2 and Table 1. The FTIR spectra of the ruthenium complexes in the 1650–2100 cm^{-1} region are shown in Fig. S28.† All complexes show a signal at 1735 cm^{-1} which is assigned to the CO stretch in the acid moiety on the dcb peripheral ligand. The IR spectrum for $[\text{Ru}(\text{dcb})(\text{bpy})(\text{bpt})\text{Re}(\text{CO})_3\text{Cl}](\text{PF}_6)$ contains stretching vibration at 1735 (COOH), 1896, 1908, 1930, 2020 and 2036 cm^{-1} . Based on the relative shapes of these stretches we assign these signals to a mixture of two different fac-Re(CO)₃Cl complexes. Similar observations have been observed by other groups for Ru–Re assemblies, and attributed to isomers.^{82–84}

The chromophores developed in this study have absorption and emission properties analogous to other molecular systems developed around a ruthenium trisbipyridyl sensitizer, with metal-to-ligand charge transfer bands observed between 400 and 500 nm.^{63,85} As can be observed from Fig. S29,† the Re(Hbpt)(CO)₃Cl catalyst has very limited absorbance in the visible region of the solar spectrum. The RuRe supramolecular





Fig. 2 Left panel: steady-state absorption spectra of the photocatalysts in acetonitrile. Right panel: absorption spectra of the same systems when adhered to NiO electrodes following dye-bath deposition.

Table 1 Summary of basic optical parameters for the photocatalysts used in this study. The molar absorption coefficient ϵ was determined at the absorption maximum, in acetonitrile

Compound	$\lambda_{\text{max}}^{\text{abs}}$ (MeCN)/nm	$\lambda_{\text{max}}^{\text{fluor}}$ (MeCN)/nm	$\lambda_{\text{max}}^{\text{abs}}$ (NiO)/nm	E_{00}/eV (MeCN)	$\epsilon/\text{M}^{-1} \text{cm}^{-1}$	Dye loading/mol cm^{-2}
PC1	470	671	470	2.05	9400	4.33×10^{-8}
PC2	474	665	467	1.98	12 300	2.50×10^{-8}
PC3	490	720	480	1.99	2700	3.70×10^{-8}

assembly **PC1**, however, has a significantly extended absorbance range, which should enable this assembly to deliver improved photoelectrocatalytic activity compared to the $\text{Re}(\text{Hbpt})(\text{CO})_3\text{Cl}$ complex by itself.

The dyes were adsorbed on the NiO surface by immersing the electrodes in an acetonitrile solution containing 3 mM photocatalyst overnight. A 10–20 nm blue shift in the absorption spectra was observed for each photocatalyst, which is consistent with the behaviour of the ester equivalents.⁶³ The loading of the photocatalyst on the freshly prepared surface of the NiO was quantified by desorbing the photocatalyst using 1 M NaOH solution and recording the UV-visible absorption spectrum (Table 1). The loading was approximately 25–50 nmol cm^{-2} (highest for **PC1**), which is much higher than the ester analogues (4–7 nmol cm^{-2}).^{27,28,61,86} This means that the light harvesting efficiency is sufficient with a thin NiO film (1.5 μm). Weak fluorescence was readily observed in solution (see ESI[†]), but this was very hard to detect reliably from the NiO surface. Therefore, fluorescence quenching could not be used effectively to characterize charge-transfer at the NiO|PC interface. Instead, transient absorption spectroscopy was employed (see below).

For charge-transfer to occur from the NiO to the photocatalyst following light absorption, the HOMO of the dye needs to be at more positive potential than the potential of the valence band edge, which is $E_{\text{VB}} = ca. -0.12 \text{ V vs. Fc}$ for NiO in MeCN.⁸⁷ To estimate the ample driving force for charge transfer at the NiO|PC interface, cyclic voltammetry (CV) for **PC1–3** was performed and the data acquired is provided in the ESI.[†] The CVs were not as straightforward as for the ester moieties, which show three reversible reductions and a reversible oxidation, due to the non-innocent behaviour of the acid moieties,⁷⁶ as the acids can undergo deprotonation following electrochemical

reduction forming the carboxylates, causing irreversibility of the reduction process.

The CV for **PC3** contains three reduction processes at -1.7 , -2.0 and -2.2 V vs. Fc , which can be assigned to reduction of the three ligands. Based on previous reports, the reduction takes place in the order $\text{dcb} > \text{bpy} > \text{bpt}$.^{80,88,89} The first reduction appears to be irreversible while the second and third reduction have some reversible character. A fourth irreversible reduction at -2.8 V vs. Fc , may be assigned to a second reduction taking place on the bpt bridging ligand, as previously reported by Vos *et al.* for the $\text{Ru}(\text{bpy})_2(\text{bpt})$ complex. A reversible oxidation at *ca.* 0.75 V vs. Fc is assigned to the $\text{Ru}^{\text{II/III}}$ oxidation. A small shouldering irreversible oxidation at 0.44 V was also observed for the Ru-ester analogue ($\text{Ru}(\text{dcb})_2(\text{bpt})$) and can be assigned to residual $\text{Ru}(\text{dcb})(\text{bpy})(\text{Hbpt})$ in which the triazole moiety on the ruthenium complex is still protonated.⁹⁰ No reductive signals were observed for this protonated species as the proton is lost under slight reductive conditions forming the deprotonated species which is consequently observed in the reductive region of the CV.

The CV for **PC2** was more complicated than that of the Ru mononuclear species, **PC3**. The separate reduction processes observed between -1.7 and -2.2 V for **PC3** were less resolved for **PC2** and became fully irreversible. The reversible oxidation for the Ru centre was also shifted anodically to 0.85 V vs. Fc . A second irreversible oxidation was observed at 1.4 V vs. Fc , which relates to oxidation of the Pd centre. As for **PC2**, the CV for the RuRe analogue **PC1** contained additional reductive and oxidative features compared to **PC3**. The reduction processes on the three ligands of the Ru complex also became irreversible and were observed at -1.8 , -2.0 and -2.1 V . Reversibility was also partially lost for the $\text{Ru}^{\text{II/III}}$ oxidation which is observed at 0.9 V



vs. Fc. A second oxidation at 1.1 V is likely to be associated with the $\text{Re}^{\text{I/III}}$ oxidation.⁹¹

To determine the feasibility of CO_2 reduction by the supramolecular assembly, **PC1** and the corresponding $\text{Re}(\text{Hbpt})(\text{CO})_3\text{Cl}$ fragment were tested for electrocatalytic reduction of CO_2 . First, both catalysts were tested in a homogeneous system using DMF as the solvent and bubbled either with N_2 (blank test) or CO_2 (catalytic test), respectively. Both **PC1** and the $\text{Re}(\text{bpt})(\text{CO})_3\text{Cl}$ complex showed a similar increase in current under a CO_2 atmosphere with an I_{cat}/I_0 of 3.31 for the **PC1** assembly and 3.04 for the $\text{Re}(\text{Hbpt})(\text{CO})_3\text{Cl}$ complex (Fig. S40 & S41[†]). The observed current increase for $\text{Re}(\text{Hbpt})(\text{CO})_3\text{Cl}$ matches well with the value observed for the similar $\text{Re}(\text{Hph})(\text{CO})_3\text{Cl}$ (3.58) complex, which was previously reported for electrocatalytic CO_2 reduction by Nguyen *et al.*⁹² The fairly small difference in catalytic activity between the $\text{Re}(\text{bpt})(\text{CO})_3\text{Cl}$ fragment and the supramolecular assembly **PC1** is consistent with the limited influence of the Ru fragment on the Re catalytic centre as observed for the analogous supramolecular assembly utilising the bpt bridging ligand but with ester rather than acid groups on the peripheral bpy ligands.⁸⁰

Kinetic experiments

Ultrafast UV-visible transient absorption spectroscopy (TA) and time-resolved infrared spectroscopy (TRIR) measurements were used to explore the excited state dynamics of **PC1–3** in solution and when adsorbed on the NiO substrates. The spectra at selected time delays after excitation at 500 nm are shown in Fig. 3 and the results from global analysis of the data are provided in the ESI[†] with extracted time constants provided in Table 2. The spectra are consistent with the formation of a triplet metal-to-ligand charge transfer (MLCT) states.⁹³ For **PC3**, a transient signal was observed below 430 nm, a bleach corresponding to the ground state absorption below 560 nm,

and a broad transient absorption above 560 nm. The spectral features changed little within the timescale of the experiment (6 ns). Global analysis of the data revealed two components, $\tau_1 \sim 200$ ps, and $\tau_2 \gg 1$ ns.

TA measurements of NiO|**PC3** films revealed very different dynamics and global analysis revealed characteristically heterogeneous behaviour for dyes adsorbed on mesoporous semiconductor surfaces, which can be described by a minimum of three components. The two short-lived components were similar in shape and lifetime compared to **PC3*** in solution, $\tau_1 = 13$ ps, and $\tau_2 = 260$ ps, and these are likely to be associated with relaxation of the excited states and charge-transfer from the NiO to **PC3*** to form $\text{NiO}^+|\text{PC3}^-$. $\tau_3 \gg 1$ ns, is assigned to the reduced photocatalyst (**PC3⁻**), with a weak transient absorption < 400 nm and >510 nm. Dietzek *et al.* showed that MLCT states localized on the carboxylic acid-functionalised bipyridine are unfavourable for charge-separation.⁹⁴ In our previous work, we observed the presence of both charge-separated ($\text{NiO}^+|\text{PC}^-$) and excited states ($\text{NiO}|\text{PC}^*$) for photocatalysts with diester-functionalised bipyridine ligands.^{62,63} In this case, for **PC3**, although most of the signal decays within 1 ns, there is evidence for successful charge-separation with the formation of $\text{NiO}^+|\text{PC3}^-$.

The transient absorption spectra for **PC1** in MeCN and **PC1**|NiO followed the same trend as **PC3** and **PC3**|NiO, which suggests that the Re-centre has little impact on the photo-physics of the ruthenium complex. However, **PC2** behaved differently. The excited states for **PC2** in MeCN decayed mostly within the 6 ns window of the experiment. RuPd complexes typically have a low-lying triplet state which quenches the ³MLCT states. **PC2**|NiO contained spectral features and dynamics that were consistent with **PC1**|NiO and **PC3**|NiO, suggesting that charge-transfer from NiO to **PC2*** takes place, but the yield of the charge-separated state, $\text{NiO}^+|\text{PC2}^-$ could be

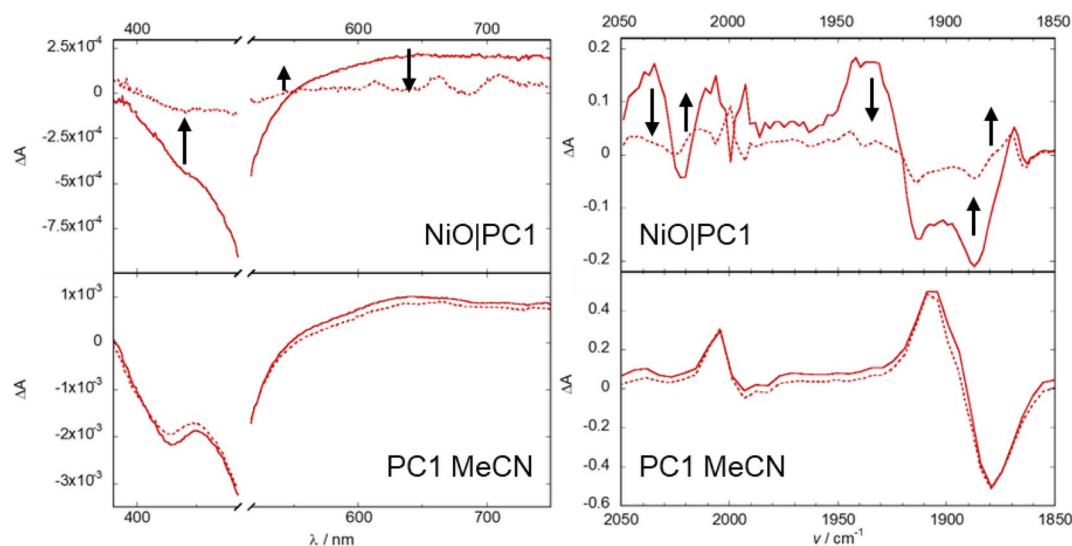


Fig. 3 UV-visible transient absorption spectra (left) and time resolved infrared spectra (right) for **PC1**, 1 ps (solid line) and 1 ns (dotted line) following excitation at 500 nm in acetonitrile (bottom) and on a NiO thin film (top). The transient features at 1 ns spectrum for NiO/**PC1** are imprints of the probe spectrum.



Table 2 Summary of lifetimes estimated by fitting the evolution of the TA and TRIR spectra using global analysis

	PC1	PC2	PC3
TAS in MeCN	$\tau_1 = 160$ ps, and $\tau_2 \gg 1$ ns	$\tau_1 = 35$ ps, and $\tau_2 = 980$ ps, $\tau_3 \gg 1$ ns	$\tau_1 = 200$ ps, and $\tau_2 \gg 1$ ns
TAS on NiO	$\tau_1 = 4.6$ ps, $\tau_2 = 110$ ps, and $\tau_3 \sim 5.1$ ns	$\tau_1 = 6.0$ ps, $\tau_2 = 120$ ps, and $\tau_3 \gg 1$ ns	$\tau_1 = 13$ ps, and $\tau_2 = 260$, $\tau_3 \gg 1$ ns
TRIR in MeCN	$\tau_1 = 130$ ps, and $\tau_2 \gg 1$ ns	$\tau_1 = 22$ ps, and $\tau_2 = 1.2$ ns, $\tau_3 \gg 1$ ns	$\tau_1 = 14$ ps, and $\tau_2 \gg 1$ ns
TRIR film	$\tau_1 = 2.4$ ps, $\tau_2 = 150$ ps, and $\tau_3 \gg 1$ ns	$\tau_1 = 7.0$ ps, $\tau_2 = 120$ ps, and $\tau_3 \gg 1$ ns	$\tau_1 = 7$ ps, and $\tau_2 = 81$, $\tau_3 \gg 1$ ns

Table 3 Average current density (J), faradaic efficiencies (η_{Far}) and turnover frequency per hour (TOF) averaged over 3 hours for CO and H₂ production during photoelectrocatalysis for NiO|PC1 under AM1.5 illumination (100 mW cm⁻²). The applied bias to these systems was -0.2 V vs. Ag/AgCl

PC1 NiO @ -0.2 V vs. Ag/AgCl	pH 5 acetate (under CO ₂)	pH 6.6 50 mM NaHCO ₃ (under CO ₂)	pH 8 phosphate buffer (under CO ₂)	pH 9.2 carbonate buffer (under CO ₂)
$J/\mu\text{A cm}^{-2}$	4.6	2.9	3.8	3.1
$\eta_{\text{Far}}/\%$ (CO)	54.3	94.0	61.8	20.2
$\eta_{\text{Far}}/\%$ (H ₂)	0.46	0.33	0.32	0.1
[CO]/ $\mu\text{mol h}^{-1} \text{cm}^{-2}$	0.12 (± 0.014)	0.27 (± 0.022)	0.11 (± 0.015)	0.09 (± 0.016)
[H ₂]/ $\text{nmol h}^{-1} \text{cm}^{-2}$	1.06 (± 0.13)	0.93 (± 0.074)	0.54 (± 0.076)	0.40 (± 0.072)
TOF (CO) h ⁻¹	0.75	1.63	0.63	0.52
TOF (H ₂) h ⁻¹	0.007	0.007	0.003	0.003

lower to that of the other systems because of the shorter lifetime of the excited state of the photocatalyst.

TRIR measurements provided information regarding the distribution of electron density in the excited state (Fig. 3 and S49–S53[†]). The TRIR spectra for PC1 in MeCN are shown in Fig. 3 and S49.[†] Depletion of the ground state bands at 1735, 1880 and 1993 cm⁻¹, together with formation of transient absorption bands at 1706, 1909 and 2004 cm⁻¹ were observed within the laser pulse. The signals persisted beyond the 5 ns duration of the experiment, which is consistent with the properties of the ester analogue of PC1.²⁸ Global fitting (ESI[†]) of the TRIR data yielded two components but very little spectral change between species, with $\tau_1 = 130$ ps, and $\tau_2 \gg 1$ ns, consistent with the TA data. The transient IR-absorption bands for the Re-CO shift to higher energy relative to the ground state bleach, consistently with electron density removed from the Re centre, resulting in a decrease in π -back bonding to the CO ligands. The transient IR-absorption of the $\nu(\text{CO})$ of the bpy-C(O)OH ligand shifts to higher energy relative to the ground state bleach. This indicates that electron density moves towards the dcbl ligands in the excited state rather than towards the catalytic centre. The TRIR spectra for PC2 and PC3 in MeCN (see ESI[†]) are similar to those of PC1, showing a ground state bleach at 1735 cm⁻¹ and a transient absorption at 1700 cm⁻¹, consistent with the formation of a ³MLCT state. The lifetime of the excited state of PC2 (major component $\tau_2 = 1.2$ ns) was short relative to PC3 in MeCN (major component $\tau_2 \gg 1$ ns). The efficiency of charge-injection could therefore be limited by the comparatively short excited-state lifetime. The results from the solution studies were consistent with our prior results for the ester derivatives.²⁸

We also performed the TRIR experiments for the immobilised photocatalysts, NiO|PC. The data were affected by typical scattering from dye-sensitized films, but the key features were

observed. For NiO|PC3 and NiO|PC2, there was no observable signal in the Re(CO) carbonyl region. It was expected that the C=O stretch for the acid anchoring group would be absent in the IR spectra of photocatalysts adsorbed onto NiO.⁹⁵ Instead, we observed a broad absorption across the fingerprint region corresponding to changes in electron density in the aromatic ligands following excitation. These features decayed on time-scales broadly consistent with the transient absorption spectra. For NiO|PC1, we were able to observe the characteristic bleach and corresponding absorption for the CO ligands on the Re centre. Interestingly, the spectral shape changed during the duration of the experiment, unlike in the solution TRIR. The transient absorption shifts to lower wavenumber at increasing delay time after excitation and this is consistent with an increase in electron density on the Re centre, leading to increased π -back bonding to the CO ligands. This is consistent with the reduction of the photocatalyst and the shift in electron density towards the Re centre would favour catalysis. We attribute this difference to a change in the relative energy of the ligands when the photocatalysts are immobilised on NiO. An increase in the LUMO energy on binding to NiO would be consistent with the blue shift in the absorption spectrum observed (above).

Photoelectrocatalysis

The photocurrent for NiO|PC1 was recorded under chopped light illumination in the presence of an electron acceptor in solution (10 mM tris(ethylenediamine)cobalt(III)) (see ESI[†]). This provides an idealized case to test the photocathodic response where electrons are extracted from the photocathode. The observed current density of approx. 20 $\mu\text{A cm}^{-2}$ confirmed that PC1 transfers charge from the NiO electrode to the electron acceptor in solution in the presence of light.



Photoelectrochemical experiments were then performed with PC1|NiO without the electron acceptor but, instead, bubbling with CO₂ or Ar to evaluate whether PC1 could drive photoelectrocatalysis. The photocurrents and evolved gasses were evaluated under a range of pH between 5 to 9.2. An ionic liquid, 1-ethyl-3-methylimidazolium bis(trifluoromethylsulfonyl) imide, was also tested due to its CO₂-solvation ability.^{96,97} A summary of results for the evaluated systems is provided in Table 3. Similar current densities were recorded for each, under simulated 100 mW cm⁻² intensity AM1.5 irradiation. Fig. 4 and S65–S73† illustrate the photocurrent observed in the devices tested.

At the beginning of each experiment a characteristic spike in the dark current was recorded which has been associated with charging of the electrode surface.⁹⁸ Results from PEC experiments under two different applied bias, -0.2 and -0.5 V vs. Ag/AgCl, are presented in the ESI (Fig. S65–S73†). These potentials are more positive than the flat band potential of TiO₂ that would typically serve as a photoanode.⁹⁹ Control experiments with unsensitized NiO films confirmed that the majority of photocurrent generated arose from light absorption by the chromophore. Typical photocurrent densities were approximately 2–5 μA cm⁻² for all systems in aqueous electrolytes where the applied bias was -0.5 V vs. Ag/AgCl. At a lower applied bias of -0.2 V vs. Ag/AgCl, the dark current for all systems was smaller and surface charging was less pronounced. The photocurrent density for PC2|NiO was remarkably lower than that attained with the ester analogue (*ca.* 40 μA cm⁻²).⁶³ However, the photocurrent density for PC1|NiO was *ca.* 6 times higher than the ester analogue (1.3 μA cm⁻²).⁸⁰ Both were higher than the catalyst-free control, PC3|NiO.

Bubbles formed on the electrode surface during extended periods of irradiation (AM1.5, 100 mW cm⁻²). Care was taken with gas sampling experiments to compare rigorously with appropriate control experiments so that any decomposition of the photocatalyst could be accounted for. Evolved gases from NiO|PC1, NiO|PC2 and NiO|PC3 and the NiO control were tested

over a three-hour irradiation experiment in the three aqueous electrolytes. The headspace was periodically sampled, and the products were quantified by gas chromatography. This required occasional shaking of the cell to promote mixing. The electrolyte was tested for liquid products using ion chromatography, but no formate was detected. The results are provided in Table 3 and the ESI.† A very small amount of H₂ was evolved from NiO|PC2. CO was the main product from NiO|PC1 and in pH 5 acetate buffer and -0.5 V vs. Ag/AgCl, CO was produced with a faradaic efficiency of approx. 91% and a turnover per catalytic centre of 5.4 in 3 hours. The turnover frequency for this system was equal to that reported by Ishitani *et al.* for a RuRe catalyst with a protective polymer layer.⁶⁹ Hydrogen was also detected, but in very small amounts. To confirm the origin of the CO was the gas rather than decomposition of the photocatalyst, the electrolyte was purged with ¹³CO₂, and gaseous products were analysed by mass spectrometry. The major product under both -0.5 V and -0.2 V vs. Ag/AgCl in pH 5 acetate buffer was ¹³CO (see ESI†).

To further optimise the system, the electrolyte was replaced with pH 6.6 NaHCO₃ solution. At -0.2 V, the average photocurrent density was 2.9 μA cm⁻², the faradaic efficiency for CO was 94%, and a turnover of 4.88 per catalyst over 3 hours was attained. To our knowledge, this is the highest selectivity for CO production with a dye-sensitized photocathode. The summary of results for NiO|PC1 in different buffers is shown in Fig. 5.

Surface analysis

After the photoelectrocatalysis experiments, the UV-visible absorption spectra were re-recorded. There was an approx. 20% reduction in the absorption maxima for the three photocatalysts on average according to UV-vis absorption difference spectra. Some of this may be accounted for by the reduction of surface Ni³⁺ states during the reduction (see comment on current spikes above), but it appears that some of the photocatalyst was lost during the reaction. To further characterize the

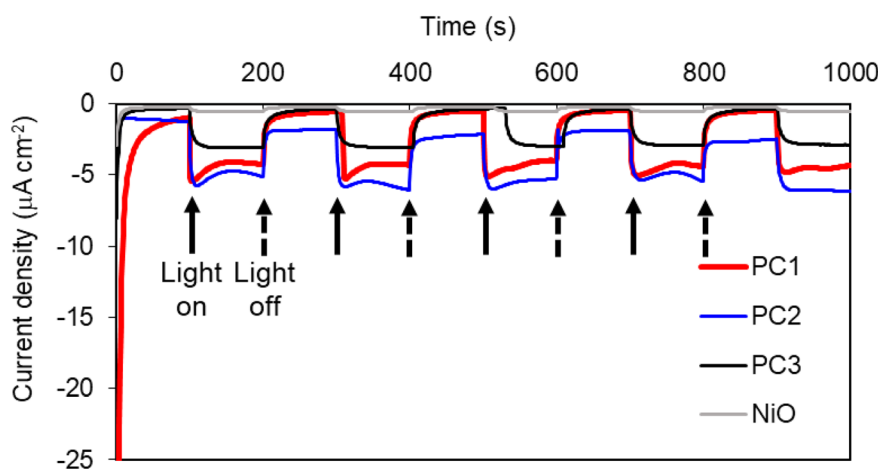


Fig. 4 Representative example of photocurrent in 0.1 M pH 5 acetate buffer under chopped light illumination for photoelectrocatalysis experiments of NiO films sensitized with photocatalysts. The applied bias was -0.2 V vs. Ag/AgCl. NiO, PC1|NiO and PC3|NiO are saturated with CO₂, PC2|NiO is saturated with Ar.





Fig. 5 Evolved CO detected by gas chromatography for PC1/NiO thin films of 0.79 cm^2 working area with an applied bias of -0.2 V vs. Ag/AgCl under 1 sun AM 1.5, (100 mW cm^{-2}) irradiation in various pH buffer solutions as described earlier. Note the pH 6.6 is 50 mM NaHCO_3 solution.

surface of the photocathode before and after the experiments, X-ray photoelectron spectroscopy was used.

The results from the X-ray photoelectron spectroscopy provided evidence the photocatalysts were indeed deposited on the NiO surface and enabled us to determine changes/losses of chromophore after a standard photocatalysis reaction. No noticeable change in the oxidation state of the key elements was determined for our experiments, but there was a slight decrease in signal intensity following photocatalysis for 15 minutes (note these were different samples from the same batch). PC2 was observed to be less stable than PC1, with some loss of ruthenium after the photoelectrocatalysis experiments (Fig. S108 ESI†). This is consistent with the relatively poor faradaic efficiency for PC2 as compared to PC1 (Table 3).

Conclusions

In summary, we have evaluated the photoelectrocatalytic performance of two supramolecular photocatalysts with carboxylic acid anchoring groups, integrated in NiO photocathodes for CO_2 reduction and hydrogen evolution. The RuRe compound PC1, performed better than the ester analogue previously reported.⁶⁰ In contrast, the RuPd compound PC2 was significantly less active for hydrogen generation than the ester analogue.⁶³ The main reason for the change in behaviour is attributed to the difference in lifetime of the excited state. This is much shorter in the case of PC2, which could limit light-induced charge transfer at the NiO|PC2 interface. The NiO|PC1 system is, however, a promising, highly selective system for dye-sensitized NiO photocathodes for CO_2 reduction. Further tuning of PC2 could improve the photocurrent and TON. For example, modifying the bridging ligand to promote charge transfer towards the catalytic metal and away from NiO. The absorption coefficients of the photocatalysts are quite low compared to state-of-the-art photosensitizers in NiO-based DSSCs, which are typically organic systems rather than coordination complexes. A hybrid system with an organic chromophore may lead to much higher light harvesting efficiency and, consequently, better performance. Finally, the solubility of CO_2 in the solvent should be greater to support the higher photocurrent densities and further work on alternatives to MeCN are necessary.

Conflicts of interest

There are no conflicts to declare.

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