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# Effect of sulfonamidoethylenediamine substituents in Ru<sup>II</sup> arene anticancer catalysts on transfer hydrogenation of coenzyme NAD<sup>+</sup> by formate<sup>†</sup>

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A series of neutral *pseudo*-octahedral Ru<sup>II</sup> sulfonamidoethylenediamine complexes [( $\eta^6$ -*p*-cym)Ru(*N,N'*)Cl] where *N,N'* is *N*-(2-(R<sup>1</sup>,R<sup>2</sup>-amino)ethyl)-4-toluenesulfonamide (TsEn(R<sup>1</sup>,R<sup>2</sup>)) R<sup>1</sup>,R<sup>2</sup> = Me,H (**1**); Me,Me (**2**); Et,H (**3**); benzyl,H (Bz, **4**); 4-fluorobenzyl,H (4-F-Bz, **5**) or naphthalen-2-ylmethyl,H (Naph, **6**), were synthesised and characterised including the X-ray crystal structure of **3**. These complexes catalyse the reduction of NAD<sup>+</sup> regioselectively to 1,4-NADH by using formate as the hydride source. The catalytic efficiency depends markedly on the steric and electronic effects of the *N*-substituent, with turnover frequencies (TOFs) increasing in the order: **1** < **2** < **3**, **6** < **4**, **5**, achieving a TOF of 7.7 h<sup>-1</sup> for **4** with a 95% yield of 1,4-NADH. The reduction rate was highest between pH\* (deuterated solvent) 6 and 7.5 and improved with an increase in formate concentration (TOF of 18.8 h<sup>-1</sup>, 140 mM formate). The calculations suggested initial substitution of an aqua ligand by formate, followed by hydride transfer to Ru<sup>II</sup> and then to NAD<sup>+</sup>, and indicated specific interactions between the aqua complex and both NAD<sup>+</sup> and NADH, the former allowing a preorganisation involving interaction between the aqua ligand, formate anion and the pyridine ring of NAD<sup>+</sup>. The complexes exhibited antiproliferative activity towards A2780 human ovarian cancer cells with IC<sub>50</sub> values ranging from 1 to 31  $\mu$ M, the most potent complex, [( $\eta^6$ -*p*-cym)Ru(TsEn(Bz, H))Cl] (**4**, IC<sub>50</sub> = 1.0  $\pm$  0.1  $\mu$ M), having a potency similar to the anticancer drug cisplatin. Co-administration with sodium formate (2 mM), increased the potency of all complexes towards A2780 cells by 20–36%, with the greatest effect seen for complex **6**.

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## 1. Introduction

Nicotinamide adenine dinucleotide (NAD<sup>+</sup>) and its reduced form (NADH), as well as their phosphorylated derivatives, NADP<sup>+</sup> and NADPH, play a vital role in biological systems as redox coenzymes.<sup>1</sup> More than 400 enzymatic redox reactions rely on the action of nicotinamide enzymes, in which the transformation of NAD(P)<sup>+</sup> to NAD(P)H is involved.<sup>2–4</sup> The reduction of pyridinium salts (*e.g.* NAD<sup>+</sup>) to dihydropyridine com-

pounds (*e.g.* NADH) is of critical importance for energy storage and release in cell metabolism.<sup>3,4</sup> Transition metal-mediated catalytic reduction of NAD<sup>+</sup> to NADH, using hydrogen,<sup>5</sup> 2-propanol,<sup>6,7</sup> glycerol<sup>8</sup> and sodium formate as hydride donors, has been intensively studied for the last three decades.<sup>9–13</sup> Compared to reduction with H<sub>2</sub> (hydrogenation), transfer hydrogenation (TH) reactions have the advantage of being simpler, without the need for any high external pressure and use readily available, safer-to-handle, hydride sources.<sup>14</sup> Also, TH reduction of NAD(P)<sup>+</sup> artificially has attracted wide interest as an *in vitro* mimic for enzymatic reactions performed under biologically relevant conditions.<sup>15</sup>

The pathways of hydride transfer between pyridinium salts and dihydropyridine compounds are also of interest. The first mechanistic study of the TH reduction of BNA<sup>+</sup> (1-benzylnicotinamide), as a model for NAD<sup>+</sup> was reported by Steckhan and Fish *et al.* using [( $\eta^5$ -Cp\*)Rh(bipy)Cl] as the catalyst and sodium formate as a hydride source in aqueous media in the 1990s.<sup>10–12,16,17</sup> They proposed a catalytic cycle involving a

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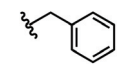
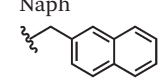


ring-slippage  $\eta^4$ -Cp\* intermediate with Rh coordinated to the amide of the pyridine ring.<sup>18</sup> Knör *et al.* reported a Rh-coordinated poly(arylene-ethynylene)-*alt*-poly(arylene-vinylene) polymer as photocatalyst for the reduction of NAD<sup>+</sup>; involving a possible photoexcited polymer chain being quenched and transferring an electron to the Rh<sup>III</sup> active centre.<sup>19</sup> More recently, Yoon *et al.* described a mechanism involving hydride transfer to Cp\* and formation of the Rh<sup>I</sup> intermediate  $[(\eta^4\text{-Cp}^*\text{-H})\text{Rh}((\text{CH}_2\text{OH})_2\text{-bipy})]^+$  followed by hydride transfer from the *endo* orientation of the C–H bond to maintain the 1,4-regioselectivity of NADH.<sup>20</sup>

The half-sandwich ruthenium complex  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsDPEN})\text{Cl}]$  (TsDPEN: *N*-((1*S*,2*S*)-2-amino-1,2-diphenylethyl)-4-methylbenzenesulfonamide) was first reported by Noyori and coworkers in 1995.<sup>21,22</sup> Potent catalytic activity has been shown in asymmetric TH reduction of aromatic ketones. Most recently, the 16-electron Os analogues  $[(\eta^6\text{-arene})\text{Os}(\text{TsDPEN})]$  of Noyori type complexes were reported to reduce pyruvate enantioselectively to (*D*- or *L*-) lactate *via* asymmetric transfer hydrogenation in human cancer cells.<sup>23</sup> Nonetheless, the hydrophobic nature of the two phenyl groups on the ethylene backbone limits its application as a possible catalyst for TH reduction of NAD<sup>+</sup> under biologically relevant conditions. Complexes with chelating diamine ligands such as complex 7 in Fig. 1, display good aqueous solubility but poor catalytic activity in TH reduction of NAD<sup>+</sup>.<sup>24</sup> However, *p*-cymene (*p*-cym) complexes with functional sulfonyl substituents such as  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$  (*e.g.* complex 8 in Fig. 1),<sup>25</sup> exhibit good solubility in water and improved catalysis for NAD<sup>+</sup> reduction to NADH in aqueous media. Moreover, co-administration of  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$  with low non-cytotoxic doses of sodium formate led to an enhancement of the antiproliferative activity against A2780 human ovarian cancer cells by up to 50%.<sup>15,26</sup>

Here we investigate the effect on catalytic reduction of NAD<sup>+</sup> using formate as a hydride source upon variation of substituents on the amino group of the *N,N*-chelating TsEn ligand in Ru<sup>II</sup> complexes  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{N,N}')\text{Cl}]$  where *N,N'* is *N*-(2-(methylamino)ethyl)-4-toluenesulfonamide (TsEn(Me,H), 1), *N*-(2-(dimethylamino)ethyl)-4-toluenesulfonamide (TsEn(Me,Me), 2), *N*-(2-(ethylamino)ethyl)-4-toluene sulfonamide (TsEn(Et,H), 3), *N*-(2-(benzylamino)ethyl)-4-toluenesulfonamide (TsEn(Bz,H), 4), *N*-(2-((4-fluorobenzyl)amino)ethyl)-4-toluenesulfonamide (TsEn(4-F-Bz,H), 5) and *N*-(2-(naphthalen-2-ylmethyl)amino)ethyl)-4-toluenesulfonamide (TsEn(Naph,H), 6) (Table 1). In addition, the catalytic mechanism was investigated both experimentally and by density functional theory

Table 1 Ru<sup>II</sup> complexes studied in this work

Complex	R <sup>1</sup>	R <sup>2</sup>
1	Me	H
2	Me	Me
3	Et	H
4	Bz	H
5		H
6		H

(DFT) calculations. We also explored the effect of co-administration of formate on the antiproliferative activity of these complexes against A2780 human ovarian cancer cells.

## 2. Experimental

### 2.1 Materials

Ruthenium(III) trichloride hydrate was purchased from Precious Metals Online (PMO Pty Ltd) and used as received. Toluenesulfonyl chloride, sodium formate and  $\beta$ -nicotinamide adenine dinucleotide hydrate (NAD<sup>+</sup>) were obtained from Sigma-Aldrich. Magnesium sulfate, potassium hydroxide, sodium chloride, and hydrochloric acid were obtained from Fisher Scientific.  $\alpha$ -Phellandrene was purchased from SAFC. The Ru<sup>II</sup> precursor dimer  $[(\eta^6\text{-}p\text{-cym})\text{RuCl}_2]_2$  was prepared following literature methods,<sup>27</sup> as were the ligands 4-methyl-*N*-(2-(methylamino)ethyl)benzene sulphonamide (TsEn(Me,H))<sup>28</sup> and *N*-(2-(dimethylamino)ethyl)-4-methylbenzenesulfonamide (TsEn(Me,Me)).<sup>29</sup> The solvents used for NMR spectroscopy were purchased from Sigma-Aldrich and Cambridge Isotope Laboratories Inc. Non-dried solvents used in synthesis were obtained from Fisher Scientific. Solvents were used as received, except in the case of ethanol, 2-propanol, and methanol, which were degassed prior to use by bubbling with nitrogen.

A2780 human ovarian carcinoma cells were obtained from the European Collection of Cell Cultures. The cell line was grown in Roswell Park Memorial Institute medium (RPMI-1640) supplemented with 10% of foetal calf serum, 1% v/v of 2 mM glutamine and 1% v/v penicillin/streptomycin (10 000 units). All cells were grown as adherent monolayers at 310 K in a 5% CO<sub>2</sub>-humidified atmosphere and passaged at *ca.* 70–80% confluency.

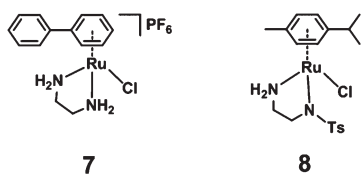


Fig. 1 Organometallic Ru<sup>II</sup> complexes  $[(\eta^6\text{-biph})\text{Ru}(\text{en})\text{Cl}]\text{PF}_6$  (7) and  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$  (8).



## 2.2 Instruments

NMR spectra were acquired on Bruker HD-300, HD-400, HD-500, and AV III 600 spectrometers.  $^1\text{H}$  NMR chemical shifts were internally referenced to TMS *via* 1,4-dioxane in  $\text{D}_2\text{O}$  ( $\delta = 3.75$  ppm) or residual protiated  $\text{d}_4\text{-MeOD}$  ( $\delta = 3.31$  ppm), or  $\text{CDCl}_3$  ( $\delta = 7.26$  ppm). 1D spectra were recorded using standard pulse sequences. Typically, data were acquired with 16 transients into 32k data points over a spectral width of 14 ppm and, for the kinetic experiment, 32 transients into 32k data points over a spectral width of 30 ppm using a relaxation delay of 2 s.

Elemental analysis were performed by Warwick Analytical using an Exeter Analytical elemental analyzer (CE440).

Positive ion electrospray mass spectra were obtained on an Agilent 6130B ion mass spectrometer. High resolution mass spectrometry data were obtained on a Bruker Maxis Plus Q-TOF instrument.

X-ray crystallographic diffraction data were collected on an Oxford Diffraction Gemini four-circle system with a Ruby CCD area detector. The structure was refined by full-matrix least-squares against  $F^2$  using SHELXL 9735 and solved by direct methods using SHELXS36 (TREF) with additional light atoms found by Fourier methods. The atoms from the sulfonamide nitrogen to the end of the chain (C10 C11 N12 C13) were modelled as disordered over two positions related by a small ruffle in the chain. The occupancy of the two positions was linked to a free variable which refined to 86 : 14. The minor component was refined isotropically. X-ray crystallographic data for complex **3** has been deposited in the Cambridge Crystallographic Data Center (CCDC) under the accession number CCDC 1571331.†

ICP-OES analysis were carried out on a PerkinElmer Optima 5300 DV series Optical Emission Spectrophotometer. The water used for ICP-OES analysis was doubly deionized (DDW) using a Millipore Milli-Q water purification system and a USF Elga UHQ water deionizer. The ruthenium Specpure plasma standard (ruthenium chloride,  $1004 \pm 5 \mu\text{g mL}^{-1}$  in 10% v/v hydrochloric acid) was diluted with 3.6% v/v  $\text{HNO}_3$  to freshly prepare calibrants at concentrations of 50–700 ppb. Calibration standards were adjusted to match the sample matrix by standard addition of sodium chloride (TraceSELECT®). Total dissolved solids did not exceed 0.2% w/v. Data were acquired and processed using WinLab32 V3.4.1 for Windows.

ICP-MS analysis were carried out on an Agilent Technologies 7500 series ICP-MS instrument. The water used for ICP-MS analysis was double-deionized (DDW) using a Millipore Milli-Q water purification system and a USF Elga UHQ water deionizer. The Ruthenium Specpure plasma standard (ruthenium chloride,  $1004 \pm 5 \mu\text{g mL}^{-1}$  in 10% v/v hydrochloric acid) was diluted with 3.6% v/v  $\text{HNO}_3$  to prepare calibrants freshly at concentrations of 0.1–1000 ppb. The ICP-MS instrument was set to detect  $^{101}\text{Ru}$  in no gas mode. Total dissolved solids did not exceed 0.1% w/v. An internal standard of  $^{166}\text{Er}$  (50 ppb) was used. Data were acquired using ICP-MS-TOP and proceeded using Offline Data Analysis (ChemStation version B.03.05, Agilent Technologies, Inc.).

pH values were measured using a Minilab IQ125 pH meter equipped with a ISFET silicon chip pH sensor and referenced in KCl gel.  $\text{pH}^*$  values (pH meter reading without correction for the effect of deuterium on the sensor) of NMR samples in  $\text{D}_2\text{O}$  were measured at 310 K. Relative hydrophobicity measurements were performed utilising the Agilent 1200 HPLC system with a VWD and 50  $\mu\text{L}$  loop. The column was an Agilent Zorbax 300SB C18,  $150 \times 4.6$  mm with a 5  $\mu\text{m}$  pore size. The mobile phase was  $\text{H}_2\text{O}$  (50 mM NaCl)/ $\text{H}_2\text{O}/\text{CH}_3\text{CN}$  1 : 1 (50 mM NaCl), with a flow of  $1 \text{ mL min}^{-1}$ . The detection wavelength was set at 254 nm with the reference wavelength at 360 nm.

## 2.3 Turnover frequency determination

**UV-vis spectroscopy.** In a typical experiment, 330  $\mu\text{L}$  of each solution (84  $\mu\text{M}$  complex in  $\text{MeOH}/\text{H}_2\text{O}$  1 : 9 v/v, 102 mM sodium formate and 510  $\mu\text{M}$   $\text{NAD}^+$  in  $\text{H}_2\text{O}$ ) was added to a 1 mL cuvette, and the pH adjusted to 7.2, bringing the total volume to 1 mL (final concentrations: Ru complex 28  $\mu\text{M}$ ;  $\text{NAD}^+$  170  $\mu\text{M}$ ;  $\text{NaHCO}_2$  34 mM; molar ratio 1 : 6 : 1200). UV spectra were recorded and the absorbance at 340 nm was monitored every 5 min until completion of the reaction.

**NMR spectroscopy.** Complexes were dissolved in  $\text{d}_4\text{-MeOD}/\text{D}_2\text{O}$  (1 : 4, v/v) (1.4 mM) in a glass vial. Solutions of sodium formate (35 mM) and  $\text{NAD}^+$  (5.6 mM) in  $\text{D}_2\text{O}$  were also prepared and then incubated at 310 K,  $\text{pH}^* 7.2 \pm 0.1$ . An aliquot of 200  $\mu\text{L}$  from each solution was added to a 5 mm NMR tube, giving a final volume of 0.64 mL (Ru complex 0.44 mM;  $\text{NAD}^+$  1.75 mM;  $\text{NaHCO}_2$  10.94 mM; molar ratio 1 : 4 : 25). A  $^1\text{H}$  NMR spectrum was recorded at 310 K every 162 s until the completion of the reaction. Further experiments under similar conditions using different concentrations of sodium formate (complex **4**,  $\text{NAD}^+$  and sodium formate in ratio of 1 : 4 :  $X$ , where  $X = 10, 25, 50$ , and 100 mol equiv.) and different concentrations of  $\text{NAD}^+$  (complex **4**,  $\text{NAD}^+$  and sodium formate in ratio of 1 :  $Y$  : 25, where  $Y = 2, 4, 6$  and 10) were also studied. Another series of experiments using different  $\text{pH}^*$  values of the reaction solutions (5, 6, 7, 8 and 9) were also performed. Molar ratios of  $\text{NAD}^+$  and  $\text{NADH}$  were determined by integrating  $^1\text{H}$  NMR peaks corresponding to  $\text{NAD}^+$  (9.33 ppm) and 1,4- $\text{NADH}$  (6.96 ppm). The turnover number (TON) for the reaction was calculated as follows:

$$\text{TON} = \frac{I_{6.96}}{I_{6.96} + I_{9.33}} \frac{[\text{NAD}^+]}{[\text{Catalyst}]}$$

where  $I_n$  is the integral of the signal at  $n$  ppm and  $[\text{NAD}^+]$  is the concentration of  $\text{NAD}^+$  at the start of the reaction.

## 2.4 Cell growth inhibition assays

The antiproliferative activity of complexes **1–6** was determined in A2780 ovarian cancer cells. Briefly, 96-well plates were used to seed 5000 cells per well. Cells were incubated in drug-free medium at 310 K for 48 h before addition of tested compounds (prepared by serial dilution in culture medium containing 5% DMSO, typically 6 concentrations in the range: 0.01–100  $\mu\text{M}$ ). Exact Ru concentrations were determined by



ICP-OES and the maximum concentration of DMSO to which cells were exposed never exceeded 0.5% v/v. A drug exposure period of 24 h was allowed. After this, supernatants were removed by suction and each well was washed with PBS. A further 72 h were allowed for the cells to recover in drug-free medium at 310 K. The sulforhodamine B (SRB) assay was used to determine cell viability.<sup>30</sup> IC<sub>50</sub> values, as the concentration that causes 50% cell death, were determined as duplicates of triplicates in two independent sets of experiments and their standard deviation were calculated. Data were processed using Microsoft Excel and sigmoidal curves fitted using Origin 9.1.

## 2.5 Co-administration of Ru complexes with formate

Cell viability assays were carried out with complexes 1–6 with co-administration of sodium formate in A2780 ovarian cancer cells. These experiments were carried out as above (*in vitro* growth inhibition assay) with the following modifications: a fixed equipotent concentration of each Ru complex equal to  $1/3 \times \text{IC}_{50}$  in that cell line was used in coadministration with three different concentrations of sodium formate (0.5, 1.0 and 2.0 mM). Drug stock solutions (*ca.* 100  $\mu\text{M}$ ) were prepared and they were further diluted using media until working concentrations were achieved. Separately, a stock solution of sodium formate was prepared in saline. The complex and formate solutions were added to each well independently, but within 5 minutes of each other. All other experiment conditions were kept unchanged (drug exposure and cell recovery time, as well as, end point assay used).

## 2.6 Cellular accumulation

The accumulation studies for Ru complexes 1–6 were performed on A2780 ovarian cancer cells.  $1.5 \times 10^6$  cells were seeded on a six-well plate using 2 mL of cell culture medium. After 24 h of pre-incubation in drug-free medium at 310 K, cells were exposed to complexes at equipotent IC<sub>50</sub> concentrations for 24 h (prepared by serial dilution of a *ca.* 100  $\mu\text{M}$  stock solution, prepared using culture medium containing 5% DMSO. This solution was analysed by ICP-OES to determine Ru concentration before treatment of cells with Ru complex). After this time, drug solutions were removed by suction, cells were washed with PBS and then treated with trypsin–EDTA. A suspension of single cells was counted, and cell pellets were collected. Each pellet was digested overnight in freshly-distilled concentrated nitric acid (200  $\mu\text{L}$ , 72% v/v) at 353 K; the resulting solutions were diluted with double-distilled water to a final concentration of 3.6% v/v HNO<sub>3</sub>, and the amount of Ru in A2780 ovarian cells was determined by ICP-MS. These experiments did not include any cell recovery time in drug-free media; they were carried out in triplicate, and the standard deviations were calculated. Data were processed using Microsoft Excel and reported as ng Ru  $\times 10^6$  cells.

## 2.7 ROS determination

Flow cytometry analysis of ROS/superoxide induction in A2780 cells caused by exposure to complexes 1 and 4 was carried out using the Total ROS/Superoxide detection kit (Enzo-Life

Sciences) according to the manufacturer's instructions. Briefly,  $1.0 \times 10^6$  A2780 cells per well were seeded in a six-well plate. Cells were preincubated in drug-free media at 310 K for 24 h in a 5% CO<sub>2</sub> humidified atmosphere, and then drugs were added to triplicates wells at IC<sub>50</sub> concentration. After 24 h of drug exposure, supernatants were removed by suction and cells were washed with PBS and harvested. Staining was achieved by re-suspending the cell pellets in buffer containing the orange/green fluorescent reagents. Cells were analysed in a Becton Dickinson FACScan flow cytometer using FL1 channel Ex/Em: 490/525 nm for the oxidative stress and FL2 channel Ex/Em: 550/620 nm for superoxide detection. Data were gated using positive-stained (pyocyanin positive control), untreated-stained and untreated-unstained control samples, acquired as instrumental triplicates, and were processed using FlowJo V10 for Windows. At all times, samples were kept under dark conditions to avoid light-induced ROS production.

## 2.8 DFT calculations

The DFT calculations of electronic energy levels of the catalytic cycle were based on the crystal structure of complex 3. The method of the calculation was functional CAM-B3LYP<sup>31</sup> with basis set CEP-31G,<sup>32–34</sup> using Gaussian 16 software.<sup>35</sup> Ultrafine grid of integration was used in each case. The starting geometry was taken from X-ray data for 3, with an appropriate change of substituents for other systems. All given energy values are the result of the full geometry optimisation with subsequent frequency calculations. Optimisations were performed with modelling of water as solvent, within the continuous polarisation model with integral equation formalism variant (IEFPCM keyword of Gaussian). Grimme empirical corrections for dispersion were applied (keyword GD3). The optimisations were performed using the solvent-accessible surface option and the final energy was calculated with using solvent-excluding surface options (keywords surface = sas and ses, respectively). NAD<sup>+</sup> was modelled with an effective charge of –1, with two deprotonated phosphate groups; the same protonation of phosphate was used for NADH, giving an effective charge of –2.

## 2.9 DNA binding

The reactions of complex 4 (*ca.* 2 mM) with nucleobases (9-EtG and 5'-AMP) were studied typically by addition of an aqueous solution of nucleobase (3 mM, 1.5 mol equiv.) in 10% of d<sub>4</sub>-MeOD and 90% of D<sub>2</sub>O, pH\* 7.2  $\pm$  0.1, and monitored by <sup>1</sup>H NMR at 310 K. Solutions of double-helical calf thymus DNA (ct-DNA) at a concentration of 32  $\mu\text{g mL}^{-1}$  were incubated with complex 4 at *r*<sub>1</sub> value of 0.1 in 10 mM NaClO<sub>4</sub> at 310 K (*r*<sub>1</sub> is defined as the molar ratio of free ruthenium complex to nucleotide phosphates at the onset of incubation with DNA). The concentration of ruthenium associated with DNA in these samples was determined by flameless atomic absorption spectrometry (FAAS). The concentrations of DNA were determined by absorption spectrophotometry. Plasmid DNA pBR322 (28  $\mu\text{g mL}^{-1}$ ) and complex 4 in various molar ratios (*r*<sub>1</sub> = 0.05–1) were incubated in 0.01 M NaClO<sub>4</sub> at 310 K for 24 h in the dark.





Then the samples were directly mixed with the loading buffer and loaded onto a 1% agarose gel running at 298 K in the dark with Tris-acetate-EDTA (TAE) buffer and the voltage set at 25 V. No separation step was included before loading the samples into the gel to allow detection of potential noncovalent binding (if any). The gels were then stained with EtBr, followed by photography with a transilluminator.

## 3. Results and discussion

### 3.1 Synthesis and characterisation

Ru<sup>II</sup> complexes 1–6 were synthesised using a similar procedure to that reported for related complexes (Scheme 1).<sup>25</sup> Typically, triethylamine (4 mol equiv.) and ligands (*ca.* 2 mol equiv.) were added to a solution of  $[(\eta^6\text{-}p\text{-cym})\text{RuCl}_2]_2$  in degassed isopropanol, and the reaction was stirred under a N<sub>2</sub> atmosphere at 365 K for 12 h. All synthesised complexes were characterised by <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy, mass spectrometry (ESI-MS) and elemental analysis (CHN). A crystal of complex 3 suitable for X-ray analysis was obtained by diffusion of diethyl ether into a solution of the complex in methanol at ambient temperature. Selected bond lengths and angles for complex 3 are listed in Table 2. Crystallographic data are presented in Table S1,<sup>†</sup> and the structure of complex 3 is shown in Fig. 2. Complex 3 adopts a *pseudo*-octahedral geometry with the  $\eta^6$ -bonded aromatic ring occupying 3 coordination sites. The chelating ligand is deprotonated and bonded as a monoanionic bidentate ligand. The CH<sub>2</sub>CH<sub>2</sub>N-Et atoms from N,N' chelated ligand (C10 C11 N12 C13) were modelled as disordered over two positions whose occupancy refined to 86 : 14. Compared to reported ruthenium ethylenediamine complexes (either neutral or +1 charge),<sup>25,36,37</sup> the Ru–N<sup>−</sup> bond length (N9,

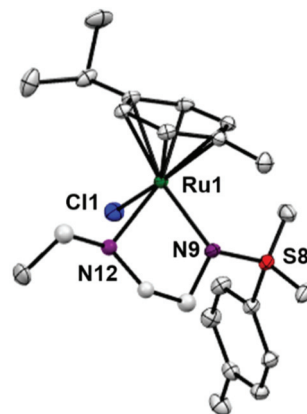


Fig. 2 ORTEP diagrams for complex 3. Ellipsoids are shown at the 50% probability level. All hydrogen atoms have been omitted for clarity.

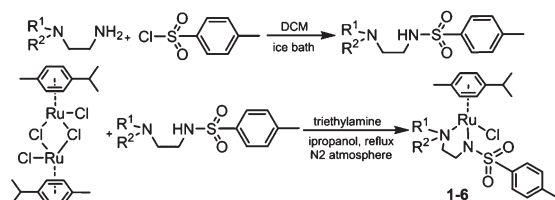
2.126(9)) is within the expected range of 2.11–2.14 Å,<sup>37</sup> but the Ru–N12 length (2.1702(11) Å) is longer than the neutral analogue  $[(\eta^6\text{-biph})\text{Ru}(\text{TsEn})\text{Cl}]$  (2.122(3) Å),<sup>25</sup> suggesting that the presence of *N*-ethyl substituent causes a slight weakening of this Ru–N bond. The remaining bond length and angles show no significant difference.

### 3.2 Hydrolysis and pK<sub>a</sub><sup>\*</sup> determination

The hydrolysis of complex 4 was studied by dissolving the Ru<sup>II</sup> complex in d<sub>4</sub>-MeOD/D<sub>2</sub>O (1.4 mM, 1 : 9 (v/v)). The <sup>1</sup>H NMR spectrum remained unchanged after 24 h and the hydrolysis was assumed to be rapid since the peaks could be assigned to the aqua Ru<sup>II</sup> species (4a) by comparison to those from the aqua species generated in a reaction with silver nitrate in D<sub>2</sub>O (1 mol equiv.). The pK<sub>a</sub><sup>\*</sup> (pK<sub>a</sub> value determined in deuterated solvent) of complex 4a was determined by a pH\* (meter reading) titration ranging from 2 to 12 by addition of NaOD or DNO<sub>3</sub> solutions as appropriate. Changes in the chemical shift of a tosyl <sup>1</sup>H NMR resonance were followed and the data were fitted to the Henderson–Hasselbalch equation, giving a pK<sub>a</sub><sup>\*</sup> value of 9.73 ± 0.06 (Fig. 3).

### 3.3 Kinetics of transfer hydrogenation reactions

The ratio of coenzyme NAD<sup>+</sup>/NADH greatly influences the intracellular potential and can drive many reactions *in vivo*.<sup>38</sup> The reduction of coenzyme nicotinamide adenine dinucleotide (NAD<sup>+</sup>) to NADH was investigated in an aqueous medium using complexes 1–6 as catalysts and sodium formate as the hydride source. Initially, the TH reactions were studied by UV-visible spectroscopy under conditions of pH 7.2 ± 0.1, 310 K and MeOH/H<sub>2</sub>O (1 : 9, v/v, Table 3); in all the cases, an increase in intensity of the band at 340 nm was observed, which is assignable to formation of NADH (Fig. S1, ESI<sup>†</sup>). The kinetics of conversion were also monitored by <sup>1</sup>H NMR at 310 K and pH\* 7.2 ± 0.1. The reactions were performed in a mixed solvent d<sub>4</sub>-MeOD/D<sub>2</sub>O (1 : 4, v/v), due to the poor aqueous solubility of complexes 5 and 6, although the presence of methanol in such reactions is known to enhance the reaction rate.<sup>25</sup>

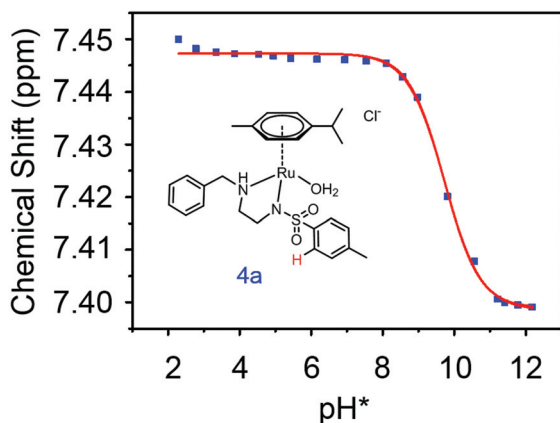


Scheme 1 Synthetic routes for diamine ligands and Ru<sup>II</sup> complexes 1–6.

Table 2 Selected bond lengths (Å) and angles (°) for complex 3

Bonds	Length/angle
Ru1–N9	2.1256(9)
Ru1–N12	2.1702(11)
Ru1–N12A	2.157(8)
Ru1–Cl1	2.4173(3)
Ru1–arene (centroid)	1.664
N9–Ru1–N12	78.74(4)
N9–Ru1–N12A	76.1(2)
N9–Ru1–Cl1	89.47(3)
N12–Ru1–Cl1	87.55(4)





**Fig. 3** Dependence of the  $^1\text{H}$  NMR chemical shift of a tosyl proton (red) on  $\text{pH}^*$  of aqua complex **4a**. The red curve is the best fit to the Henderson–Hasselbalch equation corresponding to a  $\text{pK}_a^*$  of  $9.73 \pm 0.06$ .

**Table 3** Turnover frequencies for transfer hydrogenation reactions using Ru complexes **1–6** as catalysts

Complex	$\text{R}^1, \text{R}^2$	$\text{TOF}^a$ ( $\text{h}^{-1}$ )	$\text{TOF}^b$ ( $\text{h}^{-1}$ )
<b>1</b>	Me, H	$2.97 \pm 0.04$	$4.0 \pm 0.3$
<b>2</b>	Me, Me	$3.9 \pm 0.1$	$4.1 \pm 0.1$
<b>3</b>	Et, H	$4.3 \pm 0.1$	$5.9 \pm 0.2$
<b>4</b>	Bz, H	$7.4 \pm 0.1$	$7.7 \pm 0.3$
<b>5</b>	4-F-Bz, H	$7.1 \pm 0.1$	$6.5 \pm 0.4$
<b>6</b>	Naph, H	$6.1 \pm 0.9$	$4.9 \pm 0.5$

<sup>a</sup> By UV-vis spectroscopy. <sup>b</sup> By NMR spectroscopy.

In general, the introduction of substituents on the terminal nitrogen improved the catalytic activity. The bulkier the substituents on the terminal nitrogen, the higher the TH reaction rate becomes. The turnover frequency reaches a maximum (*ca.*  $7.54 \text{ h}^{-1}$ ) when the substituent on the terminal N is benzyl (complex **4**), making it as efficient as the  $\text{Rh}^{\text{III}}$  complex  $[(\eta^5\text{-Cp}^*)\text{Rh}(\text{bipy})\text{Cl}]\text{PF}_6$ .<sup>16</sup> Interestingly, the TOF decreases when the substituent is *para*-fluoro-benzyl (complex **5**) or naphthalene (complex **6**), probably, because these ligands hamper the approach of  $\text{NAD}^+$  to the Ru centre. Compared to the en complex with unsubstituted nitrogens  $[(\eta^6\text{-biph})\text{Ru}(\text{en})\text{Cl}]\text{PF}_6$ , the turnover frequency of complex **4** is  $41\times$  higher,<sup>24</sup> and  $2.7\times$  higher compared to  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$ .<sup>25</sup>

The NH proton of the chelated diamine ligand appears to be essential for the TH reduction of ketones to alcohols;<sup>39</sup> normally,  $\text{Ru}^{\text{II}}$  catalysts for TH of ketones form 16-e intermediates.<sup>40,41</sup> It has been reported that a  $\text{Ru}^{\text{II}}$  complex with two *N*-alkyl groups (*R,R*)- $[(\eta^6\text{-benzene})\text{Ru}(\text{TsDPEN-Me}_2)\text{Cl}]$  exhibited poor catalytic reactivity in TH reaction of ketones.<sup>40</sup> However, complex **2**  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{Me}, \text{Me}))\text{Cl}]$  exhibited good catalytic activity towards the TH reduction of  $\text{NAD}^+$  to  $\text{NADH}$  ( $\text{TOF} = 4.1 \text{ h}^{-1}$ , Table 3), despite not having an NH proton, which suggests, as expected, that an N–H is not essential in the transfer reduction of  $\text{NAD}^+$  to  $\text{NADH}$ .

The dependence of the rate of catalysis on pH was determined. Six  $\text{pH}^*$  values ranging from 5 to 9 were studied for complex **4** at a mol ratio complex **4** :  $\text{NAD}^+$  : formate of 1 : 4 : 25, in the same mixed solvent at 310 K (Fig. S2, ESI†). The TOF was relatively insensitive to  $\text{pH}^*$  over the range  $\text{pH}^*$  6–8 (*ca.*  $7.5 \text{ h}^{-1}$ ), but decreased slightly at lower and higher  $\text{pH}^*$  ( $5.6 \text{ h}^{-1}$  at  $\text{pH}^*$  5,  $6.6 \text{ h}^{-1}$  at  $\text{pH}^*$  9).

The dependence of turnover frequency on the concentrations of sodium formate and  $\text{NAD}^+$  was also investigated for complex **4** in  $\text{d}_4\text{-MeOD}/\text{D}_2\text{O}$  (1 : 4) at 310 K. The mol ratio of complex **4** :  $\text{NAD}^+$  : formate was 1 : 4 : *X*, where *X* = 5, 10, 25, 50 and 100 (Fig. S3, ESI†). The TOF increased steadily from  $2.2 \text{ h}^{-1}$  to  $18.8 \text{ h}^{-1}$  as the concentration of formate was increased from 7 mM to 140 mM. Next the dependence of TOF on the  $\text{NAD}^+$  concentration was studied for mol ratio complex **4** :  $\text{NAD}^+$  : formate = 1 : *Y* : 25, where *Y* = 2, 6 and 10. The TOF was found to be independent of  $\text{NAD}^+$  concentration ( $7.7 \pm 0.5 \text{ h}^{-1}$ ).

The Michaelis–Menten kinetic behaviour is apparent from a plot of turnover frequency *versus* formate concentration. A reciprocal plot of turnover frequency *versus* formate concentration gave a Michaelis constant of  $K_M = 0.086 \text{ mM}$  (Fig. S3 and S4, ESI†). The maximum turnover frequency  $\text{TOF}_{\text{max}}$  for complex **4** ( $30.3 \text{ h}^{-1}$ ) is *ca.*  $5\times$  higher than for  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$  (complex **8**,  $\text{TOF}_{\text{max}} = 6.4 \text{ h}^{-1}$ )<sup>25</sup> and  $20\times$  higher than for the complex  $[(\eta^6\text{-hmb})\text{Ru}(\text{en})\text{Cl}]\text{PF}_6$  ( $\text{TOF}_{\text{max}} = 1.46 \text{ h}^{-1}$ ).<sup>24</sup> The much lower Michaelis–Menten constant ( $K_M = 0.086 \text{ mM}$ ) for the *N*-benzyl complex **4** indicates a stronger affinity of the complex for formate compared to  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn})\text{Cl}]$  ( $K_M = 27.8 \text{ mM}$ )<sup>25</sup> and  $[(\eta^6\text{-hmb})\text{Ru}(\text{en})\text{Cl}]\text{PF}_6$  ( $K_M = 58 \text{ mM}$ ).<sup>24</sup>

The maximum turnover frequency was observed at  $\text{pH}^*$  6 ( $\text{TOF}_{\text{max}} = 7.7 \text{ h}^{-1}$ ) (Fig. S2, ESI†). The TOF for complex **4** gradually decreased when the  $\text{pH}^*$  was raised above 6. Transfer hydrogenation was halted below  $\text{pH}^*$  4 because of the decomposition of the complex.

### 3.4 Antiproliferative activity and anticancer activity with formate

Ruthenium complexes have shown promise for their activity against various types of cancer cells.<sup>42</sup> The antiproliferative activity of complexes **1–6** towards A2780 human ovarian cancer cells was determined and compared with the clinically approved drug cisplatin, Fig. 4. The  $\text{IC}_{50}$  values (50% inhibition of cell growth) range from 1 to  $6.5 \mu\text{M}$  for complexes containing aromatic R substituents (**4–6**), whereas those containing aliphatic R substituents were less potent with  $\text{IC}_{50}$  values of  $12\text{--}31 \mu\text{M}$ . The complex  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{Bz}, \text{H}))\text{Cl}]$  (**4**) ( $\text{IC}_{50}$ ,  $1.0 \mu\text{M}$ ) has a potency similar to cisplatin in this cell line (CDDP,  $1.20 \pm 0.02 \mu\text{M}$ ). It is apparent that the presence of aromatic substituents on the chelated ligands of complexes **4–6** give rise to more potent cytotoxicity than aliphatic substituents in complexes **1–3**, most probably due to their higher lipophilicity.

Combination treatment with formate can greatly increase the antiproliferative activity of  $\text{Ru}^{\text{II}}$  arene sulfonyl diamine complexes, which offers a potential new strategy for cancer



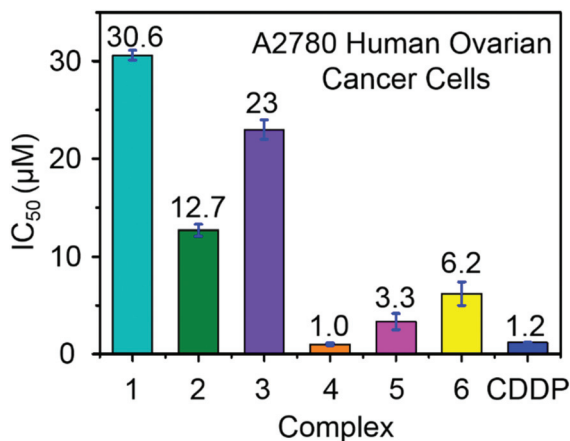


Fig. 4 Antiproliferative activity of Ru<sup>II</sup> complexes 1–6 and cisplatin towards A2780 human ovarian cancer cells.

treatment.<sup>15</sup> In this work, the antiproliferative activity of Ru<sup>II</sup> complexes in A2780 human ovarian cancer cells in the presence of sodium formate was studied (Fig. 5). Firstly, the cytotoxicity of sodium formate alone towards A2780 human ovarian cancer cells was investigated. No significant toxicity was found up to formate concentrations of 2 mM which is in agreement with the previous report.<sup>15</sup> Then, A2780 human ovarian cancer cells were coincubated with equipotent concentrations of complexes 1–6 ( $1/3 \times IC_{50}$ ) and three different concentrations of sodium formate (0.5, 1 and 2 mM) in order to observe the formate-concentration dependence of the cell viability. The antiproliferative activity of complexes 1–6 increased significantly upon coincubation with 2 mM formate. The formate-induced decrease in viability of A2780 cells ranged from 20% to 36% in the presence of complexes 1–6.

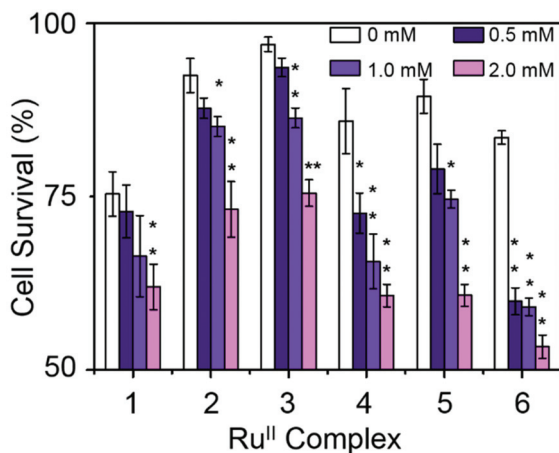


Fig. 5 Percentage of cell survival when equipotent concentrations of complexes 1–6 ( $1/3 \times IC_{50}$ ) were co-administered with different concentrations of sodium formate, *p*-values were calculated after a *t*-test against the negative control data (without sodium formate), \**p* < 0.05, \*\**p* < 0.01.

Interestingly for complex 6, a 28% decrease in cell viability was observed with only 0.5 mM formate present (Fig. 5, for percentage of viability decrease see Table S3, ESI<sup>†</sup>). The largest decrease of cell survival was 31% for complex 6 in the presence of 2 mM sodium formate, followed by 29% and 32% for the other two complexes with aromatic substituents, complexes 4 and 5, respectively. Complexes 1–3 with aliphatic functional groups showed an increase in potency of 18%, 21% and 22%, respectively.

### 3.5 Cell accumulation and relative hydrophobicity

Hydrophobicity and cellular accumulation are often important factors that play key roles in the potency of organometallic and other anticancer drugs.<sup>43</sup> The cellular accumulation, as an equilibrium between uptake and efflux, of ruthenium in A2780 human ovarian cancer cells after exposure to complexes 1–6 at their  $IC_{50}$  equipotent concentrations was determined by inductively coupled plasma mass spectrometry (ICP-MS) and is shown in Fig. 6.

Complex 4 gave the lowest cellular accumulation ( $0.52 \pm 0.08$  ng of Ru per  $10^6$  cells), while complex 6 with moderate anticancer activity, exhibited the highest extent of cell accumulation with  $4.5 \pm 0.2$  ng of Ru per  $10^6$  cells at  $IC_{50}$  concentration, 8.6× higher than complex 4. Complexes 1–3 and 5, gave rise to similar cell uptake  $2.4 \pm 0.3$  ng,  $1.2 \pm 0.2$  ng,  $3.0 \pm 0.2$  ng and  $1.3 \pm 0.2$  ng per  $10^6$  cells, respectively, following the order:  $4 < 2, 5 < 1 < 3 < 6$ .

The relative hydrophobicity of complexes 1–6 was determined by RP-HPLC. The more hydrophobic complexes have longer retention times on a reverse-phase  $C_{18}$  column.<sup>44</sup> To ensure solubility of the Ru<sup>II</sup> complexes in water, methanol was used as co-solvent (MeOH/H<sub>2</sub>O, 1 : 9 v/v) together with NaCl (50 mM) to suppress hydrolysis of the complexes. The HPLC solvents were also prepared with 50 mM NaCl (measurements see Fig. S5, ESI<sup>†</sup>). The resulting retention times are shown in Table 4, and follow the order: 1, 2, 3 < 4, 5 < 6. Complex 3 shows the shortest retention time (least hydrophobic) of

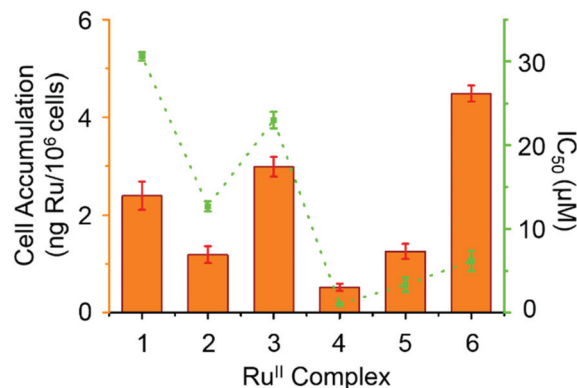


Fig. 6  $IC_{50}$  values (μM) for complexes 1–6 against A2780 human ovarian cancer cells (orange bars) and cellular accumulation of Ru in A2780 cancer cells at equipotent  $IC_{50}$  concentrations in the absence of sodium formate (in green).



**Table 4** Retention times ( $t_R$ ) of Ru<sup>II</sup> complexes 1–6 by RP-HPLC and cellular accumulation (at equipotent of IC<sub>50</sub> concentrations) in A2780 cells

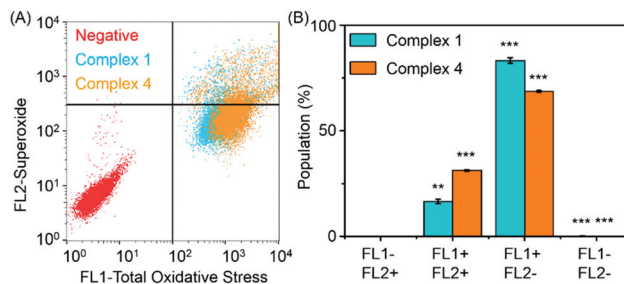
Complex	$t_R$ (min)	Cellular-Ru (ng per 10 <sup>6</sup> cells)
1	15.4 ± 0.9	2.4 ± 0.3
2	14.5 ± 0.3	1.2 ± 0.2
3	14.0 ± 0.3	3.0 ± 0.2
4	17.4 ± 0.2	0.52 ± 0.08
5	17.27 ± 0.08	1.3 ± 0.2
6	20 ± 1	4.5 ± 0.2

14.0 min, while complex 6 shows the longest retention time (most hydrophobic), 20.9 min.

It is evident from Table 4 that the Ru<sup>II</sup> complexes with aromatic substituents (complexes 4–6) exhibit higher hydrophobicity than complexes with aliphatic substituents (complexes 1–3). The most hydrophobic complex (6) shows the highest cell accumulation. Nonetheless, there is no linear correlation between the hydrophobicity of complexes 1–6 and their cellular accumulation. This has been observed before.<sup>45</sup> In these cases, the chemistry and the mechanism of action of each particular complex has a higher impact on the compound's anticancer activity than cellular accumulation *per se*. However, complex 4 has the lowest extent of cell uptake, but the most potent antiproliferative activity, suggesting that it is the chemical properties of the intracellular drug that are more important for activity than the total amount of Ru entering the cell. In general, a high hydrophobicity could facilitate interaction between the organometallic complex and cell membranes, and also correlate with the potency of the complex, but that is not always the case.<sup>43,45</sup>

### 3.6 ROS induction

Reactive oxygen species (ROS) are metabolic byproducts of aerobic respiration and are responsible for maintaining redox homeostasis in cells.<sup>46</sup> ROS also play a significant role in the mechanism of action of anticancer agents.<sup>47,48</sup> Some organometallic complexes, *e.g.* Ir and Os,<sup>49–51</sup> can generate high ROS levels or bursts of superoxide in cancer cells to induce cell apoptosis,<sup>49</sup> but by comparison, other complexes are known to induce cell death by reductive stress.<sup>15</sup> The levels of reactive oxygen species (ROS) were determined in A2780 human ovarian cancer cells for complexes 1 and 4 at IC<sub>50</sub> concentrations by flow cytometry fluorescence analysis (Fig. 7). This included the monitoring of H<sub>2</sub>O<sub>2</sub>, peroxy and hydroxyl radicals using a green probe, and superoxide levels using the orange channel. Induction of total ROS and superoxide were determined in A2780 cells after 24 h exposure to complexes 1 and 4 when compared to the negative untreated control. The populations of cells that show high fluorescence in both FL-1 and FL-2 channels (both high total ROS and high superoxide generation) for complexes 1 and 4 are 16.5 ± 1.0% and 31.3 ± 0.3%, respectively, which indicates a higher induction of superoxide by complex 4. Remarkably, the total increase of the population in the high FL-1 green channel shows that the levels of total



**Fig. 7** ROS in A2780 cells induced by complexes 1 and 4, FL1 channel detects total oxidative stress, and FL2 channel detects superoxide production. (A) Induction of ROS by complexes 1 and 4. (B) Four different populations induced by complexes 1 and 4 at equipotent IC<sub>50</sub> concentrations. *p*-Values were calculated after a *t*-test against the negative control data, \*\**p* < 0.01, \*\*\**p* < 0.001.

ROS are induced in the majority, if not in all, of the cell population. These ROS may play a major role in killing the cancer cells (Table S4, ESI<sup>†</sup>).

### 3.7 DNA related binding for complex 4

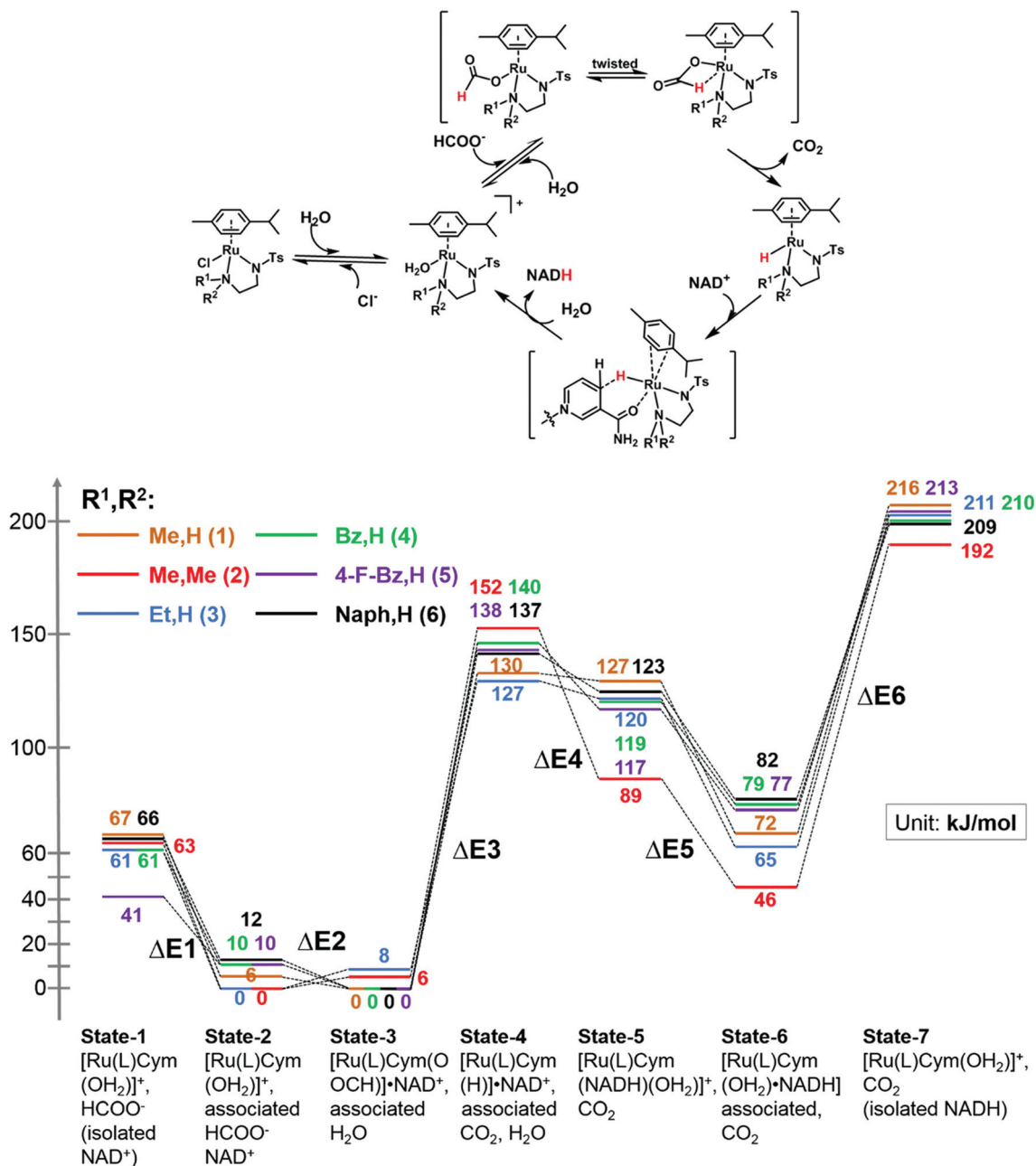
The interaction of complex 4 with DNA nucleobase models: 9-ethylguanine (9-EtG) and adenosine 5'-monophosphate (5'-AMP) was studied by <sup>1</sup>H NMR spectroscopy (Fig. S6, ESI<sup>†</sup>). The reactions were performed by adding nucleobase solution (3 mM in D<sub>2</sub>O) to Ru<sup>II</sup> complex solution (2 mM in 10% d<sub>4</sub>-MeOD/90% D<sub>2</sub>O) at 310 K, to give a final 1.5 : 1 mol ratio. The formation of adduct 4-9-EtG was confirmed by following the new set of peaks, and up to 90% yield of adduct was obtained when 1.5 mol equiv. nucleobase solution was added. However, no adduct was found when 1.5 mol equiv. of 5'-AMP was added to complex 4, even after 24 h incubation at 310 K. Reactions of double-helical calf thymus DNA (ct-DNA, 32 μg mL<sup>-1</sup>) and plasmid DNA pBR322 (28 μg mL<sup>-1</sup>) with complex 4 in various molar ratios ( $r_1 = 0.05-1$ ,  $r_1$  is the molar ratio of free Ru complex to nucleotide phosphates at the onset of incubation with DNA) were studied. Very low amounts of ruthenium (5–7% of initial Ru) were found in the samples of DNA treated with complex 4 for 24 h. No significant changes in the mobilities of supercoiled (sc) or open circular (oc) form of plasmid DNA were observed even when incubated with high concentration of complex 4 ( $r_1 = 1$ , Fig. S7, ESI<sup>†</sup>). DNA is thought to be a cellular target for the en complex 7 (Fig. 1).<sup>52,53</sup> However, for the substituted-en complex studied here, no obvious unwinding of DNA was observed after coincubation of ct-DNA with complex 4, suggesting that binding is weak, nor changes in the ratio of sc and oc forms of plasmid DNA, suggesting that complex 4 does not cleave DNA.

### 3.8 DFT calculations

We modelled the catalytic cycle by considering seven states of the reaction: (1) the initial aqua complex  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))(\text{OH}_2)]^+$  and formate (with isolated NAD<sup>+</sup>); (2)  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))(\text{OH}_2)]^+$  interacting intermolecularly with NAD<sup>+</sup> and formate; (3)  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))$







**Fig. 8** (Top) Reduction cycle for conversion of NAD<sup>+</sup> to 1,4-NADH via transfer hydrogenation with formate as the hydride donor. (Bottom) DFT energy profile for the formation of Ru formate species, Ru hydride complex and hydride transfer from ruthenium; brown line, complex 1; red line, complex 2; blue line, complex 3; green line, complex 4; purple line, complex 5; black line, complex 6. Sets of calculated structures of states 1–7 are supplied in the ESI and illustrated graphically in Fig. S8† for complex 2. To calculate the energy of states 1 and 7, the energies of the states represented in pdb files 1 and 7 were added to the energies calculated for NAD<sup>+</sup> and NADH.

(HCOO<sup>-</sup>)·NAD<sup>+</sup> and water; (4) [(<sup>η</sup><sup>6</sup>-*p*-cym)Ru(TsEn(R<sup>1</sup>,R<sup>2</sup>))H]·NAD<sup>+</sup> and water and CO<sub>2</sub>; (5) [(<sup>η</sup><sup>2</sup>-*p*-cym)Ru(TsEn(R<sup>1</sup>,R<sup>2</sup>))(OH<sub>2</sub>)NADH] and CO<sub>2</sub>; (6) [(<sup>η</sup><sup>6</sup>-*p*-cym)Ru(TsEn(R<sup>1</sup>,R<sup>2</sup>))(OH<sub>2</sub>)]NADH and CO<sub>2</sub>; (7) [(<sup>η</sup><sup>6</sup>-*p*-cym)Ru(TsEn(R<sup>1</sup>,R<sup>2</sup>))(OH<sub>2</sub>)<sup>+</sup>], water and CO<sub>2</sub>, and isolated NADH (Fig. 8).

For state 5, with ring-slipped coordinated <sup>η</sup><sup>2</sup>-*p*-cymene, the introduction of water into coordination sphere was necessary, while highly distorted complexes without coordinated water

were found *ca.* 100 kJ mol<sup>-1</sup> higher in energy. It is notable that the Ru atoms of all complexes in state 5 are coordinated to the amide oxygen atom of NADH, while only weakly bound to the (hydridic) CH<sub>2</sub> of NADH, giving a Ru–H distance of 3.11–3.12 Å for R<sup>1</sup>,R<sup>2</sup> = Me,H (1); Et,H (3); Naph,H (6) and 3.06–3.07 Å for R<sup>1</sup>,R<sup>2</sup> = Bz,H (4) and 4-F-Bz,H (5). For R<sup>1</sup>,R<sup>2</sup> = Me,Me (2) the calculations revealed a true bonding of the (hydridic) CH<sub>2</sub>, with a Ru–H distance of 1.99 Å. The results obtained are



shown in Fig. 8. Four general conclusions can be drawn from these data: (a) there is a strong interaction between the  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))(\text{OH}_2)]^+$  cation and  $\text{NAD}^+$  and  $\text{NADH}$  molecules, leading to a stabilisation of the cationic form by 60–70  $\text{kJ mol}^{-1}$  for  $\text{NAD}^+$  (41  $\text{kJ mol}^{-1}$  for  $\text{R}^1, \text{R}^2 = 4\text{-F-Bz, H}$  (5)) and 130–150  $\text{kJ mol}^{-1}$  for  $\text{NADH}$ ; (b) depending on the *N*-substituent, the species of the lowest energy is either  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))(\text{HCOO}^-)]\cdot\text{NAD}^+$  ( $\text{R}^1, \text{R}^2 = \text{Me, H}$  (1);  $\text{Bz, H}$  (4);  $4\text{-F-Bz, H}$  (5) and  $\text{Naph, H}$  (6)) or  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{TsEn}(\text{R}^1, \text{R}^2))(\text{OH}_2)]\cdot\text{NAD}^+$  ( $\text{R}^1, \text{R}^2 = \text{Et, H}$  (3) and  $\text{Me, Me}$  (2)), the difference between them being only 6–12  $\text{kJ mol}^{-1}$ ; (c) the effective  $\text{NADH}$ -hydride coordination for bulky  $\text{R}^1, \text{R}^2 = \text{Me, Me}$  (2) lowers the energy, relative to the state of lowest energy, of the species with coordinating  $\text{NADH}$  by 30–40  $\text{kJ mol}^{-1}$ , compared to other complexes; (d) the formation of the state with the  $\text{Ru-H}$  hydride bond, including the twist of formate and the elimination of carbon dioxide, corresponds to the highest energy step. These four factors seem all to influence the turnover. The energy barriers and optimized structures for the seven states of complex 2 in the cycle with  $\text{NAD}^+$  are listed in Table S5 and illustrated graphically in Fig. S8.† The structure files for the remaining complexes are supplied as ESI.†

## 4. Conclusions

In this work, we have synthesised a series of new  $\text{Ru}^{\text{II}}$  complexes of the type  $[(\eta^6\text{-}p\text{-cym})\text{Ru}(\text{N}, \text{N}')\text{Cl}]$  where  $\text{N}, \text{N}'$  are monosulfonamide chelating ligands derived from tosyl ethylenediamine, with either alkyl ( $\text{Me, H}$  (1);  $\text{Me, Me}$  (2);  $\text{Et, H}$  (3)) or aryl ( $\text{Bz, H}$  (4);  $4\text{-F-Bz, H}$  (5);  $\text{Naph, H}$  (6)) substituents on the terminal *N*. These substituents have a significant effect on the rate of transfer hydrogenation of coenzyme  $\text{NAD}^+$  with formate as hydride donor as determined by NMR and UV-vis spectroscopy. In general, the bulkier aromatic substituents gave rise to faster hydrogenation rates (Table 3). DFT calculations provided insight into the mechanism of hydride transfer from formate to  $\text{NAD}^+$  involving initial coordination of formate followed by transfer of hydride to ruthenium and then to  $\text{NAD}^+$  with release of  $\text{CO}_2$ . The calculations suggested a preorganization of the initial aqua complex, formate and  $\text{NAD}^+$  involving T-shaped adenosine NH-tosyl stacking, H-bonding of the NH of the chelated ligand and phosphate O of  $\text{NAD}^+$ , H-bonding between formate and water, and between formate and the pyridine ring of  $\text{NAD}^+$ . They also indicated strong interactions with  $\text{NADH}$  involving T-shaped adenosine NH-tosyl stacking, as well as H-bonds to phosphate and (hydridic)  $\text{CH}_2$ -tosylate O (Fig. S8, ESI†).

To investigate the possibility of achieving transfer hydrogenation mediated by formate in cells, we investigated the effect of formate on the antiproliferative activity of these complexes towards human ovarian cancer cells. In each case a dose-dependent increase in potency of the complexes (20–36%) was observed with increasing formate concentration over a range of non-toxic formate concentrations (0–2 mM). The complexes with aromatic substituents were the most

potent, the benzyl complex 4 being as potent as the anticancer drug cisplatin (Fig. 6). In general, the most hydrophobic complexes were found to be the most biologically active. However, the activity does not correlate closely with total cell accumulation of Ru or with hydrophobicity (Table 3). Although DNA can be a target for related arene  $\text{Ru}^{\text{II}}$  diamine complexes, it does not appear to be a target for these sulfonyl-en  $\text{Ru}^{\text{II}}$  catalysts since we observe very weak binding to both calf thymus and plasmid DNA (Fig. S7, ESI†).

We showed that complexes 1 and 4 can generate high levels of ROS in A2780 human ovarian cancer cells, especially 4, the most potent complex. This is consistent with interference in cellular redox pathways and possible attack on  $\text{NAD}^+$  when sodium formate is present. The enhancement of anticancer activity by low non-toxic dose of formate might be clinically useful since it introduces a new mechanism of activity which does not involve DNA attack, unlike the clinical drug cisplatin. Such a regime might therefore avoid some unwanted side-effects. Formate itself is a natural biochemical molecule enriched in some cancer cells.<sup>54</sup> However, more work remains to be done to investigate possible intracellular catalysis, especially since a range of metabolites might readily poison these catalysts in cells.

## Author contributions

Feng Chen, Joan J. Soldevila-Barreda, Abraha Habtemariam and Peter J. Sadler designed the experiments and interpreted data.

Feng Chen carried out synthesis and characterisation of ligands and complexes, investigated hydrolysis, determined the  $\text{pK}_a$  value, and TH turnover frequencies.

Isolda Romero-Canelón and Ji-Inn Song carried out the cell antiproliferative screening and related biochemical assays.

Guy J. Clarkson carried out the X-ray crystallography.

Juliusz A. Wolny and Volker Schünemann carried out all the DFT calculations.

Jana Kasparkova and Viktor Brabec carried out DNA binding studies.

James P. C. Coverdale carried out metal analyses by ICP-OES and ICP-MS, and related biological and biochemical assays.

All authors contributed to the writing of the paper.

## Conflicts of interest

The authors declare no conflicts of interest.

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## References

- U. Eisner and J. Kuthan, *Chem. Rev.*, 1972, **72**, 1–42.
- J. Gębicki, A. Marcinek and J. Zielonka, *Acc. Chem. Res.*, 2004, **37**, 379–386.
- A. McSkimming and S. B. Colbran, *Chem. Soc. Rev.*, 2013, **42**, 5439–5488.
- S. Fukuzumi and T. Suenobu, *Dalton Trans.*, 2013, **42**, 18–28.
- S. M. Barrett, C. L. Pitman, A. G. Walden and A. J. M. Miller, *J. Am. Chem. Soc.*, 2014, **136**, 14718–14721.
- M. C. Carrión, F. Sepúlveda, F. A. Jalón, B. R. Manzano and A. M. Rodríguez, *Organometallics*, 2009, **28**, 3822–3833.
- Y. Maenaka, T. Suenobu and S. Fukuzumi, *J. Am. Chem. Soc.*, 2012, **134**, 9417–9427.
- A. Wolfson, C. Dlugy, Y. Shotland and D. Tavor, *Tetrahedron Lett.*, 2009, **50**, 5951–5953.
- R. Ruppert, S. Herrmann and E. Steckhan, *Tetrahedron Lett.*, 1987, **28**, 6583–6586.
- E. Steckhan, S. Herrmann, R. Ruppert, E. Dietz, M. Frede and E. Spika, *Organometallics*, 1991, **10**, 1568–1577.
- E. Steckhan, S. Herrmann, R. Ruppert, J. Thömmes and C. Wandrey, *Angew. Chem., Int. Ed. Engl.*, 1990, **29**, 388–390.
- V. D. Westerhausen, S. Herrmann, W. Hummel and E. Steckhan, *Angew. Chem., Int. Ed. Engl.*, 1992, **31**, 1529–1531.
- R. T. Hembre and S. McQueen, *J. Am. Chem. Soc.*, 1994, **116**, 2141–2142.
- C. Wang, B. Villa-Marcos and J. Xiao, *Chem. Commun.*, 2011, **47**, 9773–9785.
- J. J. Soldevila-Barreda, I. Romero-Canelón, A. Habtemariam and P. J. Sadler, *Nat. Commun.*, 2015, **6**, 6582.
- H. C. Lo, O. Buriez, J. B. Kerr and R. H. Fish, *Angew. Chem., Int. Ed.*, 1999, **38**, 1429–1432.
- H. C. Lo, C. Leiva, O. Buriez, J. B. Kerr, M. M. Olmstead and R. H. Fish, *Inorg. Chem.*, 2001, **40**, 6705–6716.
- J. M. O'Connor and C. P. Casey, *Chem. Rev.*, 1987, **87**, 307–318.
- K. T. Oppelt, J. Gasiorowski, D. A. M. Egbe, J. P. Kollender, M. Himmelsbach, A. W. Hassel, N. S. Sariciftci and G. Knör, *J. Am. Chem. Soc.*, 2014, **136**, 12721–12729.
- V. Ganesan, D. Sivanesan and S. Yoon, *Inorg. Chem.*, 2017, **56**, 1366–1374.
- S. Hashiguchi, A. Fujii, J. Takehara, T. Ikariya and R. Noyori, *J. Am. Chem. Soc.*, 1995, **117**, 7562–7563.
- A. Fujii, S. Hashiguchi, N. Uematsu, T. Ikariya and R. Noyori, *J. Am. Chem. Soc.*, 1996, **118**, 2521–2522.
- J. P. C. Coverdale, I. Romero-Canelón, C. Sanchez-Cano, G. J. Clarkson, A. Habtemariam, M. Wills and P. J. Sadler, *Nat. Chem.*, 2018, **10**, 347–354.
- Y. K. Yan, M. Melchart, A. Habtemariam, A. F. A. Peacock and P. J. Sadler, *J. Biol. Inorg. Chem.*, 2006, **11**, 483–488.
- J. J. Soldevila-Barreda, P. C. A. Bruijninx, A. Habtemariam, G. J. Clarkson, R. J. Deeth and P. J. Sadler, *Organometallics*, 2012, **31**, 5958–5967.
- S. Betanzos-Lara, Z. Liu, A. Habtemariam, A. M. Pizarro, B. Qamar and P. J. Sadler, *Angew. Chem., Int. Ed.*, 2012, **51**, 3897–3900.
- A. Habtemariam, M. Melchart, R. Fernández, S. Parsons, I. D. H. Oswald, A. Parkin, F. P. A. Fabbiani, J. E. Davidson, A. Dawson, R. E. Aird, D. I. Jodrell and P. J. Sadler, *J. Med. Chem.*, 2006, **49**, 6858–6868.
- J. E. D. Martins, G. J. Clarkson and M. Wills, *Org. Lett.*, 2009, **11**, 847–850.
- X. Li, L. Li, Y. Tang, L. Zhong, L. Cun, J. Zhu, J. Liao and J. Deng, *J. Org. Chem.*, 2010, **75**, 2981–2988.
- V. Vichai and K. Kirtikara, *Nat. Protoc.*, 2006, **1**, 1112–1116.
- D. T. T. Yanai and N. Handy, *Chem. Phys. Lett.*, 2004, **393**, 51–57.
- W. J. Stevens, H. Basch and M. Krauss, *J. Phys. Chem.*, 1984, **81**, 6026–6033.
- W. J. Stevens, M. Krauss, H. Basch and P. G. Jasien, *Can. J. Chem.*, 1992, **70**, 612–630.
- T. R. S. Cundari, *J. Chem. Phys.*, 1993, **98**, 5555–5565.
- M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision A. 03*, Gaussian, Inc., Wallingford CT, 2016.
- R. E. Morris, R. E. Aird, M. P. Socorro, H. Chen, J. Cummings, N. D. Hughes, S. Parsons, A. Parkin, G. Boyd, D. I. Jodrell and P. J. Sadler, *J. Med. Chem.*, 2001, **44**, 3616–3621.
- A. F. A. Peacock, A. Habtemariam, R. Fernández, V. Walland, F. P. A. Fabbiani, S. Parsons, R. E. Aird, D. I. Jodrell and P. J. Sadler, *J. Am. Chem. Soc.*, 2006, **128**, 1739–1748.



- 38 J. Zhang, A. Pierick, H. M. Rossum, R. M. Seifar, C. Ras, J. Daran, J. J. Heijnen and S. A. Wahl, *Sci. Rep.*, 2015, **5**, 12846.
- 39 M. Yamakawa, H. Ito and R. Noyori, *J. Am. Chem. Soc.*, 2000, **122**, 1466–1478.
- 40 R. Soni, F. K. Cheung, G. C. Clarkson, J. E. D. Martins, M. A. Graham and M. Wills, *Org. Biomol. Chem.*, 2011, **9**, 3290–3294.
- 41 X. Wu, X. Li, W. Hems, F. King and J. Xiao, *Org. Biomol. Chem.*, 2004, **2**, 1818–1821.
- 42 Z. Adhireksan, G. E. Davey, P. Campomanes, M. Groessl, C. M. Clavel, H. Yu, A. A. Nazarov, C. H. F. Yeo, W. H. Ang, P. Droge, U. Rothlisberger, P. J. Dyson and C. A. Davey, *Nat. Commun.*, 2014, **5**, 3462.
- 43 A. J. Millett, A. Habtemariam, I. Romero-Canelón, G. J. Clarkson and P. J. Sadler, *Organometallics*, 2015, **34**, 2683–2694.
- 44 T. Bugarcic, O. Nováková, A. Halámiková, L. Zerzánková, O. Vrána, J. Kašpárková, A. Habtemariam, S. Parsons, P. J. Sadler and V. Brabec, *J. Med. Chem.*, 2008, **51**, 5310–5319.
- 45 M. Hanif, A. A. Nazarov, C. G. Hartinger, W. Kandioller, M. A. Jakupec, V. B. Arion, P. J. Dyson and B. K. Keppler, *Dalton Trans.*, 2010, **39**, 7345–7352.
- 46 A. T. Dharmaraja, *J. Med. Chem.*, 2017, **60**, 3221–3240.
- 47 D. Trachootham, J. Alexandre and P. Huang, *Nat. Rev. Drug Discovery*, 2009, **8**, 579–591.
- 48 J. Watson, *Open Biol.*, 2013, **3**, 120144.
- 49 Z. Liu, I. Romero-Canelón, A. Habtemariam, G. J. Clarkson and P. J. Sadler, *Organometallics*, 2014, **33**, 5324–5333.
- 50 Z. Liu, I. Romero-Canelón, B. Qamar, J. M. Hearn, A. Habtemariam, N. P. E. Barry, A. M. Pizarro, G. J. Clarkson and P. J. Sadler, *Angew. Chem., Int. Ed.*, 2014, **53**, 3941–3946.
- 51 Y. Fu, M. J. Romero, A. Habtemariam, M. E. Snowden, L. Song, G. J. Clarkson, B. Qamar, A. M. Pizarro, P. R. Unwin and P. J. Sadler, *Chem. Sci.*, 2012, **3**, 2485–2494.
- 52 H. Chen, J. A. Parkinson, S. Parsons, R. A. Coxall, R. O. Gould and P. J. Sadler, *J. Am. Chem. Soc.*, 2002, **124**, 3064–3082.
- 53 G. Gasser, I. Ott and N. Metzler-Nolte, *J. Med. Chem.*, 2010, **54**, 3–25.
- 54 P. M. Tedeschi, E. K. Markert, M. Gounder, H. Lin, D. Dvorzhinski, S. C. Dolfi, L. L. Y. Chan, J. Qiu, R. S. DiPaola, K. M. Hirshfield, L. G. Boros, J. R. Bertino, Z. N. Oltvai and A. Vazquez, *Cell Death Dis.*, 2013, **4**, 877.

