



Cite this: *RSC Adv.*, 2020, 10, 10673

Experimental and DFT studies of sulfadiazine 'piano-stool' Ru(II) and Rh(III) complexes†

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While sulfadiazine (HL^{SZ}) is extensively used to elaborate complexes of intriguing biological applications (e.g. topical antibiotic silvadene; silver sulfadiazine), the molecular structure modification of sulfadiazine or even other sulfa drugs by coordination to either η^5 -cymene Ru(II) or η^5 -Cp* Rh(III) motif has not been investigated. Here, half-sandwich organoruthenium(II) and organorhodium(III) compounds of the type $[(\eta^5\text{-}p\text{-cymene})\text{Ru}(\text{L}^{\text{SZ}})_2]$ (**1**) and $[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2]$ (**2**) are synthesized, characterized and evaluated for their potential antimicrobial activity. Spectroscopic and single crystal X-ray analysis showed that L^{SZ} is coordinated to Rh(III) via both the sulfonamide and pyrimidine nitrogen atoms forming "piano-stool" geometry. In **2**, the NMR equivalence clearly pointed to participation of two L^{SZ} molecules in a fluxional process in which the third bond of the base of the stool is oscillating between two equivalent sulfonamide nitrogen atoms. While **1** was biologically inactive, complex **2** was potent against Gram-positive bacteria, *Candida albicans* and *Cryptococcus neoformans*. Hen white egg lysozyme (HEWL), a model protein, reacted covalently with **2** via the loss of one L^{SZ} molecule, while compound **1** decomposed during the interaction with that protein.

Received 4th February 2020
Accepted 28th February 2020

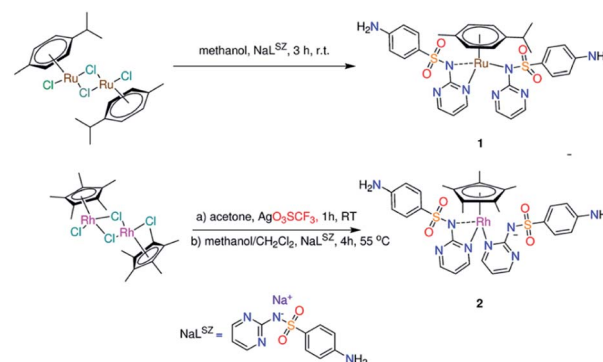
DOI: 10.1039/d0ra01085e

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Introduction

Sulfonamide derivatives are extensively used as antibacterial agents because they are cheap, have low toxicity and exhibit excellent activity against several bacterial diseases.¹ They slow down or stop bacterial growth in infected systems or in wounds without causing significant toxicity to the body tissues.² In other words, Sulfonamides are bacteriostatic rather than bactericidal. Sulfonamides interfere with the conversion of *p*-amino benzoic acid into an essential nutrient necessary for the bacterial multiplication.³ In view of growing bacterial resistance towards Sulfonamides as well as the appearance of more effective modern antibiotics, the importance of Sulfonamides has been greatly reduced. Sulfadiazine(4-amino-*N*-2-pyrimidinylbenzene sulphonamide, HL^{SZ}) was introduced in 1940 for curing of the bacterial infections in humans.⁴ It is used in combination therapy with pyrimethamine for treatment of chloroquine-resistant malaria parasite because of its long half-life time in the blood.⁵ From structural point of view, the presence of potential coordination sites in NaL^{SZ} (–NH₂, sulfonyl oxygen, sulfonamide (N_{sfd}) and pyrimidine nitrogen (N_{py})) (Scheme 1) makes it versatile ligand in the field of coordination chemistry.

Modified pharmacological and toxicological properties were observed upon coordination of HL^{SZ} to metal ions and formation of metal complexes. Zinc(II) complex of HL^{SZ} was used to inhibit the bacterial infection in burned animals,⁶ while silvadene(2-sulfanilamido-pyrimidine silver) was applied for treatment of topical burn.⁷ For HL^{SZ}, several potential coordination modes were reported,^{6–13} in which HL^{SZ} coordinated to Mⁿ⁺ as mono-, bi- and tridentate ligand. Normally, HL^{SZ} deprotonates during the interaction with Mⁿ⁺ ions. The crystallographic data of silvadene showed a polymeric arrangement structure in which Ag(I) is coordinated to tridentate L^{SZ} via N_{py}, N_{sfd} and OSO₂.⁷ For Zn(II) sulfadiazine complex, Zn(II) is coordinated to three different molecules of L^{SZ} in a tetrahedral



Scheme 1 Synthesis of compounds **1** and **2**.

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† Electronic supplementary information (ESI) available. CCDC 1940322. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ra01085e



polymeric arrangement. The ligands differently behaved towards Zn(II); the first L^{SZ} molecule binds Zn(II) via N_{pym} and sulfonyl oxygen, the second L^{SZ} interacts with Zn(II) via N_{pym} atom and the third ligand molecule coordinates to Zn(II) through N_{sfid}.⁶ Single-crystal X-ray analysis and spectroscopic characterization of other sulfadiazine complexes with Fe(III),⁸ Co(II),⁹ Ni(II),¹⁰ Cu(II),¹⁰ Cd(II),¹¹ Pt(II)¹² and Hg(II)¹³ ions were reported. For [Cu(NH₃)(L^{SZ})₂(OH₂)],^{10a} the crystal data showed that Cu(II) ion is surrounded by two L^{SZ} in a distorted N₄O₁ square-pyramidal environment; the base of the pyramid is occupied by a given *N,N*-bidentate L^{SZ}, OH₂, and NH₃, and its top is occupied with N_{pym} of the adjacent L^{SZ} molecule.

In comparison with free HL^{SZ}, the effect of complex formation on the antibacterial activity of HL^{SZ} was evaluated.^{10,14} No negative effect on the antibacterial activity of HL^{SZ} (against *Staphylococcus aureus*, *Klebsiella pneumoniae* and *Escherichia coli*) was detected by association to Cu(II) ions.^{10a} Screening of Cu(II)-L^{SZ} complexes against *Staphylococcus aureus* ATCC29213 and *Escherichia coli* ATCC 25922 indicated that the complexes were less effective than HL^{SZ}.^{10a} The -SO₂NH group of HL^{SZ} is essential for the antibacterial activity. Exchange of the ionizable proton with the metal ion may lead to diminishing of the toxicity. In other words, the truly active sulfonamide species is the ionic form; Sulfonamide drugs penetrate the bacterial cells in the neutral form and once they cross the cell wall, their toxicities may be attributed to the ionized form.¹⁴

In the present contribution, we disclose first examples of sulfa drugs based half-sandwich Ru(II) (**1**)¹⁵ and Rh(III) (**2**) organometallic compounds (Scheme 1) having either *p*-cymene or pentamethylcyclopentadienyl at the seat of piano-stool geometry. Although Sulfonamides constituted an important class of the antimicrobial agents in the world for a long time, it has not been used in synthesis of organometallic arene compounds.

The previous experimental evidences indicated that 98% of the overall administrated ruthenium, in the blood plasma, is in a protein bound form.¹⁶ The binding of cytotoxic Ru(II) complexes to the proteins may influence their toxicity, bio-distribution as well as drug delivery. Glutathione *S*-transferase P1-1, transferrin and albumin were known as probable final targets for Ru(II) complexes.¹⁷ These arguments encouraged us to study and explore the interactions between the synthesized organometallic compounds and hen white egg lysozyme (HEWL) by means of ESI-MS. HEWL was widely studied in the context of metalation of proteins because of its small size and suitability to be studied by mass spectrometry. In addition, HEWL is a perfect protein to crystallize and thus suitable for X-ray crystallography analysis. Complexes (**1**, **2**) and the metal precursors; [(η⁶-*p*-Cym)Ru(μ-Cl)Cl]₂ and [(η⁵-C₅Me₅)RhCl]₂(μ-Cl)₂ were evaluated for their inhibitory effect against the following microbes; Gram-positive bacterium (*Staphylococcus aureus* ATCC 43300), Gram-negative bacteria (*Escherichia coli* ATCC 25922, *Klebsiella pneumoniae* ATCC 700603, *Acinetobacter baumannii* ATCC 19606, *Pseudomonas aeruginosa* ATCC700603) and fungi (*Candida albicans* ATCC 90028 and *Cryptococcus neoformans* var. *grubii* H99; ATCC 208821). The complexes were assessed for their blood compatibility with the cell components

and cell viability against non-malignant cells with rapid proliferation (noncancerous human embryonic kidney cells (HEK293)).

Results and discussion

Synthesis and characterization

“Piano-stool” complex of type [(η⁶-*p*-Cym)Ru(L^{SZ})₂] (**1**) (Scheme 1) was prepared from stirring of sodium sulfadiazine (NaL^{SZ}) and [(η⁶-*p*-Cym)Ru(μ-Cl)Cl]₂ (ref. 18) in methanol at room temperature for 3 h. The proposed structure of **1**, isolated as air-stable yellow powder in 84% yield, was elucidated by elemental analysis, ESI-MS, IR and NMR (¹H, ¹³C, {¹H, ¹H} COSY, and {¹H, ¹³C} HSQC) (Fig. S1–S4†). In particular, the elemental composition, ESI-MS as well as the integral values of the ¹H NMR resonances confirm isolation of a 1 : 1 : 2 (M : *p*-Cym : L^{SZ}) complex. In the positive potential mode, the ESI-MS spectrum showed a unique peak at *m/z* = 735.1102, having a ruthenium isotope envelope, which may be allocated for the suggested mono-cationic monomeric complex of **1**. The ¹H NMR spectrum (in [D₆] DMSO) of free NaL^{SZ} shows a singlet signal at δ = 5.30 ppm (–NH₂). The signals of the pyrimidine ring are shown as triplet and doublet at δ = 6.34 and 8.06 ppm, respectively. The two doublets at δ = 6.46 and 7.48 ppm are allocated to the phenyl ring.¹⁹ The ¹H NMR signals of NaL^{SZ} are shifted downfield upon coordination to (η⁶-*p*-Cym)Ru. The well-known ¹H NMR pattern of *p*-Cym moiety is shown in the spectrum (Fig. S2†) of **1** as four doublets (at δ = 6.03, 5.89, 5.59 and 5.52 ppm) for the aryl-CH protons, a septet (at δ = 2.63 ppm) for CH(CH₃)₂, singlet methyl (at δ = 1.96 ppm) as well as two doublet methyl signals (centred at δ = 1.01 and 0.94 ppm).

The ¹³C NMR spectrum of **1** displayed the expected resonances for the *p*-Cym moiety (Fig. S2†). As previously reported, HL^{SZ} ligand exhibited different modes of coordination to Mⁿ⁺ and might be interacted as mono-, bi- and tridentate ligands in either monomeric or polymeric arrangements.^{6–13} As the *p*-Cym occupies the top of the stool of **1** by three arms, the remaining three arms around Ru(II) ion should be occupied by two deprotonated L^{SZ} molecules. Therefore, the tridentate mode of coordination for L^{SZ} should be excluded. Alternatively, solution [¹H–¹⁵N] heteronuclear multiple bond coherence (HMBC) NMR spectra of NaL^{SZ} and **1** were recorded to evaluate if the primary amino group interacts with Mⁿ⁺ ion or not. The Δ¹⁵N_{coord} parameter shows the real influence of nitrogen coordination effect; the difference between the NMR resonances of the free nitrogen and metal–nitrogen bond. The position of the NH₂ signal (δ = –313 ppm) in [¹H–¹⁵N] NMR spectrum of NaL^{SZ} doesn't change upon formation of **1**, excluding the role of NH₂ in the complex formation (Fig. S2 and S4†).

To get more insight into the coordination mode of HL^{SZ}, the vibrational spectra of NaL^{SZ} and **1** were compared. The IR spectrum of NaL^{SZ} shows ν^{as}(NH₂) at 3416 cm^{–1}, while ν^{as}(NH₂) is overlapped. The vibrations at 1630, 1292 and 1131 cm^{–1} may be assigned to ν(C=N)/(C=C), ν^{as}(SO₂) and ν^s(SO₂), respectively.¹⁹ For **1**, the observation of ν^{as}(NH₂) and ν^s(SO₂) at lower wavenumbers (3382 and 1259 cm^{–1}) than those of NaL^{SZ} may be attributed to presence of intermolecular H-bond of the type NH₂⋯OSO₂ (this is clearly shown in the crystal packing of **2**,



Fig. S5†). The shift of $\nu(\text{C}=\text{N})/(\text{C}=\text{C})$ of NaL^{SZ} from 1630 to 1624 cm^{-1} in **1** may be taken as a sign of participation of pyrimidine nitrogen in complex formation. Observation of $\nu(\text{S}-\text{N})$ mode at a higher wavenumber (972 cm^{-1}) with respect to NaL^{SZ} (977 cm^{-1}) (ref. 20) reveals presence of some sort of interaction between $\text{Ru}(\text{II})$ and sulfonamide nitrogen. While the IR changes between NaL^{SZ} and **1** are not so much as a result of change the sodium ion with $\text{Ru}(\text{II})$, the obtained IR data suggest coordination of L^{SZ} to $\text{Ru}(\text{II})$ ion *via* N_{pym} and N_{sfd} .

To conclude, the coordination sphere around $\text{Ru}(\text{II})$ in **1** is formed from *p*-cymene ring and two *N,N*-bidentate L^{SZ} molecules. As the seven-coordinates geometry is uncommon for $\text{Ru}(\text{II})$ complexes, the observed ^1H NMR equivalence clearly pointed to presence of two L^{SZ} coordinated ligands in a fluxional process in which the third bond of the base of the stool is oscillating between the two equivalent pyrimidine nitrogen atoms (Scheme 1).^{10a}

Treatment of the acetone solution of $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ ²¹ with two equivalents of silver trifluoromethane sulfonate and the subsequent removal of silver chloride yielded $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}(\text{acetone})]_2(\text{CF}_3\text{SO}_3)$. Reaction of the latter $\text{Rh}(\text{III})$ precursor with two equivalents of NaL^{SZ} was undertaken in a mixture of 1 : 1 (v/v) ($\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$) at 55 °C for 4 h. Orange precipitate of the type $[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2]$ (**2**) was obtained (Scheme 1). Analytical and spectroscopic tools (Fig. S3†) were used to get an idea about the structure of **2**. However, a clear-cut proof of the structure of **2** was gained from X-ray crystallographic analysis (Fig. 1). A singlet signal assigned to C_5Me_5 , with integration of fifteen equivalent protons, is shown in the ^1H NMR spectrum (Fig. S3†) of **2** at $\delta = 1.69$ ppm. The ^{13}C NMR spectrum (Fig. S3†) of **2** shows a doublet signal at $\delta = 94.8$ ppm ($J_{\text{C-Rh}} = 7.5$ Hz) due to coupling of $\eta^5\text{-C}_5\text{Me}_5$ carbon atoms with $\text{Rh}(\text{III})$ metal centre having $I = \frac{1}{2}$. Two characteristic fragments are observed in the ESI-MS(+) spectrum at $m/z = 759.1007$ $\{2 + \text{Na}\}^+$, 487.0669 $\{2 - \text{L}^{\text{SZ}}\}^+$. Like **1**, the ^1H NMR equivalence

indicated a fluxional behaviour between the two L^{SZ} molecules towards $\text{Rh}(\text{III})$ ion. In comparison with NaL^{SZ} , the pyrimidine-H3/H5 signals move downfield ($\delta = 8.47$ ppm) in **2** due to coordination of L^{SZ} with $\text{Rh}(\text{III})$. No change in the chemical shifts of the phenyl protons is detected, which rules out the role of either the primary amino group or SO_2 group in the formation of **2**.

Crystal structure

Diffusion of *n*-hexane into the dichloromethane solution of **2** afforded crystals suitable for single-crystal X-ray analysis. This compound crystallizes in the monoclinic space group $P2_1/m$. Applicable crystallographic parameters as well as the selected bond lengths (Å) and angles (°) are given in Fig. 1. The $\text{Rh}(\text{III})$ center is in a “three-legged piano-stool” coordination environment. While the seat of the stool is occupied by pentamethyl cyclopentadienyl ligand, the three legs are occupied with *N,N*-bidentate L^{SZ} ligand [$\text{Rh-N1}_5 = 2.137(2)$ Å, $\text{Rh-N1}_6 = 2.172(2)$ Å] and pyrimidine N [$\text{Rh-N1}_2 = 2.117(2)$ Å] from a second L^{SZ} molecule. The Rh-C bond lengths are in the range of 2.132(2)–2.167(2) Å. The N1_3 is close to Rh by 3.120 Å and thus it cannot be considered as a true bond. The N1_3 is involved in two short contacts with the methyl groups with distances of 2.678 and 2.691 Å. The $-\text{SO}_2$ group of a monodentate L^{SZ} ligand is participated in two H-bonds of different strengths with $-\text{NH}_2$ of the mono- and bidentate ligand as illustrated by the values of $\text{O}\cdots\text{N} = 2.887$ and 3.027 Å, respectively.

DFT and TD DFT calculations

Starting from the crystallographic data of **2**, ground-state geometry optimization was carried out, without any symmetry restriction, by B3LYP/Genecp method (SDD for Rh and 6-31G(d) for the rest of the elements). The local minimum structure was checked as minimum on the potential energy surface by calculating the vibrational modes. No imaginary vibrations were found. Selected optimized bond lengths and angles of **2** are given in Table S1† in comparison to the crystal data. The root mean square error of the bond lengths is 0.037 Å. While the Rh-N bonds match well with the theoretical data, the Rh-C bonds deviate largely from the calculated values by about ~ 0.05 Å. The optimized structure of **1** was obtained at B3LYP/Genecp (LANL2DZ for Ru and 6-31G(d) for the rest of the elements)

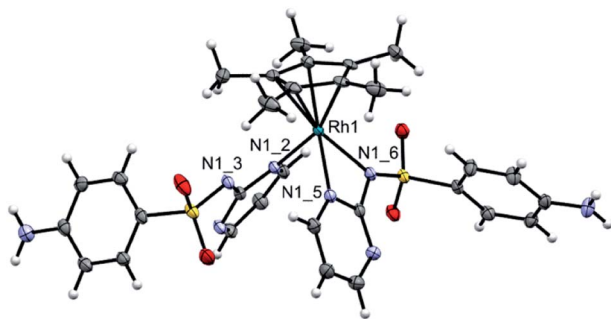


Fig. 1 Molecular structure of **2** (thermal ellipsoids are shown at 50% probability level). $\text{C}_{30}\text{H}_{33}\text{N}_6\text{O}_4\text{RhS}_2$, $M_r = 736.67$, yellow block, $0.319 \times 0.303 \times 0.120$ mm³, monoclinic space group $P2_1/n$, $a = 8.4430(19)$ Å, $b = 20.572(5)$ Å, $c = 17.935(5)$ Å, $\alpha = 90^\circ$, $\beta = 98.44(2)^\circ$, $\gamma = 90^\circ$, $V = 3081.4(13)$ Å³, $Z = 4$, $\rho_{\text{calcd}} = 1.588$ g cm⁻³, $\mu = 1.588$ mm⁻¹, $F(000) = 1512$, $T = 100(2)$ K, $R_1 = 0.0392$, $wR_2 = 0.0310$, 7062 independent reflections [$2\theta \leq 27.4879^\circ$] and 411 parameters. Selected bonds (Å) and angles (°) Rh-N1_2 2.117(2), Rh-N1_5 2.137(2), Rh-N1_6 2.172(2), Rh-C 2.132(2)–2.167(2); $\text{N1}_2\text{-Rh-N1}_5$ 82.93(7), $\text{N1}_2\text{-Rh-N1}_6$ 83.75(7), $\text{N1}_5\text{-Rh-N1}_6$ 61.12(7).

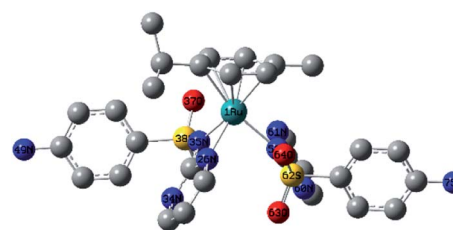


Fig. 2 Local minimum structure of **1** calculated at B3LYP/Genecp level of theory (Genecp: LANL2DZ for Ru and 6-31G(d) for the rest of the elements).



level of theory. Compound **1** adapts three-legged piano-stool geometry (Fig. 2). Cymene occupies the seat of the stool [Ru–C = 2.178–2.219 Å], while the two L^{SZ} molecules coordinate to Ru through three arms; bidentate [Ru–N26 = 2.147 Å and Ru–N35 = 2.145 Å] and monodentate [Ru–N52 = 2.168 Å] modes of L^{SZ} are observed. In contrast to the optimized and crystal structure of **2**, the monodentate ligand L^{SZ} coordinates to Ru(II) *via* sulfonamide N rather than pyrimidine N. The Ru–N bonds of the sulfonamide nitrogen are unequal; the bond to the bidentate L^{SZ} is 0.23 Å longer than that bonded to monodentate L^{SZ} . This indicates that there is not much space to accommodate two L^{SZ} in a bidentate mode of coordination.

Time dependent density functional calculations were carried out using hybrid exchange–correlation functional CAM-B3LYP, with a long-range correction term²² and LANL2DZ basis set to get an insight into the electronic structures of **1** and **2**. The first 30 singlet excited states were taken into consideration. Table S2† including the wavelengths and energies of the calculated transitions and their assignments is given in the ESI.† In 190–360 nm region, the calculated electronic spectrum of **1** (Fig. S6†) shows two bands at 232 and 352 nm (broad) as well as a shoulder at 291 nm corresponding to HOMO–3 → LUMO, HOMO–7/HOMO–4 → LUMO+1 and HOMO → LUMO/LUMO+2, respectively. Besides, two transitions assigned to HOMO → LUMO+2 and HOMO–4 → LUMO+2 (broad band) are observed at 485 and 456 nm. The reversed order of the transitions is owing to some states experience smaller solvent effect than the others. HOMO is mainly composed of d(Ru) character (Fig. 3) and thus the transitions at 485 and 291 nm have partially a nature of MLCT. The other bands may be accounted for the intra ligand transitions.

The TD DFT spectrum of **2** displays four bands at 231, 284, 383 and 429 nm with oscillator strengths of 0.1196, 0.1290, 0.0159 and 0.0145 as a result of HOMO–3 → LUMO+3, HOMO/HOMO–2 → LUMO+3, HOMO–3 → LUMO+1 and HOMO–4

→ LUMO in that order. HOMO–4 to HOMO–2 show d(Rh)/ π ligand (L^{SZ}) system character (Fig. 4). While LUMO and LUMO+1 are a mixture of d(Rh) and $\pi(Cp^*)$ orbitals, LUMO+3 has $\pi(L^{SZ})$ character. HOMO–LUMO gap of **1** is 0.59 eV wider than **2**. The transitions at 429 and 383 nm have ground-state composed of d(Rh)/ $\pi(L^{SZ})/\pi(Cp^*)$ and excited state of d(Rh)/ $\pi(Cp^*)$ forming d–d/MLCT.

Antimicrobial and cell viability

The Ru(II) and Rh(III) precursors; [RuCl(μ -Cl)(η^6 -*p*-Cym)]₂, [(η^5 -C₅Me₅)RhCl]₂(μ -Cl)₂] and their compounds (**1** and **2**) were screened for their potential antibacterial activity against *Staphylococcus aureus* ATCC 43300 (Gram-positive bacterium), *Escherichia coli* ATCC 25922, *Klebsiella pneumoniae* ATCC 700603, *Acinetobacter baumannii* ATCC 19606, and *Pseudomonas aeruginosa* ATCC 27853 (Gram-negative bacteria) using the standard broth microdilution assays.²³ The minimum inhibitory concentration (MIC, μ g mL^{−1}) values are given in Table 1. While most of ruthenium based complexes displayed significant biological activity,²⁴ [RuCl(μ -Cl)(η^6 -*p*-Cym)]₂ and its complex **1** exhibited no inhibitory activity against the tested bacteria at a concentration of 32 μ g mL^{−1}. Coordination of sulfadiazine to [RuCl(μ -Cl)(η^6 -*p*-Cym)]₂ gave rise to biologically inactive compound (**1**). The inactivity may be accounted for formation of covalent bond between Ru(II) and sulfonamide nitrogen, which should be free for sulfa drugs to be active.¹⁴

While [(η^5 -C₅Me₅)RhCl]₂(μ -Cl)₂] was inactive, compound **2** was potent against only the Gram-negative bacteria. *Acinetobacter baumannii* is two times more resistant to **2** (MIC 44 nM, equivalent to 32 μ g mL^{−1}) than the other tested Gram-negative bacteria. The inhibitory activity of **2** is associated to the cell wall structure of the bacteria, which is vital for the surviving of the bacteria. Inhibition of the synthesis of peptidoglycan may be

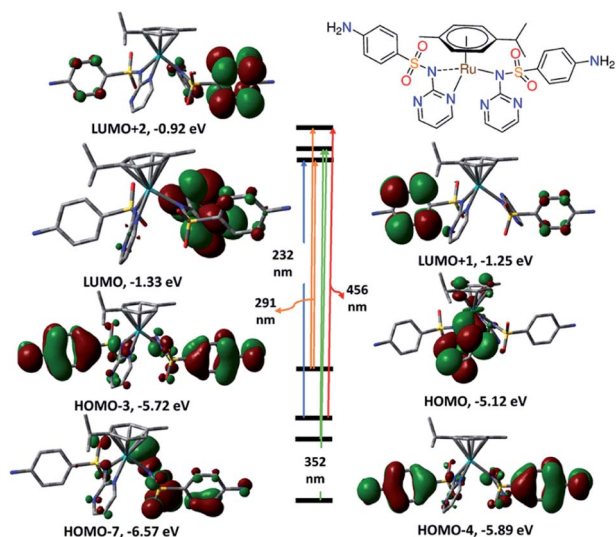


Fig. 3 Selected frontiers molecular orbitals and electronic transitions of **1** calculated at CAM-B3LYP/LANL2DZ level of theory.

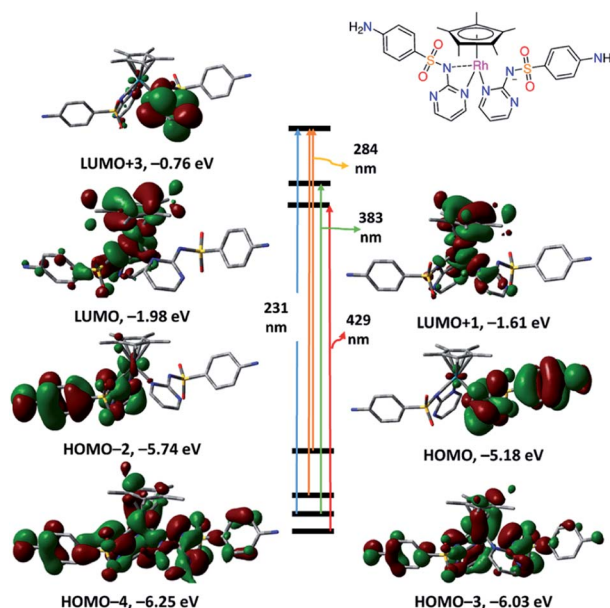


Fig. 4 Selected frontiers molecular orbitals and electronic transitions of **2** calculated at CAM-B3LYP/LANL2DZ level of theory.



Table 1 Minimum inhibitory concentration ($\mu\text{g mL}^{-1}$) determined for the synthesized complexes^a

	<i>E. coli</i>	<i>K. pneumoniae</i>	<i>A. baumannii</i>	<i>P. aeruginosa</i>
Ru ^b	>32	>32	>32	>32
Rh ^c	>32	>32	>32	>32
1	>32	>32	>32	>32
2	16	16	32	16

	<i>S. aureus</i>	<i>C. albicans</i>	<i>C. neoformans</i>
Ru ^b	>32	32	>32
Rh ^c	>32	4	2
1	>32	>32	>32
2	>32	4	4

^a Full names: *Staphylococcus aureus* ATCC 43300, *Escherichia coli* ATCC 25922, *Klebsiella pneumoniae* ATCC 700603, *Acinetobacter baumannii* ATCC 19606, *Pseudomonas aeruginosa* ATCC, *Candida albicans* ATCC 90028 and *Cryptococcus neoformans* var. *grubii* H99; ATCC 208821.

^b $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$. ^c $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$.

the job of the antibiotics. Gram-positive bacteria possess a thick cell wall comprising of several layers of peptidoglycan and teichoic acids that would be easily destroyed by drugs. Gram-negative bacteria have a comparatively thin cell wall containing a few layers of peptidoglycan surrounded by lipopolysaccharides and lipoproteins (second lipid membrane). The differences in the structure of the cell wall can afford alterations in the toxicity against the two types of the bacteria.

Based on the results of the antibacterial activity of **1** and **2**, we concluded that the antimicrobial activity cannot simply related to type and oxidation state of the metal ions, as well as type of arene moiety, other factors including lipophilicity, stability of the compounds, size of the receptor sites and electronic effect should be considered.

The antifungal activity of the synthesized complexes and their precursor metal salts was assessed against *Candida albicans* ATCC 90028 and *Cryptococcus neoformans* var. *grubii* H99; ATCC 208821. For fungi, the MIC values were determined as the lowest concentration at which the growth was completely reserved, well-defined by an inhibition of 80% and 70% for *C. albicans* and *C. neoformans* var. *grubii* H99, respectively.

While $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$ displayed antifungal activity against *Candida albicans*, its sulfadiazine complex is inactive. The antifungal activity of $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$, against *Candida albicans* (MIC 5.4 nM, equivalent to $4 \mu\text{g mL}^{-1}$), does not change by reaction with L^{SZ} to form **2**. For *Cryptococcus neoformans*, the antifungal activity of $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ is two times higher than **2** (MIC 5.4 nM, equivalent to $4 \mu\text{g mL}^{-1}$). Generally, complex **2** has the same antifungal activity against the tested fungal species.

It is believable that many organic and inorganic compounds simply have general toxicity that is not related to microbes, but also affects human cells. To discourse this issue, toxicity of the title organometallic complexes was evaluated by measuring cell viability against noncancerous human embryonic kidney cells (HEK293) and blood compatibility with the cell components.

Blood compatibility was assessed by calculating the HC^{10} (the concentration at which 10% haemolysis occurs; was calculated by curve fitting the inhibition values *versus* $\log(\text{concentration})$). Compound **2** was slightly potent to non-malignant HEK293 ($\text{CC}_{50} = 30.48 \mu\text{g mL}^{-1}$) and moderate haemoglobin release ($\text{HC}_{10} = 15.46 \mu\text{g mL}^{-1}$). Interestingly, the well-known Rh(III) precursor $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ was safe to HEK293 and induced negligible haemoglobin release ($\text{HC}_{10} > 32 \mu\text{g mL}^{-1}$). Based on the biological assays, greatest potential for further biological studies of $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ as antifungal agent is highly recommended.

Interaction with hen white egg lysozyme

Following the administration of the drugs, there is a highly chance to bind covalently or non-covalently^{25,26} to the surface-accessible histidyl of proteins. Such interactions may affect the toxicity, biodistribution and pharmacokinetics. While it is established that the target of Pd(II) and Pt(II) based drugs is DNA, recent studies present conversant and reasonable evidences that the binding of Au(III) compounds to protein is the origin for their cytotoxicity.²⁷ On the other hand, some model proteins could be used to carry the biologically active compounds into the living cells.²⁸ Besides, studying the stability of the biologically active molecules in presence of coordinating protein is essential. Therefore, HEWL was chosen for the propose of studying the possibility of bioconjugation and to investigate the stability of our compounds in presence of protein. The interactions between $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$ and some model proteins were studied with the combination of X-ray crystallography, spectroscopic and spectrometric measurements.^{29–31} Refinement of $[\text{RuCl}_2(\eta^6\text{-}p\text{-Cym})(\text{HEWL})]$ by 1.6 Å X-ray disclosed that the metalation process occurred at the His15 side-chain.²⁹ Destabilization of the adducts was observed upon mixing lysozyme with $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$.³¹ The structure of the apo-ferritin nanocage adduct with $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$ showed presence of three Ru sites per ferritin monomer.³⁰ Here, the HEWL binding affinity of **1** and **2** was investigated by positive mode electrospray ionization mass spectrometry. The reactivity of the complexes towards HEWL was investigated at the room temperature using 10 : 1 (complex : HEWL). The mass spectrometry spectra were immediately recorded after addition of the organometallic compound to the protein. Soaking lysozyme with ten equivalents of **1** produces mainly three Ru^{n+} adduct peaks at m/z 1601.0884 {HEWL + Ru^{n+} }, 1611.6432 {HEWL + $(2 \times \text{Ru})^{n+}$ } and 1622.5298 {HEWL + $(3 \times \text{Ru})^{n+}$ } as well as a weak noncovalent adduct peak^{25,26} at m/z 1671.8767 corresponding to bind of one molecule of **1** to lysozyme (Fig. 5). Destabilization of the HEWL adduct of **1** and presence of mainly three Ru binding sites per HEWL match the previously reported data of $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$.^{29–31}

Particularly, the ESI-MS of adduct **2** (Fig. 5), shows several adduct peaks at m/z 1605.5278 {HEWL + Rh^{n+} }, 1616.6437 {HEWL + $(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}^{n+}$ }, 1616.6437 {HEWL + $(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{N-pyrimidine})^{n+}$ }, 1644.4333 {HEWL + $(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})^{n+}$ } and 1655.3140 {HEWL + $\text{Rh}/(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})^{n+}$ }. The obtained results indicated that metalation of HEWL by Rh(III) fragments



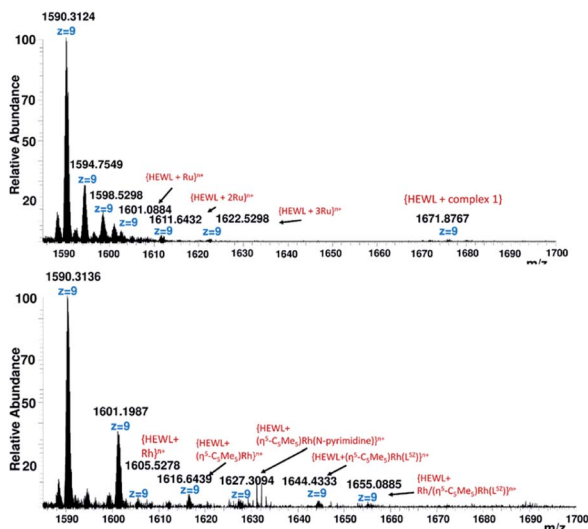


Fig. 5 Deconvoluted ESI-MS spectra of lysozyme treated with complexes 1 (up) and 2 (down).

occurs *via* the elimination of one or two sulfadiazine molecules. The fragment at m/z 1655.3140 showed presence of two metalation sites per Rh(III) HEWL adduct. In the absence of X-ray crystallography data of Ru(II) and Rh(III) adducts with lysozyme, it was difficult to assign the side-chain(s) binding sites.

Conclusion

Sulfa drugs, a well-known class of antibiotics, were extensively used to elaborate metal complexes of intriguing biological applications. However, they have not been used for synthesis of organometallic arene compounds. In the present contribution, first two examples of pentamethylcyclopentadienyl Rh(III) and *p*-cymene Ru(II) complexes bearing sulfadiazine drug (HL^{SZ}) were reported. The influence of the organometallic formation on the antibacterial activity of sulfadiazine was investigated. Single crystal X-ray analysis and spectroscopic studies indicated formation of $[(\eta^6\text{-}p\text{-Cym})\text{Ru}(\text{L}^{\text{SZ}})_2]$ and $[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2]$. The NMR equivalence clearly pointed to involvement of two L^{SZ} molecules in a fluxional process in which the third bond of the base of the “piano stool” is oscillating between two equivalent sulfonamide nitrogen atoms. The electronic structures of the complexes were investigated with the aid of quantum chemical calculations. While the coordination of sulfadiazine drug to cymene Ru(II) gave rise to antimicrobial inactive compound, Rh(III) organometallic compound showed interesting toxicity against Gram-positive bacteria as well as two fungal species; *Candida albicans* and *Cryptococcus neoformans* var. *grubii*. However, the Rh(III) compound showed slightly toxicity to non-malignant HEK293 ($\text{CC}_{50} = 30.48 \mu\text{g mL}^{-1}$) and moderate haemoglobin release ($\text{HC}_{10} = 15.46 \mu\text{g mL}^{-1}$). The precursor, $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$, was strongly potent against *Candida albicans* and *Cryptococcus neoformans* var. *grubii*, was safe to normal cells and was compatible with blood components. The interactions between the biologically active compound and

protein(s) may affect the toxicity, resistance, biodistribution as well as pharmacokinetics. Recent studies present conversant evidences that binding of some metal complexes to protein is the origin of their cytotoxicity. While $[(\eta^6\text{-}p\text{-Cym})\text{Ru}(\text{L}^{\text{SZ}})_2]$ decomposed during the reaction with hen white egg lysozyme (HEWL), a model protein, giving several adducts peaks with ruthenium ion(s), Rh(III) analogue was covalently bio-conjugated to HEWL *via* the elimination of one sulfadiazine drug. Based on the biological assays, greatest potential for further biological studies of $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ as anti-fungal agent is highly recommended.

Experimental section

Materials and instruments

The solvents were degassed and purified according to the standard published methods. Sodium(4-aminophenyl)sulfonyl-pyrimidin-2-ylazanide (NaL^{SZ}) was obtained from Sigma. $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$ (ref. 18) and $[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2$ ²¹ were synthesized by following the published procedures. Elemental compositions of the organometallic compounds were determined with Elementar Vario MICRO cube CHN analyzer nor an EA 3000 elemental analyzer from HEKtech. Electrospray mass spectra were run with a Thermo-Fisher Exactive Plus instrument with an Orbitrap mass analyzer at a resolution of $R = 70,000$ and a solvent flow rate of $5 \mu\text{L min}^{-1}$. IR spectra were recorded in the solid state on a Nicolet 380 FT-IR spectrometer equipped with a smart iFTR accessory. Electronic spectra were recorded on an Agilent 8453 diode array spectrophotometer. NMR spectra were recorded with Bruker-Avance 500 (^1H : 500.13 MHz and $^{13}\text{C}\{^1\text{H}\}$: 125.77 MHz) and Bruker-Avance 400 (^1H : 400.40 MHz, $^{13}\text{C}\{^1\text{H}\}$: 100.70 MHz) spectrometers. Assignments of NMR chemical shifts were done with the aid of $\{^1\text{H}, ^1\text{H}\}$ COSY and $\{^1\text{H}, ^{13}\text{C}\}$ HSQC.

Synthesis

$[(\eta^6\text{-}p\text{-Cym})\text{Ru}(\text{L}^{\text{SZ}})_2]$ (1). $[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$ (0.5 mmol; 315 mg) and NaL^{SZ} (2 mmol; 544 mg) were dissolved in methanol, whereupon an immediately yellow precipitate is formed. Stirring was continued for 3 h. The resulting precipitate was filtered off, washed with methanol, diethyl ether and dried under vacuum for few days. Yield (based on metal precursor): 84% (641 mg, 0.87 mmol). IR (ATR, diamond): $\tilde{\nu} = 3382$ (m, NH₂), 3335 (m, NH₂), 3231 (s), 1624 (w, CC/CN), 1592 (s), 1448 (vs), 1259 (s), 1131 (s), 1081 (s), 828 (m), 787 (s). ^1H NMR ($[\text{D}_6]$ DMSO, 500.13 MHz): $\delta = 8.64$ (br, 2H, pyrimidine-H3 or H5), 8.21 (br, 2H, pyrimidine-H3 or H5), 7.57 (d, $^3J_{\text{H,H}} = 6.1$ Hz, 4H, phenyl-H2/6), 6.68 (br, 2H, pyrimidine-H4), 6.52 (d, $^3J_{\text{H,H}} = 8.5$ Hz, 4H, phenyl-H3/5), 6.03 (d, $^3J_{\text{H,H}} = 6.0$ Hz, 1H, *p*-Cym-H2 or H6), 5.89 (d, $^3J_{\text{H,H}} = 6.6$ Hz, 1H, *p*-Cym-H6 or H2), 5.59 (d, $^3J_{\text{H,H}} = 4.1$ Hz, 1H, *p*-Cym-H3 or H5), 5.67 (br, 4H, NH₂), 5.53 (d, $^3J_{\text{H,H}} = 6.9$ Hz, 1H, *p*-Cym-H5 or H3), 2.63 (sep, $^3J_{\text{H,H}} = 6.9$ Hz, 1H, CH(CH₃)₂), 1.96 (s, 3H, CH₃), 1.01 (d, $^3J_{\text{H,H}} = 7.3$ Hz, 3H, CH(CH₃)₂), 0.94 (d, $^3J_{\text{H,H}} = 7.2$ Hz, 3H, CH(CH₃)₂) ppm. ^{13}C NMR ($[\text{D}_6]$ DMSO, 125.75 MHz): $\delta = 162.5$ (pyrimidine-C1), 159.1 (pyrimidine-C3/C5), 151.5 (phenyl-C4), 129.50 (phenyl-C1),



128.7 (phenyl-C2/6), 112.3 (pyrimidine-C4), 112.0 (phenyl-C3/5), 102.6 (*p*-Cym-C4), 100.0 (*p*-Cym-C1), 83.3 (*p*-Cym-C2 or C6), 83.2 (*p*-Cym-C6 or C2), 82.4 (*p*-Cym-C3 or C5), 80.3 (*p*-Cym-C5 or C3), 30.63 ($\text{CH}(\text{CH}_3)_2$), 22.07 ($\text{CH}(\text{CH}_3)_2$), 21.7 ($\text{CH}(\text{CH}_3)_2$), 17.8 (CH_3) ppm. ESI-MS (positive mode, acetone): $m/z = 735.1102$ $\{[(\eta^6\text{-}p\text{-Cym})\text{Ru}(\text{L}^{\text{SZ}})_2] + \text{H}^+\}^+$, 485.0577 $\{[(\eta^6\text{-}p\text{-Cym})\text{Ru}(\text{L}^{\text{SZ}})_2] + \text{H}^+ - \text{L}^{\text{SZ}}\}^+$. $\text{C}_{30}\text{H}_{32}\text{N}_8\text{O}_4\text{RuS}_2$: C 49.10, H 4.40, N 15.27, S 8.74, found, C 48.79, H 4.45, N 15.02, S 8.97.

$\{[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2] \text{ (2). Two equivalents of } \text{AgSO}_3\text{CF}_3$ (102.8 mg; 0.4 mmol), dissolved in few drops of water, was added to $\{[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2\}^{20}$ (123.6 mg, 0.2 mmol) in 15 mL acetone and stirred for 30 minutes. Silver chloride was filtered off. Removal of the solvent under vacuum yielded $\{[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}(\text{acetone})]_2\}(\text{CF}_3\text{SO}_3)$. The latter Rh(III) compound and L^{SZ} (0.8 mmol; 435 mg) were dissolved in 20 mL 1 : 1 methanol/ CH_2Cl_2 mixture and then the reaction mixture was stirred at 55 °C for 4 h. Filtration was done to remove water-soluble white precipitate (excess L^{SZ} and sodium chloride). The volume of the solution was reduced to 7 mL and diethyl ether was added. The solution was left overnight. Yellow product was filtered off, washed with diethyl ether and dried under vacuum. Yield (based on metal precursor): 44% (129 mg, 0.18 mmol). IR (ATR, diamond): $\tilde{\nu} = 3425$ (w, NH_2), 3389 (w, NH_2), 3355 (m, NH_2), 3242 (m, NH_2), 1647 (m, CC/CN), 1592, 1459, 1372, 1281, 1128, 1087, 1013, 985, 824, 786. ^1H NMR ($[\text{D}_6]$ DMSO, 500.13 MHz): $\delta = 8.47$ (br, 4H, pyrimidine-H3/H5), 7.50 (d, $^3J_{\text{H,H}} = 8.7$ Hz, 4H, phenyl-H2/6), 6.77 (t, $^3J_{\text{H,H}} = 4.9$ Hz, 2H, pyrimidine-H4), 6.48 (d, $^3J_{\text{H,H}} = 9.1$ Hz, 4H, phenyl-H3/5), 5.62 (s, 4H, NH_2), 1.69 (s, 15H, Cp*- CH_3) ppm. ^{13}C NMR ($[\text{D}_6]$ DMSO, 125.75 MHz): $\delta = 162.3$ (pyrimidine-C1), 159.1 (pyrimidine-C3/C5), 151.2 (phenyl-C4), 130.1 (phenyl-C1), 129.0 (phenyl-C2/6), 111.9 (pyrimidine-C4), 111.0 (phenyl-C3/5), 94.8 (d, $^1J_{\text{C-Rh}} = 7.5$ Hz, Cp*- CCH_3), 9.0 (Cp*- CCH_3). ESI-MS (positive mode, methanol): $m/z = 1495.2134$ $\{2 \times [(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2] + \text{Na}^+\}^+$, 759.1007 $\{[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2] + \text{Na}^+\}^+$, 487.0669 $\{[(\eta^5\text{-C}_5\text{Me}_5)\text{Rh}(\text{L}^{\text{SZ}})_2] - \text{L}^{\text{SZ}}\}^+$. $\text{C}_{30}\text{H}_{33}\text{N}_8\text{O}_4\text{RhS}_2 \cdot 0.5\text{CH}_2\text{Cl}_2$: C 47.02, H 4.55, N 14.38, S 8.23, found C 47.05, H 4.47, N 14.69, S 8.47.

X-ray diffraction analysis

Single crystals appropriate for X-ray crystallography analysis of **2** were isolated by slow diffusion of *n*-hexane into CH_2Cl_2 of the complex. Selected crystals of **2** were immersed in a film of perfluoropolyether oil, mounted on MiTeGen sample holders and transferred to a stream of cold nitrogen of the diffractometer. The crystal data were collected on a Bruker D8 QUEST diffractometer with a CMOS area detector and multi-layer mirror or graphite monochromated Mo- K_α radiation ($\lambda = 0.71073$ Å) at 100 K. All the diffracted intensities were collected using an APEX2-CCD detector and corrected for Lorentz-polarization and absorption using the Bruker AXS software. The structures were solved with the SHELXT program using intrinsic phasing method³² and refined with a full-matrix least-squares procedure using the SHELXL program.³³ All the non-hydrogen atoms were refined anisotropically refined. Hydrogen atoms were included in the structure factors calculations. Crystallographic data have been deposited with the

Cambridge Crystallographic Data Center as ESI publication no. CCDC 1940322 for compound **2**.

DFT calculations

Ground-state geometry optimization of **1** and **2** were carried at B3LYP/Genecp (LANL2DZ for Ru, SDD for Rh, and 6-31G(d) for the rest of the elements) level of theory. Time-dependent density functional theory calculations were performed by CAM-B3LYP/LANL2DZ method using PCM model to introduce the solvent effect. All the calculations were done using Gaussian 03 package³⁴ and the obtained data were visualized using Gaussview.³⁵

Interaction with hen white egg lysozyme

The reactivity of the synthesized organometallic compounds, dissolved in pure DMSO towards HEWL (2 mg mL^{-1} , H_2O) was studied by orbitrap high resolution mass spectrometer (ThermoFisher Exactive plus orbitrap) equipped with the conventional electrospray ionization source. The reaction mixtures were immediately measured after mixing. The working conditions were as follows: spray voltage 3.80 kV, capillary voltage 45 V, and capillary temperature 320 °C. For acquisition, Thermo Xcalibur qual was used.

Biological activity testing

$[\text{RuCl}(\mu\text{-Cl})(\eta^6\text{-}p\text{-Cym})]_2$, $\{[(\eta^5\text{-C}_5\text{Me}_5)\text{RhCl}]_2(\mu\text{-Cl})_2\}$ and compounds (**1** and **2**) were initially screened at a fixed concentration (usually 32 $\mu\text{g mL}^{-1}$ or 20 μM), with a maximum of 0.5% DMSO, final in assay concentration, against a series of bacterial and fungal pathogens; *Staphylococcus aureus* ATCC 43300, *Escherichia coli* ATCC 25922, *Klebsiella pneumoniae* ATCC 700603, *Acinetobacter baumannii* ATCC 19606, *Pseudomonas aeruginosa* ATCC 27853, *Candida albicans* ATCC 90028 and *Cryptococcus neoformans* var. *grubii* H99; ATCC 208821. For active compound, a follow-up hit confirmation is triggered, where the activity is measured by means of a dose-response assay against the same strains.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Antimicrobial screening was performed by CO-ADD (The Community for Antimicrobial Drug Discovery), funded by the Wellcome Trust (UK) and the University of Queensland (Australia).

References

- 1 J. H. Maren, *Annu. Rev. Pharmacol.*, 1976, **16**, 309.
- 2 *Burger's Medicinal Chemistry and Drug Discovery*, ed. M. E. Wolff, Wiley, Laguna Beach, 5th edn, 1996, vol. 2, pp. 528–576.



- 3 S. Roland, R. Ferone, R. J. Harvey, V. L. Styles and R. W. Morrison, *J. Biol. Chem.*, 1979, **254**, 10337.
- 4 A. Bult, *Met. Ions Biol. Syst.*, 1982, **16**, 261.
- 5 W. H. Wernsdorfer, *Acta Trop.*, 1994, **56**, 143.
- 6 (a) R. Yuan, R. Xiong, Z. Chen, P. Zhang, H. Ju, Z. Dai, Z. Guo, H. Fun and X. You, *J. Chem. Soc., Dalton Trans.*, 2001, 774; (b) N. C. Baenziger, S. L. Modak and C. L. Fox, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 1983, **39**, 1620; (c) C. J. Brown, D. S. Cook and L. Sengier, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 1985, **41**, 718.
- 7 (a) N. C. Baenziger and A. W. Struss, *Inorg. Chem.*, 1976, **15**, 1807; (b) D. S. Cook and M. F. Turner, *J. Chem. Soc., Perkin Trans. 2*, 1975, 1021.
- 8 M. K. Hassan, R. M. Hassan and M. A. Abd-Alla, *Monatsh. Chem.*, 1991, **122**, 829.
- 9 P. A. Ajibade, G. A. Kolawole, P. O'Brien, M. Helliwell and J. Raftery, *Inorg. Chim. Acta*, 2006, **359**, 3111.
- 10 (a) J. Tommasino, F. N. R. Renaud, D. Luneau and G. Pilet, *Polyhedron*, 2011, **30**, 1663; (b) K. K. Narang and J. K. Gupta, *Transition Met. Chem.*, 1977, **2**, 83; (c) A. Bult, J. D. Uitterdijk and H. B. Klasen, *Transition Met. Chem.*, 1979, **4**, 285; (d) E. Kremer, G. Facchin, E. Estévez, P. Alborés and E. J. Baran, *J. Inorg. Biochem.*, 2006, **100**, 1167.
- 11 (a) L. Menabue and M. Saladini, *J. Inorg. Biochem.*, 1993, **49**, 201; (b) K. K. Narang and J. K. Gupta, *Transition Met. Chem.*, 1977, **2**, 181.
- 12 A. Pasini and E. Bersanetti, *Inorg. Chim. Acta*, 1983, **80**, 99.
- 13 A. García-Raso, J. J. Fiol, G. Martorell, A. López-Zafra and M. Quirós, *Polyhedron*, 1997, **16**(4), 613.
- 14 W. O. Foye, T. L. Lemke, and D. A. Williams, *Principles of Medicinal Chemistry*, Williams & Williams, 4th edn, 1995, pp. 709–713.
- 15 (a) J. Li, M. Tian, Z. Tian, S. Zhang, C. Yan, C. Shao and Z. Liu, *Inorg. Chem.*, 2018, **57**, 1705; (b) W. Ma, L. Guo, Z. Tian, S. Zhang, X. He, J. Li, Y. Yang and Z. Liu, *Dalton Trans.*, 2019, **48**, 478.
- 16 J. M. Radmaker, D. van den Bongard, D. Pluim, J. H. Beijnen and J. H. Schellens, *Clin. Cancer Res.*, 2004, **10**, 3717; D. A. Wolters, M. Stefanopoulou, P. J. Dyson and M. Groessel, *Metallomics*, 2012, **4**, 1185.
- 17 G. Sava, S. Zorzet, C. Turrin, F. Vita, M. Soranzo, G. Zabucchi, M. Cocchietto, A. Bergamo, S. Di Giovine, G. Pezzoni, L. Sartor and S. Garbisa, *Clin. Cancer Res.*, 2003, **9**, 1898; L. Messori, P. Orioli, D. Vullo, E. Alessio and E. Iengo, *Eur. J. Biochem.*, 2000, **267**, 1206; F. Frausin, V. Scarcia, M. Cocchietto, A. Furlani, B. Serli, E. Alessio and G. Sava, *J. Pharmacol. Exp. Ther.*, 2005, **313**, 227.
- 18 S. B. Jensen, S. J. Rodger and M. D. Spicer, *J. Organomet. Chem.*, 1998, **556**, 151.
- 19 A. M. Mansour, F. A. Soliman, O. R. Shehab and N. T. Abdel-Ghani, *RSC Adv.*, 2017, **7**, 39989; O. R. Shehab, F. A. Soliman, N. T. Abdel-Ghani and A. M. Mansour, *Spectrochim. Acta, Part A*, 2020, **28**, 117821.
- 20 (a) G. M. Golzar-Hossain, A. J. Amoroso, A. Banu and K. M. A. Malik, *Polyhedron*, 2007, **26**, 967; (b) A. M. Mansour, *Inorg. Chim. Acta*, 2013, **394**, 436; (c) A. M. Mansour, *J. Coord. Chem.*, 2013, **66**(7), 1118.
- 21 C. White, A. Yates, P. M. Maitlis and D. M. Heinekey, *Inorg. Synth.*, 1992, **29**, 228.
- 22 T. Yanai, D. P. Tew and N. C. Handy, *Chem. Phys. Lett.*, 2004, **393**, 51.
- 23 M. A. Blaskovich, J. Zuegg, A. G. Elliott and M. A. Cooper, *ACS Infect. Dis.*, 2015, **1**, 285.
- 24 F. Li, F. J. G. Collins and R. Keene, *Chem. Soc. Rev.*, 2015, **44**, 2529.
- 25 (a) A. M. Mansour and O. R. Shehab, *Dalton Trans.*, 2018, **47**, 3459; (b) A. M. Mansour, O. R. Shehab and K. Radacki, *Eur. J. Inorg. Chem.*, 2020, 299; (c) A. M. Mansour and K. Radacki, *Polyhedron*, 2020, **175**, 114175.
- 26 G. Ferraro, A. M. Mansour and A. Merlino, *Dalton Trans.*, 2018, **47**, 10130.
- 27 L. Engman, M. McNaughton, M. Gajewska, S. Kumar, A. Birmingham and G. Powis, *Anticancer Drugs*, 2006, **17**, 539.
- 28 (a) M. Chaves-Ferreira, I. S. Albuquerque, D. Matak-Vinkovic, A. C. Coelho, S. M. Carvalho, L. M. Saraiva, C. C. Romão and G. J. L. Bernardes, *Angew. Chem., Int. Ed.*, 2015, **54**, 1172; (b) H. Tabe, K. Fujia, S. Abe, M. Tsujimoto, T. Kuchimaru, S. Kizaka-Kondoh, M. Takano, S. Kitahawa and T. Ueno, *Inorg. Chem.*, 2015, **54**, 215.
- 29 I. W. McNae, K. Fishburne, A. Habtemariam, T. M. Hunter, M. Melchart, F. Wang, M. D. Walkinshaw and P. J. Sadler, *Chem. Commun.*, 2004, 1786.
- 30 Y. Takezawa, P. Bockmann, N. Sugi, Z. Wang, S. Abe, T. Murakami, T. Hikage, G. Erker, Y. Watanabe, S. Kitagawa and T. Ueno, *Dalton Trans.*, 2011, **40**, 2190.
- 31 M. P. Sullivan, M. Groessel, S. M. Meier, R. L. Kingstone, D. C. Goldstone and C. G. Hartinger, *Chem. Commun.*, 2017, **53**, 4246.
- 32 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 3.
- 33 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112.
- 34 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, V. G. Zakrzewski, J. A. Montgomery, R. E. Stratmann, J. C. Burant, S. Dapprich, J. M. Millam, A. D. Daniels, K. N. Kudin, M. C. Strain, O. Farkas, J. Tomasi, V. Barone, M. Cossi, R. Cammi, B. Mennucci, C. Pomelli, C. Adamo, S. Clifford, J. Ochterski, G. A. Petersson, P. Y. Ayala, Q. Cui, K. Morokuma, D. K. Malick, A. D. Rabuck, K. Raghavachari, J. B. Foresman, J. Cioslowski, J. V. Ortiz, A. G. Baboul, B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi, R. Gomperts, R. L. Martin, D. J. Fox, T. Keith, M. A. Al-Laham, C. Y. Peng, A. Nanayakkara, C. Gonzalez, M. Challacombe, P. M. W. Gill, B. G. Johnson, W. Chen, M. W. Wong, J. L. Andres, M. Head-Gordon, E. S. Replogle and J. A. Pople, *GAUSSIAN 03 (Revision A.9)*, Gaussian, Inc., Pittsburgh, 2003.
- 35 A. Frisch, A. B. Nielson and A. J. Holder, *Gaussview User Manual*, Gaussian, Inc, Pittsburgh, PA, 2000.

