

Comparing Activity and Effectiveness of (Ru/Pd), (Ru/Pt) Photocatalysts for Hydrogen Generation, Photochemical Reaction with HPLC

Hamid M. Younis^{*1} and Johannes G. Vos²

¹Faculty of Sciences, Chemistry Department, Sirte University, Sirte-Libya,

²SRC for Solar Energy Conversion, School of Chemical Sciences, Dublin City University, Dublin 9, Ireland.

Received date here; revised date here; accepted date here, Available online date here

المخلص

في هذه الدراسة تم مقارنة وتقييم التطورات الأخيرة في تصميم وتطبيق المحفزات الضوئية Ru / Pd و Ru / Pt لتوليد الهيدروجين عبر العمليات الجزيئية. ومعرفة المفاهيم الأساسية لمثل هذه التمرکزات مع نظرة عامة على العوامل الجزيئية التي تؤثر على إمكاناتها كمحفزات ضوئية. وتم تحديد مدى تمرکز حالة الإثارة الانبعاثية ومعرفة التفاعل بين المركزين المعدنيين بواسطة الكروماتوغرافيا HPLC و التحليل الضوئي والطيفي للمطابقة.

الكلمات المفتاحية:

الروثينيوم/البلاديوم والروثينيوم/البلاتين – المحفزات الضوئية –الكروماتوغرافيا- التحليل الطيفي.

Abstract

In this contribution recent developments in the design and application of Ru/Pd, Ru/Pt photocatalysts for the generation of hydrogen via intramolecular processes are assessed. The basic concepts of such assemblies are discussed together with an overview of the factors and molecular issues that affect their potential as photocatalysts. Issues that so far have limited progress are discussed and suggestions for future directions are made. The extent of localization of the emissive excited state was determined by HPLC chromatography and same wavelengths; however, whereas for Ru/Pd the emission is based on the Ru(tpy)Cl⁻ center, for Ru/Pt the emissive state is localized on the Ru(bpy)₂ unit. This indicates that also in the excited state there is efficient interaction between the two metal centers was characterized by both HPLC, Spectroscopy.

Keywords: Ru/Pd, Ru/Pt, photocatalysts, HPLC, Photolysis, Spectroscopy.

1. INTRODUCTION

Ru-polypyridyl complexes are widely selected as the photosensitizer, while Pt- or Pd-based moieties bound via a bridging ligand are attractive catalytic centers due to their low overpotential for proton reduction¹. As proton reduction into H₂ generation requires the accumulation of two electrons at the catalyst, the common design strategy is based on the bridging ligand functioning as an electron storage reservoir for the first photo-excited electron^{2,3,4,5,6}. The reactivity of such assemblies depends on the structure of the individual components^{7,8,9,10}. Ideally, light-induced electron transfer from the photosensitizer via the bridging ligand to the catalyst should occur with a high quantum yield for efficient H₂ generation. However, this is not always the case. We have

recently observed for a series of Ru/Pd and Ru/Pt photocatalysts that photoexcitation not only leads to population of Franck-Condon states localised on the bridging ligand, but also of states localised on the peripheral ligands^{11,12,13}. A similar effect has been observed by others for a related Ru/Pd complex¹⁴ the synthesis were reported¹⁵.

The characterization of two binuclear complexes of ruthenium Pd, ruthenium and Pt containing both (bpy) (2,2'-bipyridine)

and tpy (2,6-bis(pyridin-2'-yl)-pyridine) as peripheral ligands of the general formula The compound [(tbbipy)₂Ru(tpphz)PdCl₂](PF₆)₂ **1**, where tbbipy = 4,4'-di-*tert*-butyl-2,2'-bipyridine and tpphz = tetrapyrido [3,2-a:2',3'-c:3'',2''-h:2''',3'''-j]phenazine **1**, and Ru/Pt complex with 4,7-diphenyl-1,10-phenanthroline peripheral ligands RuPt **2**. (Figure 1). In compounds **1** and **2** the bridging ligand 3,5-bis(pyridin-2'-yl)-1,2,4-triazolato (bpt) is egatively charged.^{6,7} In contrast to related pseudo-symmetric systems^{7,8}, in complex **1** and complex **2** the non-bridging polypyridyl ligands at each center are geometrically and photophysically distinct, i.e. (bpy)₂ vs (tpy)Cl. The extent and directionality of the interaction between the triazolato bridged metal centers is found to be highly dependent on the nature of the metal center. The corresponding deuteriated isotopologues complex **1** and complex **2** were prepared to facilitate assignment of the chemical shifts of the protons in the bridging (bpt) ligand and (tpy) peripheral ligand as well as the vibrational and electronic spectroscopic features. The results obtained show that in these systems the location of the lowest emissive excited state and the first oxidation can be switched by judicious choice of the metal centers. Importantly this can be done with only minimal perturbation of the photophysical properties including electronic absorption and emission spectra¹⁶.

*Correspondence:

Dr. Hamid M. Younis
Faculty of Science, University of Sirte, Libya.
hamid.younis@su.edu.ly

2. Experimental Section

Material and Methods

All complexes investigated were synthesized by and received from Prof Sven Rau research group Erlangen, Germany. All samples were used as received, with no further purification. All solvents employed in spectroscopic measurements were of spectroscopic grade (Sigma-Aldrich). All other solvents were of HPLC grade or better. $\text{cis-Ru(bpy)}_2\text{Cl}_2 \cdot 2\text{H}_2\text{O}$,¹⁷ $\text{Ru(d8bpy)}_2\text{Cl}_2 \cdot 2\text{H}_2\text{O}$, $\text{Pt(bpy)}_2\text{Cl}_2$ ¹⁸ and Ru(tpy)Cl_3 ,¹⁹ d8-bpy ²⁰ and 3,5-bis(pyridin-2-yl)-1,2,4-triazole (Hbpt)²¹ were prepared by reported procedures. Gold colloid for SERS measurements were prepared by standard methods.²²

Table 1 Spectroscopic and electrochemical data for complexes 1 and 2 and related analogs, solutions purged with Ar at 298 K.

UV/V is absorption and emission spectroscopy:

UV–Vis absorption spectra were recorded on a Shimadzu 3100 UV-Vis instrument with 1-cm quartz cells in spec grade DCM, ethanol or acetonitrile (Aldrich). Emission spectra were collected on a Perkin-Elmer LS50B luminescence spectrometer equipped with a red sensitive Hamamatsu R928 detector using 1-cm quartz 4 sided cell. Emission and excitation slit widths were typically 3, 5 or 10nm depending on individual circumstances.

Electronic absorption and emission data for the mononuclear and dinuclear complexes are listed in Table 1. All complexes exhibit absorption and emission properties that are characteristic of ruthenium(II) and Platinum (II) based polypyridyl complexes with triazolato containing bridging ligands²³.

The absorption spectra for complex 1 and complex 2 are shown in Figure 2. The shoulder at ca. 522 nm is typical for ruthenium (II) (N5) chlorido complexes^{23,24}. The absorptions >570 nm are assigned to ³MLCT transitions in the case of complex 2, however for complex 1 a significant absorption is observed in

complexes		Absorption	Emission
		$\lambda_{\text{max}}/\text{nm}$ ($\epsilon / 10^4\text{M}^{-1}\text{cm}^{-1}$)	$\lambda_{\text{max}}/\text{nm}$
RuPd	1	470 (0.95), 465 (1.55)	755
RuPt	2	481 (0.69), 606 (0.40)	766

this region also which are reminiscent of the absorption spectrum of $[\text{Ru(tpy)}_2]^{2+}$ ²⁵.

In the UV region (< 350 nm) the absorptions are assigned to $\pi-\pi^*$ intra-ligand electronic transitions associated with the (bpy), (tpy) and bridging triazolato ligands,^{26,27} whereas those in the visible region (400–570 nm) are assigned to ¹MLCT (metal ligand charge transfer) transitions²⁸.

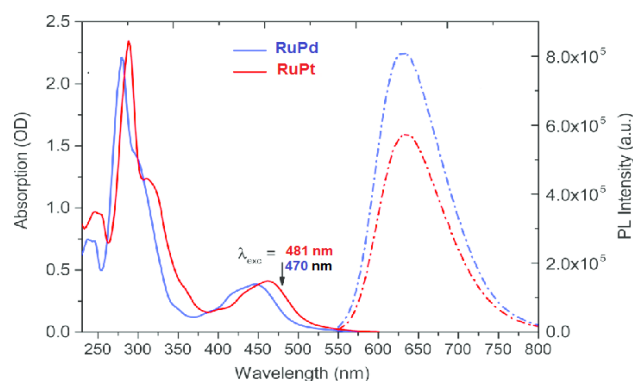


Figure 1. Absorption, Emission spectra photolysis of complex 1 RuPd λ_{max} 470 nm (blue line) and RuPt complex 2 λ_{max} 481 nm (Red line) in CH_3CN .

The emission spectra of both complex 1 and complex 2 show maxima at 755 nm and 766 nm, respectively (Figure 2). In both cases the emission energy and lifetime (Table 1) is typical of emission from a ³MLCT excited state, either localized on a (bpy) or on a (tpy) ligand. The near coincidence in energy of the emission of the two complexes could suggest that the emission is localized at the same metal center for both complexes, i.e. the $[\text{Ru(tpy)Cl}]^+$ center.

However, the differences in emission lifetimes and intensities indicate otherwise. For complex 1, the emissive ³MLCT state is assigned to be localized on the $[\text{Ru(tpy)Cl}]^+$ moiety on the basis that the related complex^{29,30}, $[(\text{tbbipy})_2\text{Ru(tpphz)PdCl}_2](\text{PF}_6)_2$ 1 has an emission maximum at 755 nm (Table 1), the substitution of the (bpy) ligand of $[\text{Ru(tpy)(bpy)Cl}]^+$ with the strongly σ -donating triazolato ligand will raise the energy of the ³MC excited state which is typically the major excited state deactivation channel for ruthenium(II) polypyridyl complexes. Furthermore the relatively higher intensity of emission for complex 2 over complex 1 despite the shorter excited state lifetime (Figure 2) indicates that complex 2 has a shorter radiative lifetime in accordance with a switch from a ruthenium(tpy) to a Platinum(bpy) centered emission between complex 1 and complex 2.

The concentrations were approximately $1\text{--}2 \times 10^{-5}$ M. A direction of energy and optical electron transfer in a heterobimetallic polypyridyl complex is switched without significant changes in electronic absorption or emission properties³¹.

3. High Performance Liquid Chromatography

(HPLC) experiments were carried out using an analytical HPLC system consisting of a Varian Prostar HPLC pump fitted with a 10 μL injection loop, a Varian Prostar PDA detector connected to a dedicated PC, and a HiChrom Partisil P10SCX-3095 cation exchange column. The mobile phase used was acetonitrile : water 80:20 (v:v) containing 0.01–0.04 M LiClO_4 ; the flow rate was 1.0 cm^3/min . The monitoring wavelength used was 280 nm. Photostability of complexes 1 and 2 was monitored by HPLC. Samples were irradiated at 470 nm with a 9 W LED-array.

Photochemical Studies

The photochemical experiments were carried out at room temperature in air-equilibrated solution in a 1 cm quartz, Light sources For illumination, we used LED light emitting diodes $\lambda =$

470 nm LAMP type L-7113PBC-BLUE. (sacrificial agent) and took 5 ml sample of above concentration solution for irradiation at 470nm (blue light) (irradiation time).

¹H-NMR Spectroscopy.

¹H NMR (400 MHz) and ¹³C NMR (100 MHz) spectra were obtained on a Bruker Avance 400 NMR Spectrometer in deuterated solvents with either TMS or residual solvent peaks as reference ³².

The ¹H NMR spectroscopic and mass spectral data for the mononuclear complexes are in agreement with previous reports (28). For the dinuclear complexes the availability of complex **1** and **2** allow for analysis of the proton chemical shifts in (bpt) and (tpy) ligands. The coordination mode of the Ru(tpy)Cl⁻ unit in complex **1** and **2** can, potentially, be such that the chlorido ligand is trans to either the triazolato or the pyridyl ring of the (bpt) bridging ligand (Figure 3).

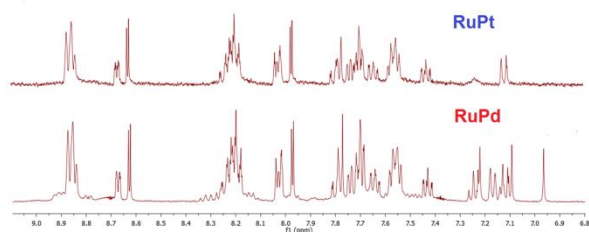


Figure 3. ¹H-NMR of complex **1** RuPd. (bottom) and complex **2** RuPt (top) in dmsO-d₆. The signals arising from the complexes are identical. The additional signals in the spectra of the monomer are due to toluene and ammonium salt impurities.

4. RESULTS AND DISCUSSION

Synthesis. The dinuclear complexes **1** and **2** were prepared by addition of the soluble mononuclear complexes [(tbbipy)₂Ru(tpphz)PdCl₂](PF₆)₂ **1**, and RuPt, respectively to a solution containing the mononuclear complex Ru(tpy)Cl₃ in EtOH/H₂O followed by heating at reflux and isolation by chromatography. (13) the first peak in HPLC at 6.18min splits up into two, corresponding to the unreacted bpy and the intermediate species. In (Figure 4) the peak at retention time 6.77min corresponds to [(tbbipy)₂Ru(tpphz)PtCl₂](PF₆)₂ (**2**).

At up to 120min most of the intermediates get converted into complex **1**, which is evident from HPLC (Figure 4) and the peaks have disappeared on going from 6.18min to 6.77min. up to 6 hours almost all of the bipyridine has reacted and we are left with the final two products [Pt(bpy)₂(bpt) Ru(tpy) PdCl₂]²⁺ complex **1**. (Figure 5).

Figure shows the absorption spectra of Ru and RuPd complex **1** dissolved in acetonitrile and dichloromethane. In acetonitrile both compounds show absorption properties which are common for the class of ruthenium polypyridine compounds. The absorption spectrum of Ru reveals four absorption maxima in the spectral range between 250 and 700 nm: The maximum at 283 nm can be assigned to a ligand-centered π-π* transition of tbbpy. The following maxima at 359 and 382 nm reflect ligand-centered π-π* transitions of tpphz. The energetically lowest lying broad transition which is pumped in our pump-probe experiments at 507 nm is assigned to MLCT transitions between

the ruthenium ion and the coordinated ligands. For RuPd complex **1** a qualitatively similar absorption spectrum with only minor differences is recorded ³³.

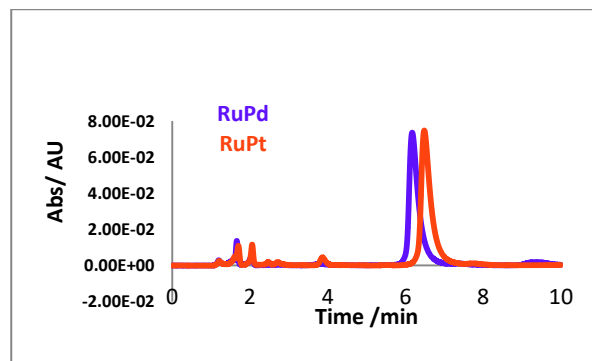


Figure 4. HPLC trace for Ru/Pd complex **1** and Ru/Pt complex **2**, in CH₃CN (mobile phase CH₃CN: H₂O with volume ratio 80:20 containing 0.1 M KNO₃). Flow rate: 1.0 cm³ min⁻¹; detection wavelength at 280 nm.

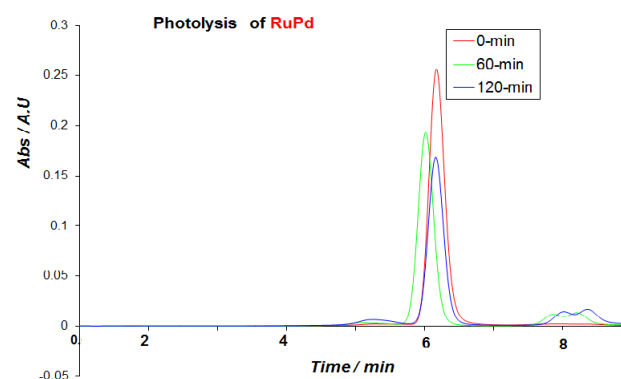


Figure 5. HPLC trace during photolysis of RuPd complex **1** in CH₃CN (mobile phase CH₃CN: H₂O with volume ratio 80:20 containing 0.1 M KNO₃). Flow rate: 1.0 cm³ min⁻¹; detection wavelength at 280 nm

We continued the study up to 120min which showed that most of the metal got charred at this higher temperature resulting in a decrease of yield to 40% at 120min in RuPt complex **2**. In (Figure 6).

Photostability, previously it was shown for complexes of the type [Ru(bpy)₂(L)Cl]⁺ where L = pyridine, (15) 4-vinylimidazole (16) and poly-4-vinyl-pyridine, (15) that rapid loss of a chlorido ligand can be observed by photochemical or thermal activation and photoinduced ligand loss has been observed for [Ru(tpy)(bpy)Cl]⁺ also (17,18). The photostability under visible excitation of [(tbbipy)₂Ru(tpphz)PdCl₂](PF₆)₂ complex **1** and RuPt complex **2**, in acetonitrile was determined by HPLC. Formation of a main photolysis product is detected only on extended photolysis after 90min and 120min irradiation for complex **1** (Figure 5), and complex **2**, (Figure 6), respectively.

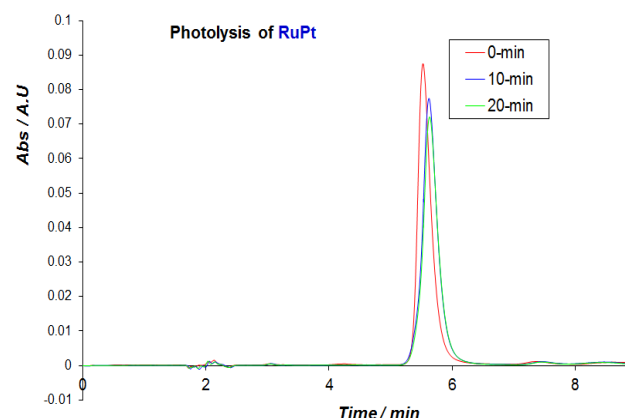


Figure 6. HPLC trace during photolysis of RuPt complex 2 in CH₃CN (mobile phase CH₃CN: H₂O with volume ratio 80:20 containing 0.1 M KNO₃. Flow rate: 1.0 cm³ min⁻¹; detection wavelength at 280 nm.

A similar result was obtained by Pt NMR spectroscopy investigations on related [(bpy)PtX₂] (X=Cl or I) complexes showing an increase in electron density (more negative chemical shift values) at the Pt nucleus induced by the softer iodide ligand compared to chloride (28). In addition, electrochemical investigations were carried out to test whether the increased electron density at the Pt center, induced by the iodide ligands, is also expressed by a corresponding change of the reduction potential of the Pt center. However, no significant difference in the reduction potentials was observed comparing photocatalyst complex 2 and complex 1.

5. CONCLUSIONS

Here, the synthesis, ¹H NMR, absorption, emission, and HPLC, characterization of the dinuclear complexes **1**, and **2**. Have been described together with their deuterated isotopologues. ¹H NMR spectroscopy indicates that only one isomer is isolated for both complexes in which the chlorido ligand is trans to the triazolato ligand. Both complexes **1** and **2** are relatively photostable in contrast to related mononuclear chlorido containing systems. Emission and HPLC indicate that for compounds **1** the emissive excited state is localized on the [Ru(tpy)Cl] center, whereas for **2** the emissive state is localized on the Ru(bpy)₂ unit. HPLC and UV/vis/NMR spectroelectrochemistry

This product manifests itself in the appearance of a peak in the HPLC chromatogram at a retention time longer than observed for complex **1** and complex **2** in (Figures 5 and 6), which indicates that the photoproduct is more positively charged. Comparison of the absorption spectra of complex **1** and complex **2** and the primary photolysis product indicates the loss of the chloride ligand. This indicates that cleavage of the Ru-Cl bond reaction of Ru and Pt metal is unlikely to be significant on the time scale of the absorption, Emission and Raman spectroscopic studies^{34,35}.

We completely succeeded in eliminating quite a numerous side products in the conventional reaction of Ru and Pt metal with (bpy) by employing a higher temperature resulting in separation of only two products at the end of the reaction clearly proved using the HPLC. Note that in these experiments a proton donor was not present and hence stabilisation of such a negative charge by hydrogen bonding would be expected under catalytic conditions. The results presented here demonstrate that the

light-induced processes lead to a reduction of the poth PdII, PtII, center already within the first few nanoseconds of the excited-state lifetime and thus much faster than has been previously suggested in the framework of a catalytic mechanism³⁶.

6. REFERENCES

1. Barber J. Biological solar energy, Philos. Trans. Royal Soc. A 2007, 365: 1007-1023.
2. Esswein AJ, Nocera DG. Hydrogen Productio by Molecular Photocatalysis, Chem. Rev. 2007, 107: 4022-4047.
3. Martis M, Mori K, Kato K, Sankar G, Yamashita H. What Are the Active Species in the Photoinduced H₂ Production with Terpyridyl Pt(II) Complexes. An Investigation by in Situ XAFS, ChemPhysChem 2013, 14:1122-1125.
4. Meyer TJ. Plants produce oxygen from water, but the same chemical reaction is hard to achieve synthetically. A new family of catalysts could breathe fresh life into the quest for artificial photosynthesis.. Nature 2008, 451: 778-779.
5. Balzani V, Credi A, Venturi M. Photochemical conversion of solar energy. Chemsuschem 2008, 1: 26-58.
6. Magnuson A, Anderlund M, Johansson O, Lindblad P, Lomoth R, Polivka T, Ott S, Stensjo K, Styring S, Sundstrom V, Hammarstrom L. Biomimetic and Microbial Approaches to Solar Fuel Generation Acc. Chem. Res. 2009, 42: 1899-1909.
7. Tachibana Y, Vayssieres L, Durrant JR. Artificial photosynthesis for solar water-splitting, Nat. Photonics 2012, 6: 511-518.
8. Juris A, Balzani V, Barigelletti F, Campagna S, Belser P, Vonzelewsky. Absorption and electrochemical properties of ruthenium(II) dyes, studied by semiempirical quantum chemical calculations. A., Coordin. Chem. Rev. 1988, 84: 85-277.
9. Yeh AT, Shank CV, McCusker JK. Ultrafast electron localization dynamics following photo-induced charge transfer. Science 2000, 289: 935-938.
10. Cannizzo A, van Mourik F, Gawelda W, Zgrablic G, Bressler C, Chergui M. Broadband femtosecond fluorescence spectroscopy of [Ru(bpy)₃]²⁺. Angew. Chem. Int. Ed. 2006, 45: 3174-3176.
11. Campagna S, Puntoriero F, Nastasi F, Bergamini G, Balzani V. Photochemistry and Photophysics of Coordination Compounds I Top. Curr. Chem. 2007, 280: 117-214.
12. Ozawa H, Haga MA, Sakai, K. A Photo-Hydrogen-Evolving Molecular Device Driving Visible-Light-Induced EDTA-Reduction of Water into Molecular Hydrogen, J. Am. Chem. Soc. 2006, 128: 4926-4927.
13. Rau S, Schafer B, Gleich D, Anders E, Rudolph M, Friedrich M, Gorgs H, Henry W, Vos JG. A supramolecular photocatalyst for the production of hydrogen and the selective hydrogenation of toluene, Angew. Chem. Int. Ed. 2006, 45: 6215-6218.
14. Morales-Guio CG, Stern LA, Hu XL. Nanostructured hydrotreating catalysts for electrochemical hydrogen evolution, Chem. Soc. Rev 2014, 43: 6555-6569.
15. Karnahl M, Kuhnt C, Ma F, Yartsev A, Schmitt M, Dietzek B, Rau S, Popp J. Tuning of Photocatalytic Hydrogen Production and Photoinduced Intramolecular Electron

- Transfer Rates by Regioselective Bridging Ligand Substitution. *Chemphyschem* 2011, 12: 2101-2109.
16. Bindra GS, Schulz M, Paul A, Groarke R, Soman S, Inglis J. L., Browne WR, Pfeffer MG, Rau S, MacLean BJ, Pryce MT, Vos JG. The role of bridging ligand in hydrogen generation by photocatalytic Ru/Pd assemblies, *Dalton Trans.* 2012, 41: 13050-13059.
 17. Bindra GS, Schulz M, Paul A, Soman S, Groarke R, Inglis J, Pryce MT, Browne WR, Rau S, Maclean BJ, Vos JG. The effect of peripheral bipyridine ligands on the photocatalytic hydrogen production activity of Ru/Pd catalysts, *Dalton Trans.* 2011, 40: 10812-10814.
 18. Michael G Pfeffer., Wächter M, Guthmüller J, Dietzek B, Vos JG, Rau S., Optimization of hydrogen-evolving photochemical molecular devices., *Angew Chem Int Ed Engl*, 2015., 54: 6627-6631.
 19. Knoll JD, Higgins SH, White TA, Brewer KJ. Subunit Variation to Uncover Properties of Polyazine-Bridged Ru(II), Pt(II) Supramolecules with Low Lying Charge Separated States Providing Insight into the Functioning as H₂O Reduction Photocatalysts to Produce H₂, *Inorg. Chem.* 2013, 52: 9749-9760.
 20. Kowacs T, Pan Q, Lang P, O'Reilly L, Rau S, Browne WR, Pryce MT, Huijser A, Vos JG. Supramolecular Bimetallic Assemblies for Photocatalytic Hydrogen Generation from Water, *Faraday Discuss.* 2015, 185: 143-170.
 21. Pan Q, Mecozzi F, Korterik JP, Sharma D, Herek JL, Vos JG, Browne WR, Huijser AJ. The Critical Role Played by the Catalytic Moiety in the Early-Time Photodynamics of Hydrogen-Generating Bimetallic Photocatalysts, *ChemPhys.Chem.* 2014, 118: 20799-20806.
 22. Tschierlei S, Karnahl M, Presselt M, Dietzek B, Guthmüller J, Gonzalez L, Schmitt M, Rau S., Popp, J. Photochemical fate: the first step determines efficiency of H₂ formation with a supramolecular photocatalyst, *Angew. Chem. Int. Ed.* 2010, 49: 3981-3984.
 23. Huey F, Suen, Wilson SW. Poimerantz, M. and Walsh, J. L., Photosubstitution reactions of terpyridine complexes of ruthenium(II), *Inorg. Chem.* 1989, 28: 786-791.
 24. Patra S, Sarkar B, Ghumaan S, Patil MP, Mobin SM, Sunoj RB, Kaim W, Lahiri GK. Isomeric ruthenium terpyridine complexes [Ru(tpy)(L)Cl]ⁿ⁺ containing the unsymmetrically bidentate acceptor L = 3-amino-6-(3,5-dimethylpyrazol-1-yl)-1,2,4,5-tetrazine. Synthesis, structures, electrochemistry, spectroscopy and DFT calculations, *Dalton Trans.* 2005, 1188-1194.
 25. Rau, S., Schafer, B., Gleich, D., Anders, E., Rudolph, M., Friedrich, M., Gçrls, H., Herny, W., Vos, J. G., A supramolecular photocatalyst for the production of hydrogen and the selective hydrogenation of tolane, *Angew. Chem.* 2006, 118: 6361 –6364; *Angew. Chem. Int. Ed.* 2006, 45: 6215 –6218.
 26. Halpin Y, Dini D, Ahmed HMY, Cassidy L, Browne WR, Vos JG. Excited State Localization and Internuclear Interactions in Asymmetric Ruthenium(II) and Osmium(II) bpy/tpy Based Dinuclear Compounds, *Inorg. Chem.* 2010, 49: 2799–2807.
 27. Pan TK, Q, Lang P, O'Reilly L, Sven R., Browne WR, Mary T, Pryce MT, Jannetje M. Huijser A, Vos JG, Supramolecular bimetallic assemblies for photocatalytic hydrogen generation from water, *Faraday Discuss.*, 2015, 185: 143-170.
 28. Knoll, J. D., Higgins, S. H., White, T. A., Brewer, K. J., Subunit Variation to Uncover Properties of Polyazine-Bridged Ru(II), Pt(II) Supramolecules with Low Lying Charge Separated States Providing Insight into the Functioning as H₂O Reduction Photocatalysts to Produce H₂, *Inorg. Chem.* 2013, 52: 9749-9760.
 29. Pfeffer MG, Kowacs T, Chtler MW, Guthmüller J, Dietzek B. Optimization of Hydrogen Evolving Photochemical Molecular Devices, *Angew. Chem. Int. Ed.* 2015, 54: 6627 –6631.
 30. Pfeffer MG, Schafer B, Smolentsev G, Uhlig J, Nazarenko E, Guthmüller J, Kuhnt C, Wächter M, Dietzek B, Sundstrom V, Rau S. Palladium versus platinum: the metal in the catalytic center of a molecular photocatalyst determines the mechanism of the hydrogen production with visible light., *Angew. Chem. Int. Ed.* 2015, 54, 5044-5048.
 31. Tschierlei S, Presselt M, Kuhnt C, Yartsev A, Pascher T, Sundström V, Karnahl M, Schwalbe M, Schafer B, Rau S, Schmitt M, Dietzek B, Popp J. Photophysics of an Intramolecular Hydrogen Evolving Ru–Pd Photocatalyst, *Chem. Eur. J.* 2009, 15: 7678.
 32. Pan Q, Freitag L, Kowacs T, Falgenhauer JC, Korterik JP, Schlettwein D, Browne WR, Pryce MT, Rau S, González L, Vos JG, Huijser A. Peripheral ligands as electron storage reservoirs and their role in enhancement of photocatalytic hydrogen generation, *Chem. Commun*, 2016, 52: 9371.
 33. O'Reilly L, Pan Q, Das N, Wenderich K, Korterik JP, Vos, J G, Pryce M, Huijser A. Hydrogen-Generating Ru/Pt Bimetallic Photocatalysts Based on Phenyl-Phenanthroline Peripheral Ligands., *Chemphyschem.* 2018, 22: 3084-3091.
 34. Kowacs T, Pan Q, Lang P, O'Reilly L. Sven Rau, Browne, W. R., Pryce, MT Huijser A, Vos JG. Supramolecular bimetallic assemblies for photocatalytic hydrogen generation from water, *Faraday Discuss.*, 2015, 185: 143.
 35. Cao J, Zhou Y. Excited state relaxation processes of H₂-evolving Ru–Pd supramolecular photocatalysts containing a linear or non-linear bridge: a DFT and TDDFT study, *Chem Phys.* 2017, 18: 11529-11539.
 36. Karnahl M, Kuhnt C, Ma F, Yartsev A, Schmitt, M, Dietzek B, Rau S. Tuning of Photocatalytic Hydrogen Production and Photoinduced Intramolecular Electron Transfer Rates by Regioselective Bridging Ligand Substitution, *ChemPhysChem* 2011, 2101-2109.