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[View Journal](#) | [View Issue](#)Cite this: *Dalton Trans.*, 2023, **52**, 169741,2-Azolyamidino ruthenium(II) complexes with DMSO ligands: electro- and photocatalysts for CO₂ reduction†Murphy Jennings,^a Elena Cuéllar,^b Ariadna Rojo,^b Sergio Ferrero,^b Gabriel García-Herbosa,^c John Nganga,^d Alfredo M. Angeles-Boza,^{a,d} Jose M. Martín-Alvarez,^b Daniel Miguel^b and Fernando Villafañe^{*b}

New 1,2-azolyamidino complexes *fac*-[RuCl(DMSO)₃(NH=C(R)az*-κ²N,N)]OTf [R = Me (**2**), Ph (**3**); az* = pz (pyrazolyl, **a**), indz (indazolyl, **b**)] are synthesized *via* chloride abstraction from their corresponding precursors *cis*,*fac*-[RuCl₂(DMSO)₃(az*H)] (**1**) after subsequent base-catalyzed coupling of the appropriate nitrile with the 1,2-azole previously coordinated. All the compounds are characterized by ¹H NMR, ¹³C NMR and IR spectroscopy. Those derived from MeCN are also characterized by X-ray diffraction. Electrochemical studies showed several reduction waves in the range of −1.5 to −3 V. The electrochemical behavior in CO₂ media is consistent with CO₂ electrocatalytic reduction. The catalytic activity expressed as [*i*_{cat}(CO₂)/*i*_p(Ar)] ranged from 1.7 to 3.7 for the 1,2-azolyamidino complexes at voltages of *ca.* −2.7 to −3 V vs. ferrocene/ferrocenium. Controlled potential electrolysis showed rapid decomposition of the Ru catalysts. Photocatalytic CO₂ reduction experiments using compounds **1b**, **2b** and **3b** carried out in a CO₂-saturated MeCN/TEOA (4:1 v/v) solution containing a mixture of the catalyst and [Ru(bipy)₃]²⁺ as the photosensitizer under continuous irradiation (light intensity of 150 mW cm^{−2} at 25 °C, λ > 300 nm) show that compounds **1b**, **2b** and **3b** allowed CO₂ reduction catalysis, producing CO and trace amounts of formate. The combined turnover number for the production of formate and CO is *ca.* 100 after 8 h and follows the order **1b** < **2b** ≈ **3b**.

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Introduction

Research on the catalytic reduction of CO₂ has been receiving intense attention since it is a key component to a sustainable future. A variety of transition metal complexes have been reported to display electrochemical CO₂ reduction activities;

however, novel catalytic systems are still required to cope with the social dilemma of climate change.¹ Ruthenium(II) complexes are among those that are more studied, and in fact the electrochemical CO₂ reductions catalyzed by *cis*-[Ru(bipy)₂(CO)₂]²⁺ (bipy = 2,2'-bipyridine) and *cis*-[Ru(bipy)₂(CO)Cl]⁺ were among the first systems reported, back in 1987.² Since then, a plethora of Ru catalysts have been described in the context of CO₂ electroreduction, most of them containing two bipy ligands coordinated *cis*, or other polypyridine-type ligands with similar geometries.^{1–3} The use of mono(bipy) complexes is limited to carbonyl complexes such as [Ru(N-N)(CO)₂Cl₂] (N–N = bipy or substituted bipy) or cationic derivatives derived from the substitution of the chlorido ligands by neutral ligands, and to their corresponding reduced Ru(0) species.⁴

Reduction of carbon dioxide can also be achieved by photochemical methods. The catalysts for these processes are usually metal complexes with different accessible redox states available in both the central metal and the ligands in order to drive the multi-electron reduction process for CO₂ reduction.⁵ In the case of Ru(II), both bis(bipy)⁶ and mono(bipy)⁷ complexes have been widely studied as photocatalysts for CO₂

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† Electronic supplementary information (ESI) available: Cyclic voltammograms of complexes 1–3, solution stability tests, TEM of **2b** degradation products, turnover numbers for the production of H₂, CO, and HCCOH at different times for compounds **1b**, **2b**, and **3b**, isotope labeling experiments for clarification of carbon source of CO production, ¹H NMR spectra of complexes 1–3. CCDC 2221963 and 2221964 crystal structures of complexes **2a** and **2b**. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3dt01122d>

reduction to give CO and/or formate, depending on the reaction conditions.

The activity of the catalyst largely depends on the substituents of the ligands, which are key factors influencing both the primary coordination sphere of the transition metal center and the secondary coordination effects. The latter have been taken into consideration only very recently, when the first reports on the role of pendant groups that may lead to supramolecular arrangements have been described.⁸ However, synthetic methods for substituted bipy ligands are usually difficult and/or tiresome, and thus the alternative of coupling two monodentate ligands to afford a chelating diimine ligand appears as a straightforward method in order to obtain N,N-chelating ligands by accessible synthetic paths.⁹ 1,2-Azolyamidino ligands fulfill these requirements, as they can be easily obtained *in situ* by the coupling reaction of 1,2-azoles and coordinated nitriles (Scheme 1). We have exploited this reaction, *i.e.* the metal-mediated coupling of 1,2-azoles and nitriles, so that a wide range of new 1,2-azolyamidino complexes can be thus easily obtained.¹⁰ The use of different 1,2-azoles and nitriles provides the opportunity of controlling the main features, electronic and steric, of the ligand. In particular, the electrochemical and luminescence properties of Re(I) tricarbonyl and *cis*-bis(bipyridyl)Ru(II) complexes containing 1,2-azolyamidino ligands, as well as the photocatalytic behaviour of the latter, have been reported.¹¹ The same reaction using a nucleobase such as 1-methylcytosine instead of the 1,2-azole has been revealed to be a new method to incorporate biologically relevant substrates into Re(I) tricarbonyl complexes.¹²

Another aspect of interest in the 1,2-azolyamidino ligands lies in the presence of an acidic NH group, which allows further significative reactivity, as it may be involved in noncovalent interactions. A relevant example is the ability of the *fac*-[Re(CO)₃(Hdmpz)(HN=C(Me)dmpz-κ²N,N)]⁺ cation to bind selectively with chloride anions by combining electrostatic attraction and hydrogen bonding of the NH groups present.¹³ A second exponent, this time on Mn(I), emerges from the use of a somehow uncommon nitrile, such as dicyanamide, which allowed a bimetallic Mn(I) complex containing a bridging tetradentate bis(pyrazolylamidino) ligand to be obtained, where the bromide anion plays a crucial role in its planarity, as demonstrated by DFT calculations.¹⁴

Herein, we report the synthesis and characterization of new 1,2-azolyamidino Ru(II) complexes. Their chlorido(1,2-azole) precursors are also described. The 1,2-azoles used in this work

are pyrazole (pzH) and indazole (indzH), whereas the nitriles are acetonitrile (MeCN) and benzonitrile (PhCN). This allows us to study easily obtained 1,2-azolyamidino ligands with different electronic and steric properties. The behavior of the complexes synthesized as electro- and photocatalysts for the reduction of CO₂ is described.

Results and discussion

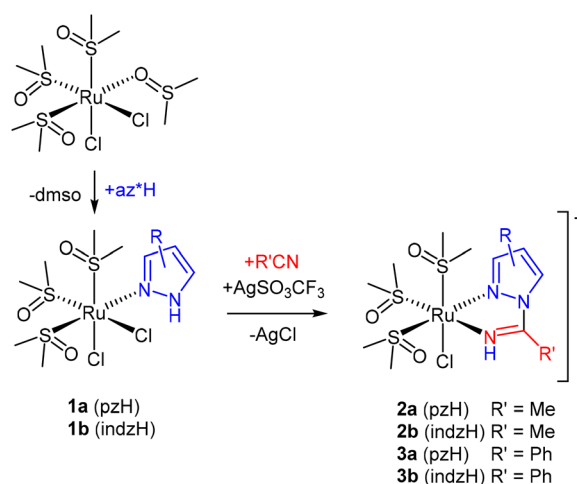
Synthesis and characterization of the complexes

The synthesis of the complexes described in this work is depicted in Scheme 2. A panel of complexes with different substituents were synthesized in order to confirm the synthetic method and to study the influence of the substituents on the electrocatalytic reduction of CO₂. Scheme 2 shows the mixed chlorido(1,2-azole) complexes *cis, fac*-[RuCl₂(DMSO)₃(az*H)] (1) (az* = pz, indz), which are precursors of the 1,2-azolyamidino *fac*-[RuCl(DMSO)₃(NH=C(R)az*-κ²N,N)]OTf (R = Me, Ph) complexes, as a result of coupling a coordinated pyrazole (pzH) or indazole (indzH) in 1 with acetonitrile (R = Me, 2) or benzonitrile (R = Ph, 3) (Scheme 1). Complexes 1a and 1b had been previously reported by Ferrer *et al.*¹⁵ and by Reisner *et al.*,¹⁶ respectively. Herein, we report a new synthetic procedure and its thorough characterization.

Chlorido(1,2-azole) complexes 1a and 1b were obtained by substituting the *O*-coordinated DMSO ligand in the parent complex *cis*-[RuCl₂(DMSO)₄] by pyrazole or indazole, respectively (Scheme 2). These synthetic methods have been improved compared to those previously described in the literature. Thus, the pyrazole complex (1a) is herein obtained at room temperature, whereas Ferrer *et al.*¹⁵ used refluxing methanol. Moreover, our reaction leading to the indazole complex 1b lasts for one hour at room temperature, whereas Reisner *et al.*¹⁶ reported an overnight reaction. The 1,2-azolyamidino complexes 2 and 3 were obtained by extracting with silver triflate one of the chlorido ligands in the presence of the nitrile,



Scheme 1 Synthesis of a 1,2-azolyamidino ligand from the (1,2-azole) precursor containing a coordinated nitrile.



Scheme 2 Synthesis of the new complexes.



followed by the coupling of nitrile and the 1,2-azole using NaOH (aq.) as catalyst, as we have previously described.^{10c}

The spectroscopic data are straightforward and support the proposed geometries (see the Experimental section). Complexes **2a** and **2b** were also characterized by single-crystal X-ray diffraction studies (Fig. 1). The distances and angles (CCDC 2221963 and 2221964†) are similar to those found in other similar Ru(II) ligands.¹⁷ Thus, the C(4)–N(3) distances

(1.265(4) Å and 1.279(4) Å, respectively, for **2a** and **2b**) are typical of C=N double bonds. In complexes **2a** and **2b**, the N-bound hydrogens of the 1,2-azolyamidino ligands are involved in hydrogen bonding with an oxygen atom of a triflate anion. The distances and angles detected for **2a** (H(3)⋯O(4) 2.050 Å, N(3)⋯O(4) 2.885 Å, N(3)–H(3)⋯O(4) 163.4°) and **2b** (H(3)⋯O(4) 2.006 Å, N(3)⋯O(4) 2.840 Å, N(3)–H(3)⋯O(4) 162.8°) may be considered as indicative of “moderate” hydrogen bonds.¹⁸

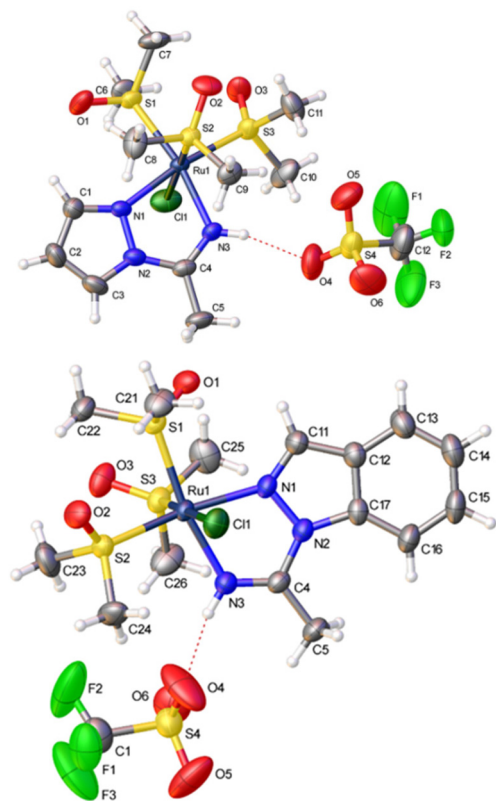


Fig. 1 Perspective views of *fac*-[RuCl(DMSO)₃(NH=C(Me)pz-κ²N,N)](OTf), **2a** (above), and *fac*-[RuCl(DMSO)₃(NH=C(Me)indz-κ²N,N)](OTf), **2b** (below), showing the atom numbering. Thermal ellipsoids are drawn at 50% probability.

Electrochemical studies

Table 1 gathers the observed potential values obtained from cyclic voltammetry (CV) experiments, referenced to the redox pair of ferrocenium/ferrocene, following the IUPAC recommendations.¹⁹ An AgCl/Ag (3 M NaCl) reference electrode was used, and ferrocene was added as internal calibrant always in the last experimental measurement. All the measurements were made in MeCN, once the stability of the complexes in this solvent was confirmed by NMR.

On scanning to negative potentials under N₂, all the complexes display several waves, as the result of successive electron transfer reductions. This is probably the result of pyrazolylamidino ligand reduction, which occurs in Ru^{II}(bipy) systems, which display one or several waves at negative potentials that have always been assigned to bipy-based reductions.^{3b,6f} When the same scans were carried out under a CO₂ atmosphere, the indazole complexes and the pyrazole complex **2a** showed a clear enhancement of the cathodic current (red lines in Fig. S1, ESI†). This electrochemical behaviour is consistent with CO₂ activation, *i.e.* electrocatalyzed reduction. The *i*_{cat}(CO₂)/*i*_p(N₂) ratio allows comparing the electrochemical activity of the complexes, with values ranging from 1.7 to 3.7 for the 1,2-azolyamidino complexes (Table 1). As a representative example, the results registered for the complex *fac*-[RuCl(DMSO)₃(NH=C(Ph)indz-κ²N,N)](OTf) (**3b**) are shown in Fig. 2. Black (N₂) and red (CO₂) traces overlap completely in the range of −1.25 to 0.0 V. Changing the atmosphere from N₂ to CO₂ generates a great enhancement of the current at potentials below −2.5 V. The rest of the experiments under the CO₂

Table 1 Electrochemical data obtained by cyclic voltammetry in this study, in MeCN and Bu₄PF₆ supporting electrolyte, and referenced to the redox system of ferrocenium/ferrocene^a

Complex	Observed <i>E</i> _{pc} values ^b (cathodic scan)	<i>i</i> _p (N ₂) ^c (at <i>E</i> _{pc})	<i>i</i> _{cat} (CO ₂) ^c (at <i>E</i> _{pc})	Ratio ^d $\frac{i_{\text{cat}}(\text{CO}_2)}{i_p(\text{N}_2)}$
1a	−2.26, −3.00 ^e	45.7 (−3.00)	79.4 (−3.10)	1.74
1b	−2.20, −2.95	34.3 (−2.95)	68.3 (−2.83)	1.99
2a	−2.11, −2.42, −3.0	27.7 (−3.00)	51.3 (−3.10)	1.85
2b	−1.93, −2.60, −3.03	27.1 (−3.03)	73.5 (−2.88)	2.71
3a	−2.10, −2.41, −2.93 ^e	38.1 (−2.93)	65.0 (−3.10)	1.71
3b	−1.93, −2.30, −2.80	53.1 (−2.80)	195 (−2.88)	3.67

^a The reduction potential mean value observed for ferrocenium/ferrocene (Fc⁺/Fc) used as an internal calibrant under the employed experimental conditions was *E*^o = 0.443 ± 0.005 V vs. the AgCl/Ag (3 M NaCl) electrode. ^b Cathodic scan peaks observed under N₂ unless stated otherwise.

^c Maximum registered cathodic current (μA) under N₂, *i*_p(N₂), or under CO₂, *i*_{cat}(CO₂) taken from the peak showing greatest enhancement with CO₂ addition. ^d Ratio between the faradaic currents observed under N₂, *i*_p(N₂), or under CO₂, *i*_{cat}(CO₂). ^e Waves where both peaks *i*_{ox} and *i*_{red} were observed. The value of *E*_{1/2} is given in those cases.



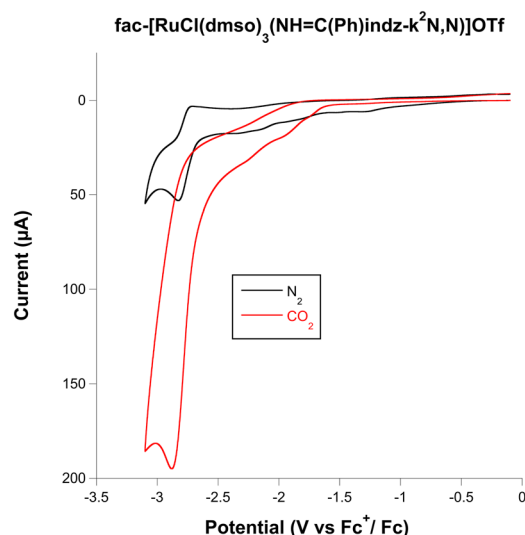


Fig. 2 Cyclic voltammograms of 0.5 mM $\text{fac}[\text{RuCl}(\text{DMSO})_3(\text{NH}=\text{C}(\text{Ph})\text{indz-}k^2\text{N,N})]\text{OTf}$ (**3b**) (glassy carbon working electrode dish 3.0 mm diameter, dry MeCN, 0.1 M Bu_4NPF_6) under N_2 (black) and after bubbling CO_2 (red).

atmosphere lead to different shapes of waves associated with the electrocatalytic reduction of CO_2 due to the competition at the electrode surface between CO_2 consumption (related to the rate-determining step of the catalytic cycle) and the arrival of new substrate by diffusion.^{1c}

The catalytic activity of these complexes is supported by the observed decrease of catalytic activity when the concentration of CO_2 is progressively substituted by bubbling Ar (Fig. S13, ESI†) and by the dependence of the electrocatalytic reduction of CO_2 on the scan rate (Fig. S2–4, ESI†) and on the concentration (Fig. S14, ESI†).

As indicated in the Introduction, there are no previous reports on the catalytic activity on CO_2 reduction for complexes structurally similar to those herein described. The closest analogues might be mono(bipy)Ru(II) complexes containing two carbonyls and two chlorido ligands, which have been used both as electro- and photocatalysts for CO_2 reduction.^{4,7} The $i_{\text{cat}}(\text{CO}_2)/i_{\text{p}}(\text{N}_2)$ ratios obtained for compounds **2** and **3** (between 1.71 and 3.67, see Table 1) are slightly below those obtained by our group for *cis*-Ru(bipy)₂ complexes and for *fac*-Re(CO)₃ complexes (between 2.1 and 10.8 and between 2.7 and 11.5, respectively).^{20,21} Although the reaction mechanism is still being discussed after more than 20 years since the first studies, there is a general consensus about the formation of reduced species that would be highly active for the reduction of CO_2 . These reduced species may contain $\text{Ru}^{\text{I}}\text{--Ru}^{\text{I}}$ or $\text{Ru}^0\text{--Ru}^0$ bonds for the less hindered diimine ligands or $\text{Ru}\text{--H}$ bonds for those containing sterically demanding substituents.^{4,7} In our case, all the attempts to reduce chemically either 1,2-azole (**1**) or 1,2-azoylamidino (**2**, **3**) complexes led to decomposition. The possibility of Ru(0) nanoparticles being active catalysts was tested in a separate experiment and is discussed below.

In order to further evaluate the electrocatalytic activity, we performed a sequence of experiments on the indz complexes (**1b**, **2b**, and **3b**). We focused our study on these complexes since they exhibit higher $i_{\text{cat}}(\text{CO}_2)/i_{\text{p}}(\text{N}_2)$ ratios and less negative potentials for the first reduction by *ca.* 0.2 V, which we can attribute to the lower energy π system on the indz fragment. First, scan rate experiments were performed on each reduction step in order to determine their redox behaviour (Fig. S2–S4, ESI†). These studies showed the general instability of these complexes under catalytic conditions. Additional experiments were run either in MeCN saturated with Cl^- (TBACl, Fig. S6, ESI†) or in DMSO (Table S1, ESI†) as solvent instead of MeCN. In all cases, the CVs point to the formation of species that show nonlinear responses to the scan rate experiments, showing the generation of non-freely diffusing redox species, which leads to the complexity seen in the CVs. All systems exhibited no reversibility, likely due to the instability of the species formed.

Controlled potential electrolysis (CPE) studies on indz complexes **1b**, **2b** and **3b** were also performed following a procedure previously described.²⁰ The experiments, carried out in MeCN using a three-electrode setup with a glassy carbon working electrode, were run at a potential 0.2 V lower than the electron addition, showing the greatest enhancement with the addition of CO_2 , as previously determined by CV for each indz complex. Each experiment was carried out for 6 h, while the MeCN solutions were maintained under a CO_2 atmosphere and stirred. Samples of the solutions were pulled and tested with CV at 0 h, 2 h, 4 h, and 6 h (Fig. S7–S9, ESI†) under a CO_2 atmosphere, bubbled with N_2 for 15 min, and the scans were then repeated. The voltammograms indicate that the electrochemical activity is not lost as the reaction progresses. However, an increase in current is detected under a N_2 atmosphere, indicating that a non-catalytical species that is accepting electrons is being formed (Fig. S7–S9, ESI†). After the 6 h CPE experiment, Hg was added to the reaction mixture, which was then stirred for 12 h more at r.t. Hg is known to bind to Ru nanoparticles from the solutions, removing them as active species.²² No observable changes to the activity behaviour of the complexes under CO_2 were detected after the addition of Hg, suggesting that the activity of the complexes is not due to the formation of nanoparticles. HPLC experiments were also performed before and after the experiment outlined above (Fig. S11, ESI†) in order to further probe the solution stability of the indz family of complexes. The chromatograms taken at 254 nm indicate that the majority of each complex is degraded to different degradation products by the end of the experiment.

In order to test the hypothesis that the rapid formation of degradation products, including Ru(0) nanoparticles, could have been on the electrode surface, CPE experiments were run for 2 h, after which CV experiments were performed, first with the electrode “as is”, and then after cleaning the electrode with a low lint delicate task wipe saturated with dry MeCN (Fig. S10, ESI†). These experiments clearly show that a catalytically active film forms on the surface of the electrode, likely containing Ru(0) nanoparticles and possibly other Ru degra-



dation products. This might be the cause of the less efficient catalytic abilities detected in the bulk-solution tests discussed above since rapid decomposition of the Ru complexes at the surface of the CPE electrode could be the cause behind their lack of activity. This degradation process was also examined by sonicating the working electrode with EtOH immediately after the 2 h CPE experiment in order to remove any solid species deposited on it. These EtOH mixtures were then drop-cast for TEM samples. The TEM images show a variety of electron-rich disordered clusters, thus supporting the hypothesis that Ru(0) particles are deposited on the electrode (Fig. S12, ESI†).

Photocatalytic activity

Given the electrochemical instability of the complexes, we turned our attention to the photocatalytic activity of these complexes. Although some complexes can act as photocatalysts and photosensitizers, this is not the case for **1b**, **2b** and **3b**. Most Ru-based photocatalysts require the assistance of photosensitizers since their direct excitation leads to ligand dissociation.²³ Shown in Fig. 3 are the results of the photocatalytic CO₂ reduction experiments using compounds **1b**, **2b** and **3b** carried out in a CO₂-saturated acetonitrile/triethanolamine (MeCN/TEOA, 4 : 1 v/v) solution containing a mixture of the catalyst and [Ru(bipy)₃]²⁺ as the photosensitizer in a glass vial with a volume of 10 mL under continuous irradiation (light intensity of 150 mW cm⁻² at 25 °C, λ > 300 nm). To validate the photocatalytic data, various control experiments were carried out under different experimental conditions under irradiation with light. In the absence of [Ru(bipy)₃]²⁺, the catalyst, or the sacrificial electron donor, TEOA, only a trace amount or no amount of product was produced, indicating that all three components are necessary for efficient CO₂ activation. The formic acid produced was quantified using the pro-

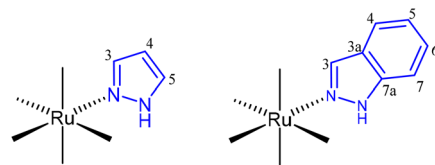


Fig. 4 Numbering of pyrazole and indazole ligands for NMR assignment.

toloc reported by Kubiak *et al.*²⁴ The results show that compounds **1b**, **2b** and **3b** are efficient CO₂ reduction catalysts under the conditions used (Table S1, ESI†), although they also produce larger quantities of dihydrogen and trace amounts of formate.^{5d} The source of protons is probably TEOA. The turnover number for the production of CO is ~100 after 8 h and follows the order **1b** > **2b** ≈ **3b**. Other complexes of the type *trans*-(Cl)-[Ru(bipy*)(CO)₂Cl₂] (where bipy* is a substituted bipy) have shown TON from *ca.* 100 to *ca.* 3000.⁷

Due to the higher photochemical activity of these complexes, a series of additional experiments were conducted, trying to establish the stability of the catalysts and to further understand the process. First, the possibility of the presence of Ru(0) nanoparticles as the active catalysts in the solution was tested by adding 3 equivalents of Hg to the reaction solutions, and checking for changes in product formation. These experiments (Fig. S7–S9, ESI†) did not show significant differences with respect to those obtained with Hg-free solutions. This result led to the conclusion that Ru(0) nanoparticles are not active catalysts in this system.

The photochemical activity of **1b** was also used to probe the carbon source of the complexes by analyzing the product formation of the CO₂ catalysis by running the photocatalysis with a solution saturated with ¹³CO₂ and analyzing the products by gas chromatography (GC), as well as by mass spectrometry coupled GC (Tables S4 and S5, ESI†). All the carbon monoxide detected was in the form of ¹³CO, showing that degradation of the complexes is not a product source; in fact, the source of CO for this reaction is the catalysis of CO₂ to CO by the complexes.

Conclusions

Base-catalyzed coupling of a nitrile and a 1,2-azole previously coordinated to Ru(II) DMSO complexes allows the synthesis of new 1,2-azolyamidino complexes *fac*-[RuCl(DMSO)₃(NH=C(R)az*-κ²N,N)]OTf (R = Me, Ph; az* = pz, indz) after chloride abstraction. The 1,2-azolyamidino complexes and the parent 1,2-azole complexes are active both as electro- and photocatalysts for the reduction of CO₂, as demonstrated by proof-of-concept trials that led to the production of CO and trace amounts of formate. These results open the door to new complexes displaying this interesting behaviour since, so far, the use of Ru(II) mono(bipy) complexes was limited to carbonyl complexes of the type [Ru(bipy)(CO)₂X₂]. The fact that 1,2-azolyamidino ligands with different electronic and steric features

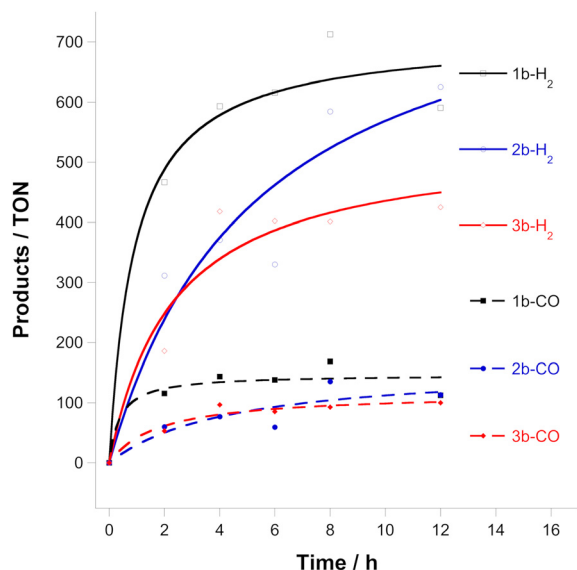


Fig. 3 Turnover numbers for CO and H₂ evolved from 0.1 mM Ru complexes, 1.6 mM [Ru(bipy)₃]²⁺ in a CO₂-saturated MeCN-TEOA solution (4 : 1 v/v) irradiated by >300 nm visible light.



may be easily obtained *in situ* may encourage future developments in this area, given the decisive role of the presence of substituents of the diimine ligand on the electrocatalytic reduction of CO₂.

Experimental section

General remarks

All manipulations were performed under an N₂ atmosphere following conventional Schlenk techniques. Solvents were purified according to standard procedures. *cis*-[Ru(DMSO)₄Cl₂]²⁵ was obtained as previously described. All other reagents were obtained from the usual commercial suppliers and used as received. Infrared spectra were recorded from solids using a Bruker Tensor 27 FTIR. Standard abbreviations are used to indicate intensity: vw = very weak, w = weak, m = medium, s = strong, and vf = very strong. NMR spectra were recorded on 500 MHz Agilent DD2 and 400 MHz Agilent MR instruments in the Laboratory of Instrumental Techniques (LTI) Research Facilities, University of Valladolid, using CDCl₃ or (CD₃)₂CO as solvents at room temperature (r.t.). ¹H and ¹³C NMR chemical shifts (δ) are reported in parts per million (ppm) and are referenced to tetramethylsilane (TMS), using the residual solvent peak as an internal reference. Coupling constants (*J*) are reported in Hz. Standard abbreviations are used to indicate multiplicity: s = singlet, d = doublet, ddd = doublet of doublet of doublets, dt = doublet of triplets, t = triplet, and m = multiplet. The full assignment of the ¹H NMR spectra was supported by typical homonuclear ¹H–¹H correlations such as COSY, TOCSY and NOESY experiments and the assignment of ¹³C{¹H} data was supported by HMBC and HSQC heteronuclear experiments. Numbering of pyrazole and indazole ligands for NMR assignments is shown in Fig. 4. Elemental analyses were performed on a Thermo Fisher Scientific EA Flash 2000.

***cis*-fac-[RuCl₂(DMSO)₃(pzH)], 1a.** This complex was prepared by a modification of the method previously described in the literature. Some spectroscopic data were also previously reported.¹⁵ A mixture of *cis*-[RuCl₂(DMSO)₄] (0.097 g, 0.2 mmol) and pzH (0.014 g, 0.2 mmol) in MeOH (10 mL) was stirred at r.t. for 3 h. The clear yellow solution was left to stand at –20 °C, giving a yellow microcrystalline solid, which was decanted, washed with Et₂O (2 × 5 mL approximately), and dried *in vacuo*, yielding 0.070 g, 74%. ¹H NMR (500 MHz, (CD₃)₂CO, r.t.) δ: 3.17 (s, 2H_{CH3} DMSO, 6 H), 3.46 (s, 2H_{CH3} DMSO, 6 H), 3.52 (s, 2H_{CH3} DMSO, 6 H), 6.42 (t, *J* = 2.3 Hz, H⁴, 1 H), 7.68 (d, *J* = 2.3 Hz, H³, 1 H), 8.55 (d, *J* = 2.3 Hz, H⁵, 1 H), 13.98 (s, NH, 1 H). ¹³C{¹H} NMR (126 MHz, (CD₃)₂CO, r.t.) δ: 45.6 (2C, 2C_{CH3} DMSO), 46.8 (2C, 2C_{CH3} DMSO), 47.4 (2C, 2C_{CH3} DMSO), 107.0 (C⁴), 130.5 (C³), 141.9 (C⁵). IR (cm^{–1}): 615 w, 675 m, 708 w, 719 w, 771 m, 816 d, 880 vw, 935 m, 951 m, 968 m, 981 m, 1011 vs, 1036 m, 1060 m, 1091 s, 1171 vw, 1359 w, 1408 m, 2838 w, 2918 w, 3122 w. Calcd for C₉Cl₂H₂₂N₂O₃RuS₃: C, 22.84; H, 4.65; N, 5.92; S, 20.30. Found: C, 22.88; H, 4.62; N, 5.77; S, 20.54.

***cis*-fac-[RuCl₂(DMSO)₃(indzH)], 1b.** This complex was prepared by a modification of the method previously described in

the literature. Some spectroscopic data were also previously reported.¹⁶ A mixture of *cis*-[RuCl₂(DMSO)₄] (0.097 g, 0.2 mmol) and indzH (0.024 g, 0.2 mmol) in MeOH (10 mL) was stirred at r.t. for 1 h. The clear yellow solution was left to stand at –20 °C, giving a yellow microcrystalline solid, which was decanted, washed with Et₂O (2 × 5 mL approximately), and dried *in vacuo*, yielding 0.078 g, 74%. ¹H NMR (500 MHz, (CD₃)₂CO, r.t.) δ: 3.51 (s, 2H_{CH3} DMSO, 6 H), 3.50 (s, 2H_{CH3} DMSO, 6 H), 3.21 (s, 2H_{CH3} DMSO, 6 H), 7.22 (ddd, *J* = 8.2, 6.9, 0.9 Hz, H⁵, 1 H), 7.47 (ddd, *J* = 8.1, 6.9, 1.1 Hz, H⁶, 1 H), 7.73 (dd, *J* = 8.1, 0.9 Hz, H⁷, 1 H), 7.85 (dd, *J* = 8.2, 1.1 Hz, H⁴, 1 H), 9.16–9.10 (m, H³, 1 H), 14.12 (s, NH, 1 H). ¹³C{¹H} NMR (126 MHz, (CD₃)₂CO, r.t.) δ: 44.9 (2C, 2C_{CH3} DMSO), 46.1 (2C, 2C_{CH3} DMSO), 46.5 (2C, 2C_{CH3} DMSO), 110.4 (C⁷), 120.8 (C⁴), 121.5 (C⁵), 127.9 (C⁶), 128.2 (C^{7a}), 138.4 (C³), 140.4 (C^{3a}). IR (cm^{–1}): 3929 mw, 3238 w, 3133 w, 3028 w, 3013 w, 2922 w, 2324 w, 2286 w, 2164 w, 2149 w, 2113 w, 2051 w, 1981 w, 1933 w, 1797 w, 1700 w, 1626 m, 1585 w, 1509 m, 1483 w, 1446 m, 1412 m, 1380 m, 1359 m, 1307 m, 1287 m, 1270 m, 1253 w, 1218 w, 1193 m, 1148 w, 1084 vs, 1059 s, 1011 s, 973 m, 952 s, 920 m, 902 m, 863 w, 852 w, 781 w, 757 m, 749 m, 720 m, 678 m, 620 w. Calcd for C₁₃Cl₂H₂₄N₂O₃RuS₃: C, 29.83; H, 4.59; N, 5.35; S, 18.36. Found: C, 29.75; H, 4.61; N, 5.19; S, 18.43.

***fac*-[RuCl(DMSO)₃(NH=C(Me)pz-κ²N,N)](OTf), 2a.** AgOTf (0.051 g, 0.20 mmol) was added to a solution of **1a** (0.095 g, 0.20 mmol) in MeCN (10 mL). 100 μL of aqueous 0.02 M NaOH (0.002 mmol) was then added and the mixture was stirred at r.t. for 12 h in the absence of light. The reaction mixture was filtered to remove solid AgCl and dried *in vacuo*. The solid was crystallized in MeCN/Et₂O at –20 °C, giving a colorless microcrystalline solid, which was decanted, washed with Et₂O (2 × 5 mL approximately), and dried *in vacuo*, yielding 0.079 g, 63%. ¹H NMR (500 MHz, (CD₃)₂CO, r.t.) δ: 2.83 (s, H_{CH3} DMSO, 3 H), 3.02 (s, N=C(CH₃), 3 H), 3.19 (s, H_{CH3} DMSO, 3 H), 3.39 (s, H_{CH3} DMSO, 3 H), 3.55 (s, H_{CH3} DMSO, 3 H), 3.57 (s, H_{CH3} DMSO, 3 H), 3.61 (s, H_{CH3} DMSO, 3 H), 6.94 (t, *J* = 2.5 Hz, H⁴, 1 H), 8.78 (d, *J* = 2.5 Hz, H³, 1 H), 8.87 (d, *J* = 2.5 Hz, H⁵, 1 H), 11.00 (s, NH, 1 H). ¹³C{¹H} NMR (126 MHz, (CD₃)₂CO, r.t.) δ: 17.8 (N=C(CH₃)), 44.6 (C_{CH3} DMSO), 45.2 (C_{CH3} DMSO), 45.7 (C_{CH3} DMSO), 46.2 (C_{CH3} DMSO), 47.0 (C_{CH3} DMSO), 47.1 (C_{CH3} DMSO), 111.0 (C⁴), 134.9 (C³), 148.7 (C⁵), 164.3 (N=C(CH₃)). IR (cm^{–1}): 633 s, 685 m, 724 w, 759 w, 779 m, 920 w, 939 m, 964 m, 977 m, 1021 vs, 1052 w, 1086 vs, 1107 s, 1107 m, 1152 m, 1173 m, 1229 s, 1244 m, 1279 w, 1411 w, 1656 m, 2923 w, 3005 w, 3102 w, 3205 w. Calcd for C₁₂ClF₃H₂₅N₃O₆RuS₄: C, 22.95; H, 3.98; N, 6.69; S, 20.40. Found: C, 22.75; H, 3.91; N, 6.72; S, 20.55.

***fac*-[RuCl(DMSO)₃(NH=C(Me)indz-κ²N,N)](OTf), 2b.** AgOTf (0.049 g, 0.19 mmol) was added to a solution of **1b** (0.100 g, 0.19 mmol) in MeCN (10 mL) and the mixture was stirred at r.t. for 12 h in the absence of light. The reaction mixture was filtered to remove solid AgCl and dried *in vacuo*. The solid was crystallized in MeCN/Et₂O at –20 °C, giving a colorless microcrystalline solid, which was decanted, washed with Et₂O (2 × 5 mL approximately), and dried *in vacuo*, yielding 0.801 g, 62%. ¹H NMR (500 MHz, (CD₃)₂CO, r.t.) δ: 3.10 (s, H_{CH3} DMSO, 3 H),



3.24 (s, H_{CH_3} DMSO, 3 H), 3.33 (s, $N=C(CH_3)$, 3 H), 3.44 (s, H_{CH_3} DMSO, 3 H), 3.60 (s, 2 H_{CH_3} DMSO, 6 H), 3.65 (s, H_{CH_3} DMSO, 3 H), 7.61 (t, $J = 8.0$ Hz, H^6 , 1 H), 7.83 (t, $J = 8.0$ Hz, H^5 , 1 H), 8.17 (d, $J = 8.2$ Hz, H^4 , 1 H), 8.19 (d, $J = 8.2$ Hz, H^7 , 1 H), 9.69 (s, H^3 , 1 H), 10.51 (s, NH, 1 H). $^{13}C\{^1H\}$ NMR (126 MHz, $(CD_3)_2CO$, r.t.) δ : 20.5 ($N=C(CH_3)$), 45.0–50.0 (6 C, C_{CH_3} DMSO), 112.9 (C^4), 123.3 (C^7), 125.3 (C^6), 125.6 (C^{7a}), 131.3 (C^5), 140 (C^{3a}), 146.3 (C^3), 165.0 ($N=C(CH_3)$). IR (cm^{-1}): 3221 w, 3117 vw, 3022 w, 2971 vw, 1690 w, 1632 m, 1584 vw, 1511 w, 1484 m, 1468 w, 1428 m, 1409 w, 1356 w, 1316 w, 1276 s, 1249 vs, 1226 m, 1196 m, 1175 m, 1156 s, 1112 s, 1100 s, 1081 s, 1029 vs, 1019 vs, 977 m, 966 m, 943 w, 914 m, 864 w, 795 w, 756 s, 719 w, 683 m, 634 vs. Calcd for $C_{16}ClF_3H_{27}N_3O_6RuS_4$: C, 28.34; H, 3.99; N, 6.20; S, 18.89. Found: C, 28.45; H, 4.00; N, 6.39; S, 18.68.

fac-[RuCl(DMSO)₃(NH=C(Ph)pz- κ^2 N,N)](OTf), 3a. AgOTf (0.051 g, 0.2 mmol) was added to a solution of **1a** (0.095 g, 0.2 mmol) in $(CH_3)_2CO$ (10 mL). 100 μ L of PhCN and 100 μ L of aqueous 0.02 M NaOH (0.002 mmol) were then added and the mixture was stirred at r.t. for 24 h in the absence of light. The reaction mixture was filtered to remove solid AgCl and dried *in vacuo*. The solid was crystallized in $(CH_3)_2CO/Et_2O$ at -20 °C, giving a yellow microcrystalline solid, which was decanted, washed with Et_2O (2×5 mL approximately), and dried *in vacuo*, yielding 0.084 g, 61%. 1H NMR (500 MHz, $(CD_3)_2CO$, r.t.) δ : 3.31 (s, H_{CH_3} DMSO, 3 H), 3.34 (s, H_{CH_3} DMSO, 3 H), 3.47 (s, H_{CH_3} DMSO, 3 H), 3.63 (s, H_{CH_3} DMSO, 3 H), 3.64 (s, H_{CH_3} DMSO, 3 H), 3.69 (s, H_{CH_3} DMSO, 3 H), 6.97 (dd, $J = 3.2$, 2.1 Hz, H^4 pz, 1 H), 7.59–8.04 (m, C_6H_5 Ph, 5 H), 8.45 (dd, $J = 3.2$, 0.6 Hz, H^3 pz, 1 H), 9.03 (dd, $J = 2.1$, 0.6 Hz, H^5 pz, 1 H), 11.21 (s, NH, 1 H). $^{13}C\{^1H\}$ NMR (126 MHz, $(CD_3)_2CO$, r.t.) δ : 44.6 (C_{CH_3} DMSO), 45.2 (C_{CH_3} DMSO), 45.7 (C_{CH_3} DMSO), 46.2 (C_{CH_3} DMSO), 47.0 (C_{CH_3} DMSO), 47.1 (C_{CH_3} DMSO), 111.6 (C^4), 129.4–133.4 (5C, C Ph), 136.4 (C^3), 149.7 (C^5), 165.0 ($N=C$ (Ph)), 166.0 (C_{ipso} Ph). IR (cm^{-1}): 3508 w, 3220 w, 3138 w, 3025 w, 2925 w, 2323 w, 2287 w, 2231 w, 2188 w, 2164 w, 2140 w, 2113 w, 2083 w, 2050 w, 1982 w, 1917 w, 1626 m, 1577 w, 1527 w, 1497 w, 1456 w, 1442 m, 1415 m, 1321 w, 1274 s, 1246 vs, 1224 s, 1156 s, 1079 s, 1027 vs, 973 m, 922 m, 896 m, 760 m, 724 w, 704 m, 684 m, 636 s. Calcd for $C_{17}ClF_3H_{27}N_3O_6RuS_4$: C, 29.54; H, 3.94; N, 6.08; S, 18.56. Found: C, 29.62; H, 3.78; N, 6.01; S, 18.47.

fac-[RuCl(DMSO)₃(NH=C(Ph)indz- κ^2 N,N)](OTf), 3b. AgOTf (0.051 g, 0.2 mmol) was added to a solution of **1b** (0.105 g, 0.2 mmol) in $(CH_3)_2CO$ (10 mL). 100 μ L of PhCN was then added and the mixture was stirred at r.t. for 24 h in the absence of light. The reaction mixture was filtered to remove solid AgCl and dried *in vacuo*. The solid was crystallized in $(CH_3)_2CO/Et_2O$ at -20 °C, giving a yellow microcrystalline solid, which was decanted, washed with Et_2O (2×5 mL approximately), and dried *in vacuo*, yielding 0.093 g, 63%. 1H NMR (500 MHz, $(CD_3)_2CO$, r.t.) δ : 3.31 (s, H_{CH_3} DMSO, 3 H), 3.34 (s, H_{CH_3} DMSO, 3 H), 3.47 (s, H_{CH_3} DMSO, 3 H), 3.63 (s, H_{CH_3} DMSO, 3 H), 3.64 (s, H_{CH_3} DMSO, 3 H), 3.69 (s, H_{CH_3} DMSO, 3 H), 6.38 (dt, $J = 8.5$, 0.8 Hz, H^6 indz, 1 H), 7.46–7.94 (m, H^4 indz, H^5 indz, C_6H_5 Ph, 7 H), 8.17 (dd, $J = 8.5$, 0.8 Hz, H^7 indz, 1 H), 9.80 (d, $J = 0.8$ Hz, H^3 indz, 1 H), 10.85 (s, NH,

1 H). $^{13}C\{^1H\}$ NMR (126 MHz, $(CD_3)_2CO$, r.t.) δ : 44.5 (C_{CH_3} DMSO), 45.2 (C_{CH_3} DMSO), 45.8 (C_{CH_3} DMSO), 46.3 (C_{CH_3} DMSO), 47.0 (C_{CH_3} DMSO), 47.4 (C_{CH_3} DMSO), 112.2 (C^6 indz), 122.7 (C^{7a}), 123.2 (C^7 indz), 125.5–133.2 (7C, $C^{4,5}$ indz, C Ph), 140.7 (C^{3a}), 147.7 (C^3 indz), 165.0 ($N=C$ (Ph)), 166.0 (C_{ipso} Ph). IR (cm^{-1}): 3264 vw, 3080 vw, 3016 vw, 2964 vw, 1310 vw, 1288 vw, 1119 m, 1094 m, 1018 m, 974 m, 918 m, 713 vw, 677 w. Calcd for $C_{21}ClF_3H_{29}N_3O_6RuS_4$: C, 34.03; H, 3.94; N, 5.68; S, 17.31. Found: C, 33.92; H, 3.75; N, 5.44; S, 17.02.

Electrochemical experiments

Electrochemical experiments were performed by using a Model 6012D or 604E electrochemical analyzer from CH Instruments, Inc. Cyclic voltammetry experiments were performed under either $N_2(g)$ or $Ar(g)$ and $CO_2(g)$ in a one-compartment cell with a glassy carbon working electrode, a platinum wire counter electrode and an Ag/Ag^+ (10 mM $AgNO_3$ in acetonitrile, DMF or DMSO) reference electrode with ferrocene as an external reference. The glassy carbon working electrode was polished with 1.0 micron alumina powder, extensively rinsed with deionized water, and then polished for 60 seconds with 0.05 micron alumina powder (CH Instruments). The electrode was again rinsed with dry MeCN/DMF prior to all electrochemistry experiments. All experiments were performed by using 0.1 M TBAPF₆ as the supporting electrolyte unless otherwise specified, acetonitrile as the solvent, and with Ru complexes at a concentration of 1.0 mM unless indicated otherwise.

The solubility of saturated CO_2 in acetonitrile has been reported to be 0.28 M at 25 °C.²⁶ Changing the atmosphere from pure N_2 (or Ar) to pure CO_2 or *vice versa* required bubbling with the new gas for no less than five minutes. Lasting such time, the CVs obtained were the same as those obtained in the first scan under a specific atmosphere. Bubbling was maintained during the interim between scans. During the scan time, the PTFE was raised and kept above the surface of the solution to avoid agitation.

Controlled potential electrolysis

Controlled potential electrolysis (CPE) experiments were performed by using a Model 6012D electrochemical analyzer from CH Instruments, Inc. with 25 mL of an MeCN/TBAPF₆ solution in a cell with a carbon-rod working electrode (~ 1.9 cm² submersed area), a silver-wire pseudoreference, and a platinum-wire counter electrode separated from the solution using a medium-porosity frit. Gas chromatography analysis of the headspace was performed on an Agilent 7820 instrument equipped with an HP-Molesieve column and a gas sampling valve, as well as a thermal conductivity detector (TCD), stainless-steel column packed with molecular sieves (60/80 mesh), and UHP He as the carrier gas (flow rate = 35 mL min⁻¹). The operating temperatures of the injection port, the oven/column, and the detector were 100 °C, 80 °C, and 100 °C, respectively. Calibration curves for carbon monoxide and hydrogen were created by injecting known quantities of CO or H_2 into the electrochemical cell and then sampling the headspace.



Photocatalysis procedure

Photochemical reactions were performed in a 20 mL reaction vessel containing 10 mL of CO₂-saturated MeCN–TEOA solution (4 : 1 v/v); the Ru catalyst was present in a concentration of 0.1 mM, whereas the initial concentration of the photosensitizer, [Ru(bipy)₃]²⁺, was 1.6 mM. The solution was irradiated by >300 nm visible light. Analysis of the resulting compounds was performed as described in the CPE section.

Crystal structure determination for compounds 2a and 2b

Crystals were grown by slow diffusion of Et₂O into concentrated solutions of the complexes in MeCN (for **2a** and **2b**) at –20 °C. Relevant crystallographic details can be found in the CIF. A crystal was attached to a glass fiber and transferred to an Agilent SuperNova diffractometer fitted with an Atlas CCD detector. The crystals were kept at 293(2) K during data collection. Using Olex2,²⁷ the structures were solved with the ShelXT²⁸ structure solution program and then the structures were refined with the ShelXL²⁹ refinement package using least squares minimization. All non-hydrogen atoms were refined anisotropically. Hydrogen atoms were set in calculated positions and refined as riding atoms, with a common thermal parameter. All graphics were made with Olex2, and distances and angles of hydrogen bonds were calculated with PARST³⁰ (normalized values).³¹

Conflicts of interest

There are no conflicts of interest to declare.

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References

- (a) J. Qiao, Y. Liu, F. Hong and J. Zhang, *Chem. Soc. Rev.*, 2014, **43**, 631–675; (b) N. Elgrishi, M. B. Chambers, X. Wang and M. Fontecave, *Chem. Soc. Rev.*, 2017, **46**, 761–796; (c) R. Francke, B. Schille and M. Roemelt, *Chem. Rev.*, 2018, **118**, 4631–4701; (d) C. Jiang, A. W. Nichols and C. W. Machan, *Dalton Trans.*, 2019, **48**, 9454–9468; (e) S. Zhang, Q. Fan, R. Xia and T. J. Meyer, *Acc. Chem. Res.*, 2020, **53**, 255–264; (f) Y. J. Sa, C. W. Lee, S. Y. Lee, J. Na, U. Lee and Y. J. Hwang, *Chem. Soc. Rev.*, 2020, **49**, 6632–6665; (g) F. Franco, C. Rettenmaier, H. S. Jeon and B. R. Cuenya, *Chem. Soc. Rev.*, 2020, **49**, 6884–6946; (h) R.-Z. Zhang, B.-Y. Wu, Q. Li, L.-L. Lu, W. Shi and P. Cheng, *Coord. Chem. Rev.*, 2020, **422**, 213436; (i) N. W. Kinzel, C. Werlé and W. Leitner, *Angew. Chem., Int. Ed.*, 2021, **60**, 11628–11686; (j) P. Saha, S. Amanullah and A. Dey, *Acc. Chem. Res.*, 2022, **55**, 134–144; (k) E. Fujita, D. C. Grills, G. F. Manbeck and D. E. Polyansky, *Acc. Chem. Res.*, 2022, **55**, 616–628; (l) G. Marcandalli, M. C. O. Monteiro, A. Goyal and M. T. M. Koper, *Acc. Chem. Res.*, 2022, **55**, 1900–1911; (m) H.-Q. Liang, T. Beweries, R. Francke and M. Beller, *Angew. Chem., Int. Ed.*, 2022, **61**, e202200723; (n) X. She, Y. Wang, H. Xu, S. C. E. Tsang and S. P. Lau, *Angew. Chem., Int. Ed.*, 2022, **61**, 202211396; (o) K. Lei and B. Y. Xia, *Chem. – Eur. J.*, 2022, **28**, e202200141; (p) W. Nie and C. C. L. McCrory, *Dalton Trans.*, 2022, **51**, 6993–7010; (q) R. Ayyappan, I. Abdalghani, R. C. Da Costa and G. R. Owen, *Dalton Trans.*, 2022, **51**, 11582–11611; (r) Y. Yamazaki, M. Miyaji and O. Ishitani, *J. Am. Chem. Soc.*, 2022, **144**, 6640–6660.
- (a) H. Ishida, H. Tanaka, K. Tanaka and T. Tanaka, *J. Chem. Soc., Chem. Commun.*, 1987, 131–132; (b) H. Ishida, K. Tanaka and T. Tanaka, *Organometallics*, 1987, **6**, 181–186.
- (a) H. Ishida, K. Fujiki, T. Ohba, K. Ohkubo, K. Tanaka, T. Terada and T. Tanaka, *J. Chem. Soc., Dalton Trans.*, 1990, **2**, 2155–2160; (b) J. R. Pugh, M. R. M. Bruce, B. P. Sullivan and T. J. Meyer, *Inorg. Chem.*, 1991, **30**, 86–91; (c) H. Nagao, T. Mizukawa and K. Tanaka, *Inorg. Chem.*, 1994, **33**, 3415–3420; (d) K. Toyohara, H. Nagao, T. Mizukawa and K. Tanaka, *Inorg. Chem.*, 1995, **34**, 5399–5400; (e) M. M. Ali, H. Sato, T. Mizukawa, K. Tsuge, M. Hagab and K. Tanaka, *Chem. Commun.*, 1998, 249–250; (f) K. Tanaka and D. Ooyama, *Coord. Chem. Rev.*, 2002, **226**, 211–218; (g) F. H. Haghighi, H. Hadadzadeh, H. Farrokhpour, N. Serri, K. Abdia and H. A. Rudbarib, *Dalton Trans.*, 2014, **43**, 11317–11332; (h) D. J. Boston, Y. M. F. Pachón, R. O. Lezna, N. R. de Tacconi and F. M. MacDonnell, *Inorg. Chem.*, 2014, **53**, 6544–6553; (i) B. A. Johnson, S. Maji, H. Agarwala, T. A. White, E. Mijangos and S. Ott, *Angew. Chem., Int. Ed.*, 2016, **55**, 1825–1829; (j) B. A. Johnson, H. Agarwala, T. A. White, E. Mijangos, S. Maji and S. Ott, *Chem. – Eur. J.*, 2016, **22**, 14870–14880.
- (a) M.-N. Collomb-Dunand-Sauthier, A. Deronzier and R. Ziessel, *J. Chem. Soc., Chem. Commun.*, 1994, 189–191; (b) M.-N. Collomb-Dunand-Sauthier, A. Deronzier and R. Ziessel, *Inorg. Chem.*, 1994, **33**, 2961–2967; (c) S. Chardon-Noblat, M.-N. Collomb-Dunand-Sauthier, A. Deronzier, R. Ziessel and D. Zsoldos, *Inorg. Chem.*, 1994, **33**, 4410–4412; (d) S. Chardon-Noblat, A. Deronzier, R. Ziessel and D. Zsoldos, *Inorg. Chem.*, 1997, **36**, 5384–5389; (e) S. Chardon-Noblat, A. Deronzier, R. Ziessel and D. Zsoldos, *J. Electroanal. Chem.*, 1998, **444**, 253–260;



- (f) C. W. Machan, M. D. Sampson and C. P. Kubiak, *J. Am. Chem. Soc.*, 2015, **137**, 8564–8571.
- 5 (a) A. J. Morris, G. J. Meyer and E. Fujita, *Acc. Chem. Res.*, 2009, **42**, 1983–1994; (b) H. Takeda and O. Ishitani, *Coord. Chem. Rev.*, 2010, **254**, 346–354; (c) Y. Yamazaki, H. Takeda and O. Ishitani, *J. Photochem. Photobiol., C*, 2015, **25**, 106–137; (d) Y. Kuramochi, O. Ishitani and H. Ishida, *Coord. Chem. Rev.*, 2018, **373**, 333–356; (e) Y.-H. Luo, L.-Z. Dong, J. Liu, S.-L. Li and Y.-Q. Lan, *Coord. Chem. Rev.*, 2019, **390**, 86–126; (f) H. Kumagai, Y. Tamaki and O. Ishitani, *Acc. Chem. Res.*, 2022, **55**, 978–990; (g) N. Nandal and S. L. Jain, *Coord. Chem. Rev.*, 2022, **451**, 214271.
- 6 (a) H. Ishida, T. Terada, K. Tanaka and T. Tanaka, *Inorg. Chem.*, 1990, **29**, 905–911; (b) J.-M. Lehn and R. Ziessel, *J. Organomet. Chem.*, 1990, **382**, 157–173; (c) K. Kobayashi, T. Kikuchi, S. Kitagawa and K. Tanaka, *Angew. Chem., Int. Ed.*, 2014, **53**, 11813–11817; (d) Y. Kuramochi, M. Kamiya and H. Ishida, *Inorg. Chem.*, 2014, **53**, 3326–3332; (e) R. R. Rodrigues, C. M. Boudreaux, E. T. Papish and J. H. Delcamp, *ACS Appl. Energy Mater.*, 2019, **2**, 37–46; (f) R. N. Sampaio, D. C. Grills, D. E. Polyansky, D. J. Szalda and E. Fujita, *J. Am. Chem. Soc.*, 2020, **142**, 2413–2428.
- 7 (a) R. Kuriki, K. Sekizawa, O. Ishitani and K. Maeda, *Angew. Chem., Int. Ed.*, 2015, **54**, 2406–2409; (b) Y. Kuramochi, K. Fukaya, M. Yoshida and H. Ishida, *Chem. – Eur. J.*, 2015, **21**, 10049–10060; (c) Y. Kuramochi, J. Itabashi, K. Fukaya, A. Enomoto, M. Yoshida and H. Ishida, *Chem. Sci.*, 2015, **6**, 3063–3074; (d) Y. Kuramochi, J. Itabashi, M. Toyama and H. Ishida, *ChemPhotoChem*, 2018, **2**, 314–322.
- 8 (a) P. L. Cheung, S. C. Kapper, T. Zeng, M. E. Thompson and C. P. Kubiak, *J. Am. Chem. Soc.*, 2019, **141**, 14961–14965; (b) M. R. Madsen, J. B. Jakobsen, M. H. Rønne, H. Liang, H. C. D. Hammershøj, P. Nørby, S. U. Pedersen, T. Skrydstrup and K. Daasbjerg, *Organometallics*, 2020, **39**, 1480–1490; (c) C. Back, Y. Seo, S. Choi, M. S. Choe, D. Lee, J.-O. Baeg, H.-J. Son and S. O. Kang, *Inorg. Chem.*, 2021, **60**, 14151–14164; (d) Z. S. Dubrawski, C. Y. Chang, C. R. Carr, B. S. Gelfand and W. E. Piers, *Dalton Trans.*, 2022, **51**, 17381–17390.
- 9 F. Villafañe, *Coord. Chem. Rev.*, 2017, **339**, 128–137.
- 10 (a) M. Arroyo, Á. López-Sanvicente, D. Miguel and F. Villafañe, *Eur. J. Inorg. Chem.*, 2005, 4430–4437; (b) N. Antón, M. Arroyo, P. Gómez-Iglesias, D. Miguel and F. Villafañe, *J. Organomet. Chem.*, 2008, **693**, 3074–3080; (c) P. Gómez-Iglesias, M. Arroyo, S. Bajo, C. Strohmman, D. Miguel and F. Villafañe, *Inorg. Chem.*, 2014, **53**, 12437–12448.
- 11 (a) P. Gómez-Iglesias, F. Guyon, A. Khatyr, G. Ulrich, M. Knorr, J. M. Martín-Alvarez, D. Miguel and F. Villafañe, *Dalton Trans.*, 2015, **44**, 17516–17528; (b) E. Cuéllar, A. Diez-Varga, T. Torroba, P. Domingo-Legarda, J. Alemán, S. Cabrera, J. M. Martín-Alvarez, D. Miguel and F. Villafañe, *Inorg. Chem.*, 2021, **60**, 7008–7022.
- 12 P. Gómez-Iglesias, J. M. Martín-Alvarez, D. Miguel and F. Villafañe, *Dalton Trans.*, 2015, **44**, 17478–17481.
- 13 M. Arroyo, D. Miguel, F. Villafañe, S. Nieto, J. Pérez and L. Riera, *Inorg. Chem.*, 2006, **45**, 7018–7026.
- 14 M. Arroyo, P. Gómez-Iglesias, J. M. Martín-Alvarez, C. M. Alvarez, D. Miguel and F. Villafañe, *Inorg. Chem.*, 2012, **51**, 6070–6080.
- 15 Í. Ferrer, J. Rich, X. Fontrodona, M. Rodríguez and I. Romero, *Dalton Trans.*, 2013, **42**, 13461–13469.
- 16 E. Reisner, V. B. Arion, A. Ruffin, I. Chiorescu, W. F. Schmid and B. K. Keppler, *Dalton Trans.*, 2005, **2**, 2355–2364.
- 17 (a) C. J. Jones, J. A. McCleverty and A. S. Rothin, *Dalton Trans.*, 1986, 109–111; (b) A. Romero, A. Vegas and A. Santos, *J. Organomet. Chem.*, 1986, **310**, C8–C10; (c) J. López, A. Santos, A. Romero and A. M. Echavarren, *J. Organomet. Chem.*, 1993, **443**, 221–228; (d) M. R. Kollipara, P. Sarkhel, S. Chakraborty and R. Lalrempuia, *J. Coord. Chem.*, 2003, **56**, 1085–1091; (e) P. Govindaswamy, Y. A. Mozharivskyj and M. R. Kollipara, *J. Organomet. Chem.*, 2004, **689**, 3265–3274.
- 18 (a) A. G. Jeffrey, in *An Introduction to Hydrogen Bonding*, Oxford University Press, New York, 1997; (b) T. Steiner, *Angew. Chem., Int. Ed.*, 2002, **41**, 48–76.
- 19 G. Gritzner and J. Kuta, *Pure Appl. Chem.*, 1984, **56**, 461–466.
- 20 E. Cuéllar, L. Pastor, G. García-Herbosa, J. Nganga, A. M. Angeles-Boza, A. Diez-Varga, T. Torroba, J. M. Martín-Alvarez, D. Miguel and F. Villafañe, *Inorg. Chem.*, 2021, **60**, 692–704.
- 21 B. Merillas, E. Cuéllar, A. Diez-Varga, T. Torroba, G. García-Herbosa, S. Fernández, J. Lloret-Fillol, J. M. Martín-Alvarez, D. Miguel and F. Villafañe, *Inorg. Chem.*, 2020, **59**, 11152–11165.
- 22 (a) P. S. Campbell, M. H. G. Precht, C. C. Santini and P.-H. Haumesser, *Curr. Org. Chem.*, 2013, **17**, 414–429; (b) Y. Na, S. Park, S. B. Han, H. Han, S. Ko and S. Chang, *J. Am. Chem. Soc.*, 2004, **126**, 250–258.
- 23 (a) J. M. Kelly, C. M. O'Connell and J. G. Vos, *J. Chem. Soc., Dalton Trans.*, 1986, 253–258; (b) M.-N. Collomb-Dunand-Sauthier, A. Deronzier and R. Ziessel, *J. Organomet. Chem.*, 1993, **444**, 191–198; (c) E. Eskelinen, M. Haukka, T. Venäläinen, T. A. Pakkanen, M. Wasberg, S. Chardon-Noblat and A. Deronzier, *Organometallics*, 2000, **19**, 163–169; A. Gabrielsson, S. Zális, P. Matousek, M. Towrie and A. Vlček Jr., *Inorg. Chem.*, 2004, **43**, 7380–7388.
- 24 P. L. Cheung, C. W. Machan, A. Y. S. Malkhasian, J. Agarwal and C. P. Kubiak, *Inorg. Chem.*, 2016, **55**, 3192–3198.
- 25 C. E. McCusker and J. K. McCusker, *Inorg. Chem.*, 2011, **50**, 1656–1669.
- 26 (a) A. Gennaro, A. A. Isse and E. Vianello, *J. Electroanal. Chem. Interfacial Electrochem.*, 1990, **289**, 203–215; (b) E. Fujita, C. Creutz, N. Sutin and D. J. Szalda, *J. Am. Chem. Soc.*, 1991, **113**, 343–353.
- 27 O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- 28 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 3–8.



- 29 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 30 (a) M. Nardelli, *Comput. Chem.*, 1983, **7**, 95–98; (b) M. Nardelli, *J. Appl. Crystallogr.*, 1995, **28**, 659–659.
- 31 (a) G. A. Jeffrey and L. Lewis, *Carbohydr. Res.*, 1978, **60**, 179–182; (b) R. Taylor and O. Kennard, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1983, **39**, 133–138.

