Photoelectrochemical hydrogen production in water using a layer-by-layer assembly of a Ru dye and Ni catalyst on NiO†

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Capture and conversion of sunlight into the storable energy carrier H2 can be achieved through photoelectrochemical water splitting using light-absorbing cathodes and anodes bearing H2 and O2 evolving catalysts. Here, we report on the development of a dye-sensitised p-type nickel oxide (NiO) photocathode with a hexaphosphonated Ru(2,2'-bipyridine)3 based dye (RuP3) and a tetraphosphonated molecular [Ni(P2N2)2]2+ type proton reduction catalyst (NiP) for the photoreduction of aqueous protons to H2. A layer-by-layer deposition approach was employed, using Zr4+ ions to link the phosphate units in RuP3 and NiP in a supramolecular assembly on the NiO photocathode. This approach keeps the dye in close proximity to the catalyst and semiconductor surface, but spatially separates NiP from NiO for advantageous electron transfer dynamics. The NiO|RuP3–Zr4+–NiP electrodes generate higher photocurrents and are more stable than photocathodes with RuP3 and NiP co-immobilised on the NiO surface in the absence of Zr4+ cations linking dye and catalyst. The generation of H2 with the NiO|RuP3–Zr4+–NiP hybrid electrode in pH 3 aqueous electrolyte solution during irradiation with a UV-filtered solar light simulator (λ > 400 nm, 100 mW cm⁻², AM1.5G) has been confirmed by gas chromatography at an underpotential of 300 mV (Eappl = +0.3 V vs. RHE), demonstrating the potential of these electrodes to store solar energy in the chemical bond of H2.

Introduction

Photovoltaic technology, such as dye-sensitised solar cells, enables the conversion of solar energy into electricity and is commercially available.¹ Solar fuel devices allow for the storage of solar energy in a chemical energy carrier, but are only in an early stage of development and have not yet reached the required efficiencies and stabilities at low cost suitable for application.² Splitting of water into its elements in a photoelectrochemical (PEC) cell is a potentially sustainable means of generating renewable H2.³ However, a lack of novel strategies to electronically couple a light-harvesting unit to a stable proton reduction catalyst is currently holding back advances in the development of high-performance PEC devices.⁴ Molecular dyes and catalysts offer several advantages over heterogeneous alternatives as their composition and activity can be more easily synthetically controlled and fine-tuned.⁵ They provide an excellent platform to develop rational strategies such as layer-by-layer deposition⁶ to integrate dyes and catalysts on electrode materials and enable detailed mechanistic studies to improve our understanding of photoelectrode assemblies.⁷ Ultimately, these insights may contribute to the establishment of advanced approaches to fabricate electrodes in PEC water splitting cells.

We have previously reported on several suspension-based photocatalytic H2-generation systems with molecular catalysts attached to light-absorbing nanoparticles to advance our understanding of efficient combination of dyes and catalysts on semiconducting surfaces.⁸ However, all of these systems relied on the use of a sacrificial electron donor (SED) to provide the reducing equivalents for fuel formation. SEDs are chemicals of relatively high value and are consumed stoichiometrically during the H2 generation process. They are disadvantageous as half-reaction promoting agents as they often form unknown decomposition products via highly reactive intermediates, which can interfere with the catalytic system, and should ultimately be avoided. One approach to bypass the need for a SED in a H2 evolution system is to immobilise the catalyst onto a photocathode material that can be incorporated into a PEC cell. The redox cycle can then be closed by use of a suitable (photo)anode for water/substrate oxidation.⁹x One of the main challenges for the assembly of molecule-based PEC devices is the design and effective integration of H2 evolving electrocatalysts and dyes onto the electrodes, which should ultimately allow for efficient solar light absorption, charge separation and chemical catalysis.

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Photocathode materials that have been studied for solar H₂ generation include p-silicon, p-GaP, and p-NiO. Of these, p-type NiO offers the possibility to produce nanostructured high-surface area electrodes without the need for specialised techniques. High roughness is a prerequisite for maximising dye and catalyst loading on a given geometric surface area. Dye-sensitised NiO photocathodes in organic or aqueous electrolyte solution have been studied with sacrificial electron acceptors (SEAs), [FeFe]-hydrogenase mimics, and rhodium and cobaloxime catalysts. While charge transfer and recombination kinetics of NiO|dye and NiO|dye-catalyst assemblies have been studied in detail with time-resolved spectroscopy, light-driven H₂ production with dye-sensitised NiO photocathodes remains challenging and only cobaloxime-based catalysts have been reported to generate H₂ with NiO. The binding of the cobaloxime catalyst in these cases was achieved through a linker tethered to NiO bearing a free pyridine moiety, which coordinates to the axial coordination sphere of the cobaloxime. Axial pyridine units coordinated to cobaloximes have been shown to be inherently labile during catalytic turnover and therefore H₂ evolution catalysts with a stable ligand framework are needed.

Nickel bis(diphosphine) type catalysts, [Ni(P₂N₂)₂]²⁺, are attractive proton reduction catalysts as they operate in water and display higher activity than cobaloximes in photocatalytic H₂ production. The [Ni(P₂N₂)₂]²⁺ complex also provides a stable ligand framework with no coordinating moieties becoming labile during the catalytic cycle, and is therefore less susceptible to catalyst dissociation from a surface due to ligand dissociation. [Ni(P₃P)₂]²⁺ catalysts have previously been shown to retain high electrocatalytic activity for H₂ generation from water when immobilised on multi-walled carbon nanotubes (CNTs) via covalent linking or π–π interactions of pyrene moieties, and when immobilised via phosphate groups on mesostructured TiO₂ electrodes. The phosphonated catalyst, NiP (Fig. 1), was also used as catalyst on TiO₂ nanoparticles sensitised with a phosphonated ruthenium-based dye (RuP, Fig. 1) in photocatalytic H₂ generation with a SED. A [Ni(P₃P)₂]²⁺ and a [Ni(PNP)₂]²⁺ type catalyst have previously been immobilised on p-Si, but photocatalytic H₂ generation was only shown for the latter assembly in acetonitrile with trifluoroacetic acid as proton source. Thus, previous work on H₂ generation with [Ni(P₃P)₂]²⁺ catalysts relied on the use of a SED, an applied electrochemical overpotential or non-aqueous conditions to generate H₂.

In this study, PEC H₂ evolution was investigated on a p-type NiO photocathode sensitised with a hexaphosphonated ruthenium tris(bipyridine) based dye (RuP3, Fig. 1) and NiP. A rational and simple procedure of sequential immersion of NiO films into solutions of RuP3, ZrOCl₂ and NiP was employed to produce electrodes of the type NiO[RuP3–Zr⁴⁺–NiP] (Fig. 2), where the catalyst is spatially separated from the semiconductor electrode to reduce inefficiencies from charge-recombination. Our study shows that increasing the distance of the catalyst from the NiO surface results in substantially enhanced PEC performance in terms of photocurrent and stability when compared to NiO[RuP3–NiP], where the dye and catalyst are co-immobilised on NiO. We demonstrate a dye-sensitised NiO hybrid electrode with an adsorbed Ni catalyst, which has the potential to store solar energy in the bond of H₂ without the decomposition of valuable sacrificial agents or an applied overpotential. Rational layer-by-layer assembly of dye and catalyst on a semiconductor electrode is therefore established as a route to produce functional PEC H₂ production systems.

**Results and discussion**

**Assembly and pre-catalysis characterisation**

Mesoporous NiO was grown on tin-doped indium oxide (ITO)-coated glass electrodes by a previously reported hydrothermal method from a solution containing Ni(NO₃)₂·6H₂O and hexamethylenetetramine (0.25 M each). NiO electrodes were sensitised by immersion in a RuP3 solution (1 mM in H₂O) overnight to give NiO[RuP3] for PEC experiments.
Immobilisation of monolayers of molecular compounds via phosphanic acids on metal oxide surfaces is a well-established tool for surface modification.\textsuperscript{25,26} NiO was immobilised on the NiO|RuP3 photocathodes in a layered assembly using Zr\textsuperscript{4+} to link the phosphonates of dye and catalyst (Fig. 2; see below for more details). Zr\textsuperscript{4+} ions bind strongly to phosphonic acid moieties and were previously used for creating multi-layered Zr\textsuperscript{4+}-phosphate structures.\textsuperscript{25,27} Zr\textsuperscript{4+} was bound by dipping NiO|RuP3 electrodes into a solution of ZrOCl\textsubscript{2} (5 mM in EtOH/ H\textsubscript{2}O, 60/40 v/v) for 2 h, rinsing with H\textsubscript{2}O and EtOH and drying under a stream of N\textsubscript{2}. The NiO|RuP3–Zr\textsuperscript{4+} electrodes were subsequently submerged in a solution of NiP (0.5 mM in MeOH) for at least 2 h to obtain the supramolecular NiO|RuP3–Zr\textsuperscript{4+}–NiP electrode assembly with one layer of dye and one layer of catalyst linked by Zr\textsuperscript{4+}. All immobilisation steps were carried out at room temperature.

In previously reported homogeneous, SED-promoted photocatalytic H\textsubscript{2} generation systems, the ratio between the dye and the catalyst was found to strongly influence the overall performance.\textsuperscript{17} We have therefore varied the ratio of co-immobilised RuP3 and NiP on the p-NiO photocathodes. The layer-by-layer deposition approach allowed the RuP3 and NiP loading on the metal oxide surface to be controlled more precisely, creating Zr\textsuperscript{4+}-linked dye/catalyst layers by repeated immobilisation cycles.\textsuperscript{25,26} Repeated deposition of Zr\textsuperscript{4+} and either RuP3 dye or NiP catalyst gave electrodes of the structure NiO|[(RuP3–Zr\textsuperscript{4+})\textsubscript{2}–NiP] (containing two layers of RuP3–Zr\textsuperscript{4+}) or NiO|[(RuP3–Zr\textsuperscript{4+}–NiP)]\textsubscript{2} (containing two layers of Zr\textsuperscript{4+}–NiP). For control experiments, NiO|RuP3 electrodes were submerged directly in NiP solution without prior immobilisation of Zr\textsuperscript{4+} ions, and the so-obtained NiO|RuP3–NiP electrodes were also studied. Additionally, RuP3 was replaced with the diphosphonated dye RuP (Fig. 1), which only allows binding to NiO and does not offer any phosphonic acid moieties to generate Zr\textsuperscript{4+}-linked phosphate networks.

The photocurrent response of NiO|RuP3 electrodes with different NiO film thickness was recorded in a PEC experiment in aqueous electrolyte solution (0.1 M Na\textsubscript{2}SO\textsubscript{4}, pH 3) in the presence of the SEA 4,4\textprime;-dithiodipyridine (DTDP, 1 mM; Fig. 3 and S1†) at room temperature. DTDP is reduced in two irreversible steps, with the first reduction occurring at $E_\text{p} = -0.06$ V vs. the reversible hydrogen electrode (RHE). Thickness, morphology and crystallinity of the NiO films was studied by varying the reaction conditions (see Table S1 and Fig. S2 and S3† for powder X-ray diffraction pattern and scanning electron microscopy images). The optimised films have an average pore size (measured as distance between sheets) of 290 ± 81 nm and a film thickness of 2 μm (Fig. 3). The flatband potential of this NiO electrode was determined via electrochemical impedance spectroscopy at 0.75 V vs. RHE from Mott–Schottky analysis (Fig. S4†) and shows Nernstian pH dependence (one electron/one proton couple) in aqueous electrolyte solution, which is in agreement with previous reports on NiO electrodes.\textsuperscript{29}

The photocurrent response of NiO|RuP3 and the loading capacity for RuP3 on NiO films with different film thickness was studied by PEC methods and desorption of the attached dye with 0.1 M NaOH and monitoring the UV-vis absorption at $\lambda = 295$ nm in solution (Table S1†). The dye loading capacity increased as expected with increasing film thickness, but the highest photocurrent density was achieved with a 2 μm thick NiO film, despite the higher dye loadings for thicker films (Fig. 3, Table S1†). This observation is consistent with the known limitation of hole mobility and lifetime in NiO,\textsuperscript{28} which suggests that the photocurrent generated with RuP3-sensitised NiO photocathodes is a balance between limiting charge transport in thick films and dye loading on thin films.

We subsequently investigated the interface between molecular components and the p-type NiO surface of the NiO|RuP3–Zr\textsuperscript{4+}–NiP electrodes by UV-vis (Fig. S5†), attenuated total reflection infrared (ATR-IR; Fig. S6†), X-ray photoelectron (XPS; Fig. S7†) and inductively coupled plasma-optical emission spectroscopies (ICP-OES). The UV-vis spectrum of a bare NiO thin film shows the typical strong absorption at $\lambda < 380$ nm, and also a broad indirect transition at $\lambda > 550$ nm.\textsuperscript{29} All RuP3-sensitised NiO electrodes exhibit the typical MLCT absorption band for Ru(bpy)\textsubscript{3}*-based dyes around $\lambda = 465$ nm and strong π-π* absorption in the UV region.\textsuperscript{34} When NiP is present on the electrodes a small shoulder at $\lambda = 520$ nm could be observed, which is typical for the d-d transition in NiP as also observed in solution spectra (Fig. S5†).

The ATR-IR spectra of the NiO|RuP3 electrodes at $r = 800$ to 1250 cm\textsuperscript{−1} in the presence and absence of Zr\textsuperscript{4+} and NiP are shown in Fig. S6†. This region is characteristic for P=O and P–O–R vibrations, which are also observed in the spectra of powdered RuP3, RuP and NiP.\textsuperscript{34} NiO|RuP3 electrodes show a broad band at $r \sim 1100$ cm\textsuperscript{−1} attributed to P=O vibrations in RuP3, which is absent in a bare NiO film. In a NiO|RuP3–Zr\textsuperscript{4+} electrode, a small additional IR band at $r \sim 983$ cm\textsuperscript{−1} appears when compared to NiO|RuP3, and the signal increases and shifts to slightly higher wavenumbers for NiO|RuP3–Zr\textsuperscript{4+}–NiP ($r \sim 986$ cm\textsuperscript{−1}) and NiO|RuP3–Zr\textsuperscript{4+}–NiP ($r \sim 992$ cm\textsuperscript{−1}). This shift is in good accordance with previously reported values for Zr\textsuperscript{4+} bound to phosphonic acid groups.\textsuperscript{25,26}

![Fig. 3 Photocurrent response of NiO|RuP3 under visible light irradiation (λ > 400 nm, AM1.5G filter, 100 mW cm\textsuperscript{−2}) after 100 s at $E_{\text{app}} = 0.3$ V vs. RHE in Na\textsubscript{2}SO\textsubscript{4} (0.1 M, pH 3) in the presence of 4,4\textprime;-dithiodipyridine (DTDP, 1 mM) as SEA and dye loading (no applied potential; cm\textsuperscript{−2} refers to the geometrical surface area of the NiO electrode) with varying thicknesses of the NiO films. Inset: SEM image of optimised p-type NiO electrode (2 μm film thickness).](image-url)
XPS analysis was used to further characterise the modified NiO surface, and the presence of Ru and Zr on NiO after each layering step was confirmed (Fig. S7†). The signal for NiP could not be distinguished from the NiO background signal. Instead, mesoporous indium-tin oxide (ITO, 3.3 μm thickness) electrodes were prepared according to a previously reported procedure24 and used here as a scaffold. The XPS signals of ITO|RuP3–NiP, ITO|RuP3–Zr4+ and ITO|RuP3–Zr4+-NiP electrodes (prepared analogously to the related NiO hybrids) were analysed. In the presence of NiP, peaks at 855.0 and 872.5 eV were observed and were assigned to the Ni 2p3/2 and Ni 2p1/2 of the molecular NiP, respectively.17,18 The 2p signal for ITO|RuP3–NiP and ITO|RuP3–Zr4+-NiP electrodes is broad (133.0–131.0 eV) and consists of signals for the phosphonic acid bound to ITO4+ and for the phosphine ligand.33 In contrast, ITO|RuP3–Zr4+ showed only a sharp XPS peak at 133.5 eV, which is assigned to the –PO(OH)2 groups.34 As the RuP3–Zr4+-NiP was assembled in identical manner on ITO and NiO, the presence of intact NiP on ITO supports its integrity on NiO prior to catalysis.

Finally, we quantified the immobilised dye and catalyst on NiO by UV-vis spectroscopy and ICP-OES analysis after desorption in aqueous NaOH (0.1 M). The overlap of RuP3 and NiP in the electronic absorption spectra prevented the simultaneous determination of dye and catalyst loading on the electrodes and ICP-OES measurements were used instead. Quantification of NiP on NiO was challenging due to the high nickel background from the electrode. Mesoporous ITO electrodes were used again as a scaffold and analysis of ITO|RuP3, ITO|RuP3–NiP, ITO|RuP3–Zr4+ and ITO|RuP3–Zr4+-NiP electrodes was carried out. The quantity of NiP on NiO was estimated from the Ru : Ni ratio found on the ITO electrodes (Table 1). The ICP-OES data for the amount of RuP3 immobilised on bare NiO (10.7 ± 0.3 nmol cm−2) correlates well with quantification by UV-vis spectroscopy after dye desorption (10.8 ± 0.8 nmol cm−2). Analysis of ITO|RuP3–NiP electrodes revealed a Ru : Ni ratio of 2.6 ± 0.04 : 1 and this ratio decreases only slightly to ~2.2 ± 1.5 : 1 on ITO|RuP3–Zr4+-NiP, indicating that a similar amount of NiP is immobilised in the presence and absence of Zr4+. This ratio allows us to estimate a NiP loading of 4.98 ± 3.54 nmol cm−2 for NiO|RuP3–Zr4+-NiP, which is comparable to the loading of a cobaloxime catalyst on CdSe-sensitised NiO (~3 nmol cm−2 for a 5.6 μm thick NiO film).11b

**Photoelectrochemical experiments**

The photocathodes were studied in PEC experiments in aqueous electrolyte solution (0.1 M Na2SO4, pH 3) in a three-electrode setup with a Ag/AgCl reference and a Pt counter electrode at room temperature. Linear sweep voltammetry (LSV) scans were recorded between E = +0.6 V and −0.15 V vs. RHE at a scan rate of 5 mV s−1 in the dark and under chopped and UV filtered irradiation with simulated solar light (λ > 400 nm, AM1.5G; Fig. 4a). All NiO electrodes (irrespective of being loaded with molecules or not) displayed a partially reversible oxidation wave between E = 0.42 and 0.48 V vs. RHE, which is assigned to the Ni3+/Ni2+ redox couple (Fig. S8†).24 In the absence of dye, only marginal photocurrent densities (reported per geometrical surface area) were observed in the studied potential range. Transient and small cathodic photocurrents were observed upon irradiating NiO|RuP3 ([j] ~ 1.4 μA cm−2 at 0.5 V vs. RHE) and NiO|RuP3–Zr4+ ([j] ~1.2 μA cm−2 at 0.5 V vs. RHE) electrodes, indicating hole injection of the excited dye, RuP3*, into the valence band of NiO (Fig. 2, for details see Mechanistic interpretation below). Quenching of RuP3* by NiO results in the formation of a reduced dye, RuP3−, which is then likely to undergo rapid charge recombination with the holes generated in NiO in absence of a suitable electron acceptor.13,15b A comparable photoresponse was observed between NiO|RuP3 and NiO|RuP3–NiP, indicating that direct binding of the NiP to NiO results in an unproductive assembly with negligible electron accumulation at NiP. When a NiO|RuP3–Zr4−NiP electrode was used, however, an initial photocurrent density up to [j] ~ 5.56 μA cm−2 was achieved under the same experimental conditions (Fig. 4a).

Subsequently, the importance of the phosphonic acid moieties in the assembly of functional electrodes was investigated (Fig. S9†). When the diphenylphosphonated dye RuP (Fig. 1c) is immobilised on NiO instead of RuP3, no phosphonic acid binding sites are available for Zr4+. A comparable photocurrent was observed for a NiO|RuP–Zr4–NiP and NiO|RuP–NiP electrode.

<table>
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<tr>
<th>Composition</th>
<th>RuP3/nmol cm−2</th>
<th>NiP/nmol cm−2</th>
<th>[j]/μA cm−2(a)</th>
<th>[j]/μA cm−2(b)</th>
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<tr>
<td>NiO</td>
<td>RuP3</td>
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<td>0.51</td>
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<tr>
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<td>0.76</td>
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<tr>
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<td>4.98 ± 3.54(c)</td>
<td>6.40</td>
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<tr>
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<td>n.d.</td>
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<td>14.3 ± 0.2</td>
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<td>41.3 ± 20.3</td>
<td>19.2 ± 9.9</td>
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(a) Measured in the absence and (b) presence of DTDP as SEA. (c) Calculated value from RuP3 : NiP ratio on ITO. (d) Not determined.
a NiO|RuP3–Zr4+–NiP electrode in the presence of DTDP in solution ([j] = 14.0 μA cm⁻²) compared to a SEA-free system ([j] = 6.4 μA cm⁻²). This suggests that surface-bound NiP is efficiently reduced by the generated photoelectrons and electron transfer from the reduced catalyst to the dissolved SEA in the electrolyte solution is faster than competing charge recombination. An increased photocurrent in the presence of DTDP suggests that the reduction of the SEA is faster than NiP-promoted catalysis.

The effect of the ratio of dye and catalyst was also studied by preparing electrodes with either two layers of dye, NiO|([RuP3–Zr4+]2–NiP), or two layers of catalyst, NiO|([RuP3–Zr4+]2–NiP)2. The photocurrent generated by a NiO|([RuP3–Zr4+]2–NiP) electrode after 100 s chronoamperometry at E_spp = 0.3 V vs. RHE ([j] = 2.18 μA cm⁻²) was almost three times lower than the photocurrent of a NiO|RuP3–Zr4+–NiP electrode (Table 1). In contrast, approximately 40% increase in photocurrent was observed for a NiO|RuP3–(Zr4+–NiP)2 electrode ([j] = 8.82 μA cm⁻²). These experiments confirmed that there is a significant dependence of the number of layers on the photocurrent response and show that transient photocurrents generated by dye-sensitised NiO electrodes are dependent on the strategy used to co-immobilise dye and catalyst on the surface. This layer-by-layer approach has previously been successful for dye-sensitised photoanodes for water oxidation, but has not yet been explored for fuel-forming photocathodes.

The formation of H2 at the NiO|RuP3–Zr4+–NiP electrode was then studied by controlled potential photoelectrolysis (CPPE; Fig. 5). Irradiation of the NiO|RuP3–Zr4+–NiP and NiO|RuP3–(Zr4+–NiP)2 electrodes at E_spp = 0.3 V vs. RHE in a two-compartment PEC cell generated 6.11 ± 0.68 and 6.8 ± 0.9 nmol H2 in the headspace (analysed by gas chromatography with N2 carrier gas) with a faradaic efficiency of 8.6 ± 2.3% and 10.1 ± 1.8%, respectively (Table S2†). Increasing the light intensity to 200 mW cm⁻² or using an E_spp = 0.5 V vs. RHE did not alter the H2 production rate nor improve the faradaic efficiency for NiO|RuP3–Zr4+–NiP. In the absence of NiP or in the dark, no H2 generation was observed. NiO|RuP3–NiP and NiO|RuP3–(Zr4+–NiP)2 electrodes produced only traces of H2 below the limit of quantification (LOQ) of the thermal conductivity detector. The low activity of the NiO|RuP3–Zr4+–NiP electrode is likely due to inefficient charge transfer dynamics at the NiO|RuP3 interface and between reduced NiP and oxidised RuP3 or NiO (Fig. 2b). Nevertheless, confirmation of H2 evolution at the photocathode is an important step towards the goal of implementing dye-sensitised NiO photocathodes in full water splitting PEC cells.11,13

Post-catalysis characterisation

The electrode stability was further examined by ICP-OES analysis of the Ru content in both electrolyte solution and on the electrodes following 2 h CPPE at E_spp = 0.3 V vs. RHE. For NiO|RuP3–Zr4+–NiP a substantial amount of the initially loaded Ru (6.3 ± 3.1 nmol cm⁻²) was recovered from the used electrodes (3.4–7.4 nmol Ru cm⁻²) upon desorption in aqueous NaOH (0.1 M). Almost no Ru leaked into the electrolyte solution of the
working or counter compartments (<0.2 nmol total Ru content). The opposite was observed for the NiO|RuP3–NiP electrodes after 2 h CPPE and 4.9–5.3 nmol total Ru content were found in the electrolyte solution of the working compartment and very little Ru (0.39–0.74 nmol cm$^{-2}$) was recovered from the used working electrode. The Zr$^{4+}$ interlayer might contribute to stabilising RuP3 on the surface of NiO electrodes under CPPE conditions. The stability of the NiO|RuP3–Zr$^{4+}$–NiP working electrodes against material degradation was also confirmed by the lack of observable changes in the cyclic voltammograms before and after CPPE (Fig. S13†) and a significant alteration for NiO|RuP3–Zr$^{4+}$ electrodes after CPPE at 0 V vs. RHE (Fig. S14†), indicating material instability of the latter.

The XPS spectra of a NiO|RuP3–Zr$^{4+}$–NiP electrode were recorded and confirmed the presence of both RuP3 (281 eV, Ru 3d$_{5/2}$) and Zr$^{4+}$ (182.6 and 185 eV for Zr 3d$_{5/2}$ and Zr 3d$_{3/2}$, respectively) before and after 2 h CPPE at $E_{\text{appl}} = 0.3$ V vs. RHE (Fig. S7†). No new peaks in the Ru 3d$_{5/2}$ region were observed after CPPE to indicate formation of Ru/RuO$_x$ particles. The presence and integrity of NiP on the NiO|RuP3–Zr$^{4+}$–NiP electrode after CPPE could not unambiguously be confirmed by XPS due to the presence of the NiO. However, the stability of NiP on the electrode surface of a mesostructured TiO$_2$ electrode under reducing conditions ($E_{\text{appl}} = -0.25$ V vs. RHE) in the dark in otherwise identical aqueous electrolyte solution (0.1 M Na$_2$SO$_4$, pH 3) has previously been confirmed.$^{22}$

**Mechanistic interpretation**

Visible light irradiation of RuP3 generates RuP3*, which can in principle be quenched by two different mechanisms in the NiO|RuP3–Zr$^{4+}$–NiP assembly: (a) oxidative quenching of RuP3* ($\epsilon([\text{RuP}3^*/\text{RuP3}]) \sim -0.78$ V vs. the normal hydrogen electrode (NHE), HOMO–LUMO gap $E_{\text{HOMO}}^{\text{RuP3}} = 2.19$ eV, $\lambda_{\text{em}} = 567$ nm, Fig. S15†) by NiP, followed by hole injection from the oxidised RuP3* ($\epsilon([\text{RuP}3^*/\text{RuP3}]) \sim 1.41$ V vs. NHE, Fig. S16†) into NiO ($E_{\text{red}}(\text{NiO}) = 0.57$ V vs. NHE at pH 3)$^{18}$ or (b) reductive quenching of RuP3* by fast electron transfer from NiO into the HOMO of the dye to form the reduced species RuP3$^-$ ($\epsilon([\text{RuP}3^-/\text{RuP3}]) \sim 0.80$ V vs. NHE, Fig. 2b).$^{29}$ RuP3$^-$ has a sufficiently negative ground state reduction potential ($\epsilon([\text{RuP3}/\text{RuP3}^-]) \sim -1.29$ V vs. NHE)$^{16}$ to provide driving force for reduction of NiP ($\epsilon([\text{NiP}]) = -0.35$ V vs. NHE at pH 4.5).$^{17}$ The reductive quenching mechanism for the NiO|RuP3–Zr$^{4+}$–NiP system is much more likely since hole injection from a dye into NiO happens on very fast timescales of hundreds of femto- to picoseconds.$^{13,15}$ Additionally, spectroscopic studies of a homogenous RuP–NiP system for photocatalytic H$_2$ generation have shown previously that RuP$^*$ does not directly inject electrons into NiP in solution or when both molecules are anchored on ZrO$_2$ as a non-injecting matrix. Reductive quenching by a SED or p-Ni yields RuP$^-$, which was shown to efficiently reduce NiP.$^{17}$ Reductive quenching of the NiO|RuP3–Zr$^{4+}$–NiP system would provide the catalyst with an overpotential of ~0.9 V (Fig. 2b).

Our results also suggest an important contribution of the Zr$^{4+}$ interlayer to the efficiency of electron transfer away from the electrode surface to the associated catalyst. As Zr$^{4+}$ is inert against reduction, it is unlikely that the Zr$^{4+}$ coordinated to RuP3 and NiP is acting as an electron relay similar to TiO$_2$ in H$_2$ evolving dye–TiO$_2$–catalyst suspension systems.$^{18}$ It is more likely that the Zr$^{4+}$ layer prevents NiP from binding directly on the NiO surface in close proximity to NiO|RuP3, and that spatial separation of the reduced catalyst from the generated holes in NiO is beneficial to avoid charge recombination which is fast when dye and catalyst are in close proximity.$^{33}$ Reports on the catalytic activity of photoanodes for water oxidation showed that stability was enhanced when the catalyst was spatially separated from the dye-sensitised electrode.$^{25,26,30}$ Spectroscopic studies of a TiO$_2$|RuP3–Zr$^{4+}$–catalyst photoanode demonstrated that electron back transfer was slowed down significantly compared to a TiO$_2$|catalyst electrode,$^{25b}$ which was assigned to increased spatial separation. Similar effects were also found for electron transfer between photoexcited TiO$_2$ and cobaloxime catalysts with different linker lengths.$^{41}$ These experiments show the high potential of using rationally designed dye–catalyst assemblies on photocathodes.

**Conclusions**

A dye-sensitised photocathode with a co-immobilised nickel catalyst for light driven H$_2$ generation in water has been reported. Co-deposition of the Ru dye RuP3 and Ni catalyst NiP on the p-type semiconductor NiO in a supramolecular assembly of Zr$^{4+}$-phosphonates provided us with a tool to control spatial arrangement of individual species without the need for elaborate chemical synthesis of dye–linker–catalyst dyads. This layer-by-layer approach keeps the catalyst in close proximity to the dye, but increases the distance to the semiconductor electrode. Our PEC experiments confirm that directed forward electron transfer from the excited dye to the catalyst is efficient and recombination kinetics are slowed down. The catalyst was therefore able to turn over and the hybrid electrode able to photo-generate H$_2$ at an electrochemical underpotential,
thereby demonstrating the potential of this system to store light in the chemical bonds of H₂. Layer-by-layer assembly of dye and catalyst is therefore established as a novel strategy to produce molecule-based photocathodes for H₂ evolution.

The efficiency of the presented system can be improved in the future by optimising the NiO|dye interface such as developing phosphonated push-pull dyes optimised for p-NiO, and fine-tuning of the H₂ evolution catalyst. Further work will also include the synthesis and investigation of dyes, which are better suited to inject holes into the NiO valence band. Optimised molecule-containing photocathodes might ultimately become an attractive component for use in PEC cells for full water splitting.¹⁴,²²

**Experimental section**

**Materials and methods**

**NiP**,²³ **RuP**,²³ and **RuP3**²⁴ (Fig. 1) were synthesised and characterised as previously reported. Chemicals for analytical measurements were purchased in the highest available purity and used without further purification. Ni(NO₃)₂·6H₂O (Fisher Scientific, extra pure, 250 g) and hexamethylenetetramine (Sigma Aldrich, ReagentPlus, 99%) were used for the preparation of NiO electrodes. ITO nanopowder (Sigma Aldrich, particle size below 50 nm) was used for the preparation of mesostructured ITO electrodes. All electrochemical and analytical measurements were performed using Milli-Q® H₂O (resistivity > 18 MΩ cm). ITO-covered glass sheets (Vision Tek Systems Ltd., R = 12 Ω cm⁻², thickness 1.1 mm) were cut into 3 x 1 cm² slides for preparation of working electrodes.

**Physical characterisation**

An FEI Phillips XL30 SFEG SEM was used in the Electron Microscopy Suite in the Cavendish Laboratory, University of Cambridge. Powder X-ray diffraction (XRD) analysis was performed using a PANalytical B.V X'Pert Pro X-ray diffractometer in the Department of Chemistry, University of Cambridge. XPS analysis was performed at the NEXUS XPS facility at Newcastle University. XPS spectra were calibrated to the C 1s signal at 284.8 eV. Quantification of Ni and Ru on the electrodes via ICP-OES (PerkinElmer Optima 2100 DV spectrometer) was carried out at the Department of Geography, University of Cambridge. High-resolution ATR-IR spectra were recorded on a Thermo Scientific Nicolet iS 50 FT-IR spectrometer with an ATR unit. UV-vis absorption spectra of electrodes were recorded in transmission mode in air, and UV-vis solution spectra were recorded in a quartz glass cuvette (1 cm path length) with a Varian Cary 50 UV-vis spectrophotometer. Fluorescence emission spectra of RuP3 in Na₂SO₄ (0.1 M, pH 3) were measured with an Edinburgh Instruments FS5 spectrofluorimeter.

**Electrochemical impedance spectroscopy**

Electrochemical impedance spectroscopy (EIS) was performed on an IviumStat potentiostat in the frequency range of 100 kHz to 0.1 Hz in aqueous Na₂SO₄ solution (0.1 M, pH 3, 5 and 7). An equivalent circuit (inset Fig. S4†) consisting of electrolyte solution resistance (Rₑ), constant phase element (CPE), and interfacial charge transfer resistance (R_Cr) was fitted to Nyquist plots obtained at different potentials in ZView® (Scribner Associates Inc.) to retrieve C_SC values required to form a Mott–Schottky plot. The Mott–Schottky equation

\[
\frac{1}{C_{SC}^2} = \frac{2}{\varepsilon \varepsilon_0 A^2 eN} \left( V - V_B + \frac{k_B T}{e} \right),
\]

where C_SC is the interfacial capacitance, A is the interfacial area, N the density of acceptors, V the applied potential, V_B the flatband potential, k_B the Boltzmann constant, T the absolute temperature, and e the electronic charge, was used to calculate the flatband potential of NiO. 1/C_SC² plotted against applied potential results in a straight line with the intercept being equal to V_B + k_B T/e. The negative slope is indicative of the p-type character of the NiO electrodes.

**Preparation of nanostructured NiO electrodes**²⁶

ITO covered glass slides (3 x 1 cm²) were cleaned by successive sonication in ethanol and acetone for 10 min, respectively. After drying in air, the conductive surface was confined to ca. 1 x 1.5 cm² with electrical tape. The ITO glass slides were placed at approximately 45° with the conductive side facing down into separate vials and a solution of Ni(NO₃)₂·6H₂O and hexamethylenetetramine (250 mM each, 6 mL) was added to each vial. For hydrothermal growth of the films the vials were placed into an oven (Thermo Scientific Heratherm) for 15, 30, 45 or 60 min at 90 °C and the reaction was stopped by the immediate addition of distilled water. The slides were rinsed with water, dried in air and the tape removed before the slides were annealed in a furnace (Carbolite) at 450 °C in air for 30 min (heating rate 20 °C min⁻¹), and left to cool to room temperature in the furnace chamber.

**Preparation of mesostructured ITO electrodes**²²

ITO covered glass slides (3 x 1 cm²) were cleaned by heating them in a mixture of NH₄OH (1 M, H₂O/H₂O/H₂O (3:10:69) (1 : 5 : 1)) at 70 °C for 30 min. The slides were rinsed with distilled water and dried in an oven at 180 °C for 1 h. The surface of the conductive side was confined with Scotch tape to circular 0.28 cm². ITO nanopowder (20% w/w) was sonicated in ETOH (5 M acetic acid) for 1 hour and then doctor bladed on the ITO support. After drying in air, the tape was removed and the electrodes were annealed in a furnace (Carbolite) at 400 °C in air for 1 h (4 °C min⁻¹ heating rate) and left to cool to room temperature in the oven chamber.

**Assembly of molecular photocathodes**

The layer-by-layer deposition of dye (RuP3 or RuP) and catalyst (NiP) was achieved by sequential immersion of the electrodes in solutions containing the molecular compounds at room temperature. NiO electrodes were submerged in solutions of RuP3 or RuP (1 mM in H₂O, overnight, washed with H₂O and EtOH), ZrOCl₂ (5 mM in ETOH (60% v/v in H₂O), 2 h, washed with H₂O and EtOH) and NiP (0.5 mM in MeOH, 2–3 h, washed with MeOH) to build up the supramolecular layered assemblies.
The electrodes were dried in a stream of N₂ after each immobilisation step. Multiple (n) catalyst and dye layers were assembled by repeating the immobilisation cycles n times. RuP₃ sensitised mesostructured ITO electrodes (ITO[RuP₃]) for controls were prepared according to the same procedure. The electrodes were stored in the dark prior use.

Quantification of immobilised compounds on NiO

(A) Quantification of RuP₃ by UV-vis spectrophotometry. A NiO[RuP₃] electrode was immersed in aqueous NaOH (0.1 M, 1.0 mL) for 5 min to desorb the phosphonated species. The absorbance of the resulting solution at λ = 467 and 297 nm was used to determine the amount of RuP₃ detached from the surface (assuming quantitative detachment of phosphonic acids in pH 13 solution).

(B) Quantification of Ru, and Ni via ICP-OES. Surface-bound species were detached by immersing the electrodes in aqueous NaOH (0.1 M, 1.0 mL) for 5 min. The electrodes were rinsed with 10 mL H₂O and then HNO₃ (2% v/v) was added to a final volume of 25 mL. Samples were prepared and analysed from at least two individual electrodes for each condition and the averages are reported. Errors are given as maximum deviation from the average.

Determination of RuP₃ oxidation potential

E[RuP₃³⁺/RuP₃] was determined by recording a CV of RuP₃ in Na₂SO₄ (0.1 M, pH 3) with a boron doped diamond working, Ag/AgCl/KCl(c sat.) reference and Pt mesh counter electrode. The HOMO-LUMO gap Eₕ₋ₐ(RuP₃) was determined as the intersection of the normalised absorption and emission spectra of RuP₃ in Na₂SO₄ (0.1 M, pH 3).²⁴ E[RuP₃³⁺/RuP₃⁺] was estimated by E[RuP₃³⁺/RuP₃⁺] = E[RuP₃³⁺/RuP₃] – Eₕ₋ₐ(RuP₃).

Photoelectrochemistry

Electrochemical measurements were performed on an Ivium CompactStat potentiostat using a three-electrode configuration in a custom-made cell with a flat borosilicate window for PEC experiments. The prepared dye-sensitised NiO slides (1 cm² geometrical surface area confined with electrical tape) were studied as working electrodes. A Pt mesh counter electrode, and a Ag/AgCl/KCl(c sat.) reference electrode were used in aqueous electrolyte solution (0.1 M Na₂SO₄, pH 3). Prior to any measurements, all solutions were purged for at least 10 min with N₂ to remove O₂. PEC experiments were performed under irradiation from the back-side of the electrodes with visible light (λ > 400 nm, UQG Optics filter) using a solar light simulator (Newport Oriel, 150 W, 100 mW cm⁻²) equipped with an AM1.5G and IR water filter.

CPPE was carried out in a custom-made 2-compartment PEC cell separated by a Nafion membrane (quartz glass window, 1.5 cm diameter). In the working compartment, 14.5 mL of electrolyte solution were used, leaving a gas headspace of 4.9 mL. In the counter compartment, 4.5 mL of electrolyte solution were used, leaving a gas headspace of 3.1 mL. Both compartments were purged prior to electrolysis with N₂ containing CH₄ (2%) as internal standard for gas chromatography measurements. The electrolyte solution [0.5 M Na₂SO₄, pH 3] in both compartments was stirred during the experiments. The amount of H₂ in the headspace of the electrolysis cell was quantified by an Agilent 7890A Series gas chromatograph equipped with a 5 Å molecular sieve column. The GC oven temperature was kept constant at 45 °C, N₂ was used as a carrier gas at an approximate flow rate of 3 mL min⁻¹ and a thermal conductivity detector was used. In a standard experiment, the working electrode was held at the respective potential (E_app) in the dark for the first h, and after confirming the absence of headspace H₂, it was exposed to visible light (λ > 400 nm, 100 or 200 mW cm⁻²) for 2 h at E_app. All CPPE experiments were carried out at least three times, unless noted otherwise. The mean values of the measurements are reported along with their standard errors.¹⁷w

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37 This work, estimated by the intersection point of the normalised absorption and emission spectra of RuP3 in Na2SO4 (0.1 M, pH 3).
38 This work, measured by electrochemical impedance spectroscopy of optimised NiO films in aqueous electrolyte solution.