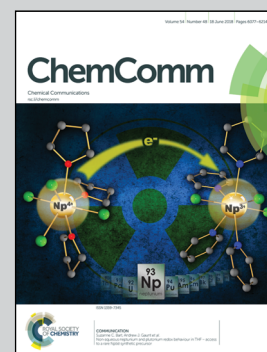


Showcasing research from Christian Hartinger's and David Goldstone's Laboratories at the Schools of Chemistry and Biological Sciences at the University of Auckland, New Zealand.

Unexpected arene ligand exchange results in the oxidation of an organoruthenium anticancer agent: the first X-ray structure of a protein–Ru(carbene) adduct

The reaction of an EPR silent Ru^{II}(N-heterocyclic carbene)-(η⁶-*p*-cymene) compound with the model protein lysozyme gave a protein adduct with the *p*-cymene ligand released. The latter has been suggested to stabilise the oxidation state of the Ru centre and upon release for the first time we could show by EPR spectroscopy that the metal centre is indeed oxidised to Ru^{III}.

As featured in:



See David C. Goldstone, Christian G. Hartinger *et al.*, *Chem. Commun.*, 2018, 54, 6120.

Cite this: *Chem. Commun.*, 2018, 54, 6120Received 27th March 2018,
Accepted 23rd April 2018

DOI: 10.1039/c8cc02433b

rsc.li/chemcomm

Unexpected arene ligand exchange results in the oxidation of an organoruthenium anticancer agent: the first X-ray structure of a protein–Ru(carbene) adduct†

 Matthew P. Sullivan,^{id ab} Michél K. Nieuwoudt,^{id ac} Graham A. Bowmaker,^a
 Nelson Y. S. Lam,^{id a} Dianna Truong,^{id a} David C. Goldstone^{id *b} and
 Christian G. Hartinger^{id *a}

The first X-ray structures of adducts formed between a Ru^{II}(N-heterocyclic carbene)(η⁶-*p*-cymene) compound and a protein are reported. Coordination to the protein induced the cleavage of the cymene ligand and EPR spectroscopy demonstrated the oxidation of the Ru centre.

Platinum-based anticancer drugs have been widely used in the treatment of tumourigenic diseases and Ru^{III} and other metal complexes are considered a promising addition to the toolbox of anticancer chemotherapeutics.^{1–4} Sodium [*trans*-tetrachlorido-bis(indazole)ruthenate(III)] is currently undergoing clinical trials, showing promising activity and good tolerability,^{2,5} while organoruthenium(II) piano stool compounds are on track toward clinical development.⁶ The latter scaffold consists of a Ru centre with a π-bound arene and either ancillary ligands and/or leaving groups to complete the coordination sphere around the metal centre. Altering the scaffold allows for fine-tuning of the pharmacological properties of the compound type.^{3,6,7} RAPTA-C [Ru^{II}(cym)(1,3,5-triaza-7-phosphatricyclo-[3.3.1.1]decane)Cl₂] (cym = η⁶-*p*-cymene) and RAED [Ru^{II}(arene)(1,2-diaminoethane)Cl]⁺ are the most notable lead structures to date, and have been shown to selectively target amino acids and DNA, respectively.^{8–11} More recently, we identified the plecstatin-targeting properties of orally-active pyridinecarbothioamide complexes.¹² Interestingly, most compounds that feature a monodentate non-leaving ligand exhibit only limited *in vitro* anticancer activity, unless the ligand itself is bioactive, while bidentate ligands as simple as 1,2-ethylenediamine often induce cytotoxicity.^{2–4,6–10,12–14} This is based on stability of

the bond(s) between the metal centre and the ancillary ligands and the lipophilicity of the ligand and complex.

N-Heterocyclic carbenes (NHCs) have been widely studied as ancillary ligands in homogeneous catalysis and form stable complexes with metal centres but they have only recently been adopted for medicinal chemistry applications.^{15,16} Early Au(NHC) complexes showed promising cytotoxic activity, with inhibition of thioredoxin reductase (TrxR) being a possible mechanism.¹⁷ An ESI-MS study on a selenopeptide derived from TrxR suggested that the inhibition is caused by binding of the Au compound to the selenocysteine residue of TrxR.¹⁸ Recently structural data for an Au(NHC)–protein adduct has been reported. Merlino *et al.* found that Au(1-butyl-3-methylimidazole-2-ylidene)Cl formed six adducts with the model protein thaumatin.¹⁹ Ott *et al.* studied Ru(NHC) complexes, such as [dichlorido(dmb)(cym)ruthenium(II)] **1** (dmb = 1,3-dimethylbenzimidazol-2-ylidene; Fig. 1), for medicinal applications and showed that, like Au(NHC) complexes, they inhibited TrxR at μM concentrations.^{20,21} In order to add structural data to the discussion on the mode of action of this compound type, we report here the first crystal structure of a Ru(NHC)–protein adduct between **1** and the model protein hen egg white lysozyme (HEWL).

HEWL has a molecular weight of 14.3 kDa (129 amino acids) and is an ideal model for studies demonstrating protein modification with metal ions.^{22–31} Crystals of HEWL are readily obtained by hanging-drop vapour diffusion using a reservoir solution containing NaCl and NaOAc at pH 4.7. The crystals were formed when the reservoir solution was mixed with an equal volume of an aqueous solution of HEWL (100 mg ml^{−1}).

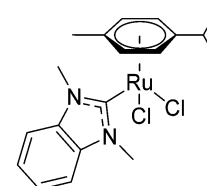


Fig. 1 Structure of [dichlorido(dmb)(cym)ruthenium(II)] **1**.

^a School of Chemical Sciences, University of Auckland, Private Bag 92019, Auckland 1142, New Zealand. E-mail: c.hartinger@auckland.ac.nz;

Web: <http://www.hartinger.auckland.ac.nz/>

^b School of Biological Sciences, University of Auckland, Private Bag 92019, Auckland 1142, New Zealand. E-mail: d.goldstone@auckland.ac.nz

^c The Photon Factory, University of Auckland, New Zealand

† Electronic supplementary information (ESI) available: Experimental procedures, literature analysis of HEWL–Ru adducts, additional mass spectrometry, X-ray crystallography, and EPR spectroscopy data. See DOI: 10.1039/c8cc02433b



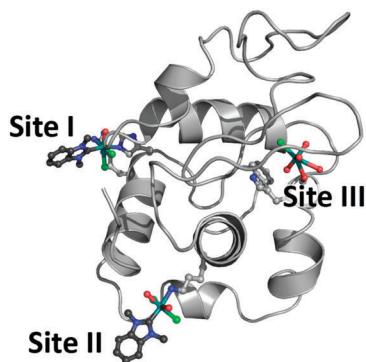


Fig. 2 Sites of metalation on HEWL after reaction with **1** (PDB ID 6BO2) indicating formation of a bidentate $\text{Ru}(\text{dmb})(\text{OH}_x)\text{Cl}_2$ -HEWL adduct at site I and a monodentate $\text{Ru}(\text{dmb})(\text{OH}_x)_2\text{Cl}_2$ -HEWL adduct at site II. Ru (teal), Cl (green), OH_x (red), and NHC (blue and dark grey) are shown in ball and stick representation while the rest of the protein is presented as a light grey cartoon.

Crystals formed within 1 d and were soaked in a solution of NaNO_3 and NaOAc containing **1** for a period of 3 d (PDB ID 6BO1) or 1 month (PDB ID 6BO2). The crystals were analysed at the Australian synchrotron MX1 beamline at a wavelength of 0.9537 Å. The phases were estimated by the molecular replacement method with a structure of HEWL (PDB ID 4NHI) as the search model. The Ru-containing fragments were identified in an anomalous difference map where they display significant anomalous signals compared with the proteinaceous light elements.³² Analysis of the data revealed that an initial adduct of the composition $\text{Ru}(\text{dmb})(\text{OH}_x)\text{Cl}_2$ -HEWL ($x = 1, 2$) was formed after a 3 d incubation with the protein acting as a bidentate ligand to the metal through the His15 and Arg14 residues (site I, Fig. 2). A second adduct was only present in a longer soak of one month, and formed at Lys33 where it was identified as $\text{Ru}(\text{dmb})(\text{OH}_x)_2\text{Cl}_2$ -HEWL with the protein monodentately coordinated to the Ru centre (site II, Fig. 2). Notably, in both structures the cym ligand is absent, while the NHC remains coordinated to the metal centre. Upon compound placement, the structures were refined to $R_{\text{work}}/R_{\text{free}}$ values of 0.20/0.23 and 0.18/0.23 for the structures of the adducts formed after 3 d and 1 month, respectively (ESI† Table S2).

A comparison of the ruthenated HEWL structures to native HEWL (PDB ID 4NHI) showed no major structural perturbations and only localised changes at sites I, II, and III in the side chain conformations to accommodate the organometallic fragment. This is supported by the small root-mean-square deviations (RMSD) of 0.164 and 0.289 Å (for overlays see Fig. S1, ESI†).

In the crystal of HEWL soaked for 3 d with **1**, a peak was observed in the anomalous difference map with a peak height of 12.2σ . This was coupled with a large area of $2F_o - F_c$ electron density (magenta) adjacent to the $\text{N}^{\epsilon 1}$ atom of His15 (Fig. S2a, ESI†). His15 is located on the surface of the protein near the C terminus. Histidine residues have been shown to be binding partners for Ru in previous protein structures.^{11,33} The anomalous density allowed for the placement of the Ru centre with a $\text{Ru}-\text{N}^{\epsilon 1}_{\text{His15}}$ bond length of 2.54 Å (site I). This is slightly longer than that

observed in the structure of HEWL modified with a $\text{Ru}(\text{cym})\text{Cl}_2$ moiety (PDB ID: 5V4G).³⁰ This interaction perturbs the side chain position of His15 resulting in a rotation of approximately 130° compared to apo-HEWL.

Five projections from the metal centre were observed in the $2F_o - F_c$ electron density map. In *trans* position to the coordinated His residue, a dagger-shaped region of density corresponding to dmb was observed. The distance between the Ru and carbene-C of 2.06 Å is in a similar range as found in $\text{Ru}(\text{NHC})$ complexes.³⁴ During refinement it became evident that Arg14 was coordinated to the metal centre ($\text{Ru}-\text{N}^{\eta 1}_{\text{Arg14}}$ 2.22 Å) and was displaced by 4.43 Å compared to a previously published structure (PDB ID: 5V4G, overlapped in Fig. S3, ESI†). In addition, density was evident for a further three spherical projections off the Ru centre. Two of these were modelled as chlorido ligands and the third as a coordinated water. The chlorido ligands are likely retained due to chloride present during crystal formation preventing ligand exchange reactions. The average $\text{Ru}-\text{Cl}$ distance was determined as 2.44 Å. The $\text{Ru}-\text{O}_{\text{water/hydroxido}}$ distance of 2.17 Å is consistent with a hydroxido/water ligand. This analysis demonstrates that the protein acts as a bidentate ligand, confirming an octahedral coordination geometry about the Ru centre. Moreover, no density was observed for the *p*-cymene ligand. This indicates that the *p*-cymene ligand was released during the binding event. The presence of the NHC ligand at the Ru centre may labilise the π -bound *p*-cymene, similar to the elongation effect NHC ligands have in complexes on bonds in *trans* position.³⁴ Consequently, the bound metallofragment at site I was identified as a $[\text{Ru}(\text{dmb})(\text{OH}_x)\text{Cl}_2]$ fragment coordinated to the $\text{N}^{\epsilon 1}_{\text{His15}}$ and $\text{N}^{\eta 1}_{\text{Arg14}}$ atoms. The refined $2F_o - F_c$ electron density map about these ligands validated the atoms in our model and the positioning of these ligands was a good fit, with an occupancy of 0.65 (Fig. S2b; for a discussion on the occupancies see Table S2, ESI†). The addition of the NHC ligand seems to weaken the $\text{Ru}-\text{cym}$ bond. This is in contrast to HEWL binding studies with other organoruthenium compounds, in which the cym ligand was retained in the protein structure.^{22,30} Notably, Ott *et al.* also showed that the dibenzyl derivative of **1** is significantly cytotoxic, demonstrating the need of the NHC ligand for the bioactivity to remain attached to the metal centre.²¹ As the arene ligand is known to stabilise the Ru centre in +2 oxidation state, we used EPR spectroscopy to investigate the impact of arene ligand exchange on the oxidation state of the Ru centre (Fig. 3). The 3 species obtained from simulated fitting of the HEWL-**1** spectrum (Fig. 3b and Table S1, ESI†) are consistent with the mononuclear Ru^{III} nature³⁵ of HEWL-**1** adducts, while neither HEWL, **1**, nor the buffer are paramagnetic. While the cleavage of cym has been seen before in the binding of RAPTA-C to macromolecules,^{36,37} this is the first instance that the ruthenation was confirmed to result in the oxidation of the Ru centre. The spectrum for HEWL-**1** KP1019 with 1.92×10^{18} spins, calculated using the area of the doubly integrated 1st-derivative EPR spectrum of a CuSO_4 standard, is much stronger than that of HEWL-**1** (5.94×10^{15} spins). This may be explained by only a small proportion of **1** undergoing ligand exchange with HEWL and oxidation to Ru^{III} , while KP1019 with its Ru^{III} centre is intrinsically paramagnetic.



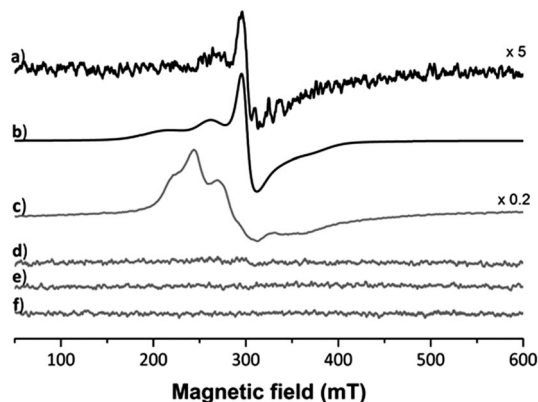


Fig. 3 EPR spectrum recorded for (a) HEWL-1 adducts, (b) simulated spectrum of the adducts EPR spectra for (c) HEWL and KP1019, (d) HEWL, (e) **1**, and (f) buffer. Simulation parameters for adducts (all rhombic 50 : 50 Gauss : Lorentzian line shapes, HEWL-1a: $g = [3.08, 2.24, 1.77]$, linewidths (mT) [45, 33, 38]; HEWL-1b: $g = [2.52, 2.38, 1.91]$, linewidths (mT) [22, 22, 23], HEWL-1c: $g = [2.18, 2.14, 2.00]$, linewidths (mT) [8, 18, 40]).

To study the impact of longer soaking times on the adduct formation and protein structure, HEWL was treated with **1** for 1 month. The crystals analysed featured the identical adduct and interaction at site I but at higher occupancy (0.90 vs. 0.65 after 3 d; Fig. 4a and b). This indicates progressively complete metalation of the protein at site I. The anomalous density (Fig. 4a) disappeared at 14.2σ , and showed similar distances (Ru-N^{ε1}_{His15} 2.13 Å, Ru-N^{η1}_{Arg14} 2.14 Å, Ru-C_{NHC} 2.05 Å, Ru-O 2.00 Å, Ru-Cl 2.44 Å) and maps as for the 3 d incubation.

In addition to the adduct at site I, the presence of anomalous density (yellow; Fig. 4c and d) was noted near Lys33 (site II) with a peak height of 12.3σ and was therefore identified as a Ru centre. The Ru centre was found with a large electron density observed in $2F_o - F_c$ maps (magenta) adjacent to the N^{ε1}_{Lys33} atom (Fig. 4c). The anomalous density allowed for the placement of the Ru centre with a Ru-N^{ε1}_{Lys33} bond length of 2.18 Å, which resulted in a shift in the side chain position of Lys33. Note that this binding site is different to site II identified for [Ru(cym)Cl₂]₂ on HEWL, which was found to form an adduct with Asp101 that acts as a bridging ligand for two Ru centres.³⁰

Around the Ru centre, five distinct projections were observed emanating from the metal centre (Fig. S5 for the coordination modes at binding sites I and II, ESI†). Four projections appeared to adopt a planar coordination arrangement including the metal centre and these were initially modelled as chlorido ligands. However, for two of the projections negative peaks appeared in the $F_o - F_c$ electron density maps, suggesting the presence of two OH_x ligands at the metal centre. Thus, these were modelled with an average Ru-Cl distance of 2.45 Å and a Ru-O_{water/OH} distance of 2.18 Å. The last area of residual electron density was located *trans* to the N^{ε1}_{Lys33} atom, as at site I, dagger-shaped and accordingly modelled as dmb with a Ru-C_{NHC} distance of 2.08 Å. The NHC forms a π -stacking interaction with the indole residue of Trp62, resulting in a twist of Trp62 by 21°, with the shortest distance between the aromatic systems being 3.44 Å. This interaction may have an additional stabilising effect on the binding of the

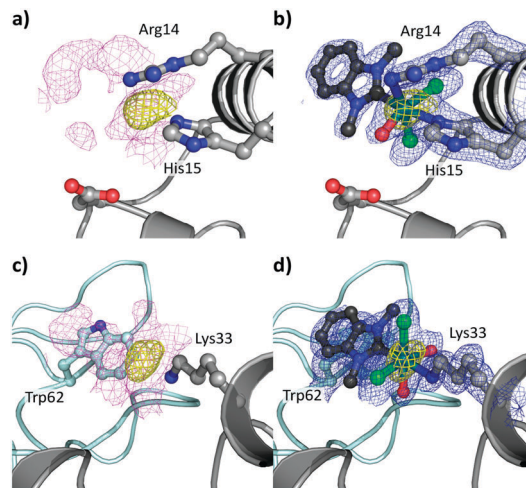


Fig. 4 Details of the binding sites for **1** on HEWL after 1 month incubation (PDB ID 6BO2), binding site I on HEWL (a and b) and binding site II (c and d). The [Ru(dmb)(OH_x)Cl₂] ($y = 1, 2$) fragments are indicated in ball and stick representation with the Ru (teal). The electron density maps are contoured at 1σ (magenta and blue maps) while the anomalous difference maps are contoured at 4σ (yellow maps). (a) Unbiased electron density (magenta) and anomalous difference map showing binding site I, as well as (b) placement of the [Ru(dmb)(OH_x)Cl₂] fragment into the refined electron density map (blue) with the anomalous difference map. The interaction between HEWL and **1** at binding site II and the adjacent protein is shown as a light teal cartoon in the background with Trp62 highlighted. (c) Unbiased electron density and the anomalous difference map. (d) Placement of the Ru(dmb)(OH_x)₂Cl₂ adduct into the refined electron density map with the anomalous difference map.

organoruthenium fragment to the protein and also impacts its positioning on the protein. Therefore, the bound metallo-fragment was identified as Ru(dmb)(OH_x)₂Cl₂ coordinated to N^{ε1}_{Lys33}. The refined $2F_o - F_c$ electron density map about these ligands validated the atoms in our model and the positioning of these ligands were a good fit, with an occupancy of 0.8 (Fig. 4d). This indicates that, as at site I, the *p*-cymene ligand underwent a ligand exchange reaction. As the interaction occurs with the N^{ε1} atom of Lys33 in a monodentate fashion, this further supports the hypothesis that the NHC ligand weakens the π -bond between the arene and the Ru centre.

A third Ru centre was identified in structural data collected from a crystal incubated for 1 month. It was found in the solvent channel with an occupancy of 0.5, however, with a coordination number of 6. We identified four OH_x ligands coordinated to the Ru centre with an average Ru-O_{water/hydroxido} distance of 2.37 Å and one chlorido ligand with a Ru-Cl distance of 2.48 Å. This [RuCl(OH_x)₄] fragment was found to interact weakly with the peptide backbone through the carbonyl oxygen of Ala107 with a Ru-O_{Ala107} distance of 2.76 Å (Fig. S4, ESI†).

Given the observed cleavage of the cym ligand from the Ru centre, we complemented the X-ray crystallographic studies with mass spectrometry experiments to understand the behaviour of **1** in presence of biomolecules in aqueous solution. Studies with His and HEWL in acetate buffer were conducted over a period of 6 d and 4 weeks, respectively, to mimic the conditions during soaking of HEWL. In the mass spectra recorded for the equimolar reaction mixture of **1** with His, the main peak was identified as



[Ru(cym)(dmb) – H]⁺ followed by [Ru(cym)(dmb) + OAc]⁺ and a formate adduct formed after dilution of the sample with water/acetonitrile/formic acid (69/30/1). In addition minor peaks were detected that were assigned to the His adducts [Ru(cym)(dmb)(His) – H]⁺ (Fig. S8, ESI[†]) and [Ru(dmb)(His)-Cl(CH₃CN) + Na]²⁺, with the latter indicating the cleavage of the cym from the Ru centre. In reaction mixtures with a 5- or 10-fold excess of **1** over HEWL, the main adduct was identified as [HEWL + Ru(cym)(NHC)] while ions could be assigned to a species after cym cleavage. This indicates the influence of the protein microenvironment and soaking conditions on the type of adducts detectable by either method.

An analysis of all the published HEWL structures modified with Ru complexes shows that the primary binding site for most Ru complexes is at His15. However, the nature of the ancillary ligands at the Ru centres determines the modality of binding and potentially the presence and nature of further interactions (see ESI[†], Table S3). Furthermore, in several of those reported structures, the original ancillary ligands, such as indazole, imidazole (and derivatives), DMSO, pyridine, and CO, at the metal centre underwent a ligand exchange reaction with most often only OH_x or Cl ligands modelled. In contrast, π -bound arenes and, in this case, the NHC ligand remain tightly attached to the metal centre, indicating that the organoruthenium compounds retain their ligands to a larger extent than coordination compounds. We have also shown that the Ru(NHC) species are more selective in terms of binding sites compared to Au(NHC) complexes, despite the long soak times. However, we acknowledge that the crystallisation conditions may influence the species formed and that the intermolecular contacts required to maintain the crystal may limit the number of interactions observed. Nevertheless, these are promising results indicating that the NHC ligand is retained in a biological setting and allows for design of stable organoruthenium complexes for medicinal application.

We thank the University of Auckland (Frederick Douglas Brown Postgraduate Science Research Scholarship to M. P. S). D. G. is supported by a Rutherford Discovery Fellowship awarded by the NZ government and administered by the RSNZ. This research was, in part, undertaken on the crystallography beamlines MX1 at the Australian Synchrotron, Victoria, Australia with support from the NZ synchrotron group.

Conflicts of interest

There are no conflicts to declare.

References

- W. A. Wani, S. Prashar, S. Shreaz and S. Gómez-Ruiz, *Coord. Chem. Rev.*, 2016, **312**, 67.
- C. G. Hartinger and P. J. Dyson, *Chem. Soc. Rev.*, 2009, **38**, 391.
- M. A. Jakupec, M. Galanski, V. B. Arion, C. G. Hartinger and B. K. Keppler, *Dalton Trans.*, 2008, 183.
- P. Zhang and P. J. Sadler, *J. Organomet. Chem.*, 2017, **839**, 5.
- H. A. Burris, S. Bakewell, J. C. Bendell, J. Infante, S. F. Jones, D. R. Spigel, G. J. Weiss, R. K. Ramanathan, A. Ogden and D. Von Hoff, *ESMO Open*, 2016, **1**, e000154.
- B. S. Murray, M. V. Babak, C. G. Hartinger and P. J. Dyson, *Coord. Chem. Rev.*, 2016, **306**, 86.
- A. A. Nazarov, C. G. Hartinger and P. J. Dyson, *J. Organomet. Chem.*, 2014, **751**, 251.
- R. E. Morris, R. E. Aird, S. Murdoch Pdel, 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.
- C. S. Allardyce, P. J. Dyson, D. J. Ellis and S. L. Heath, *Chem. Commun.*, 2001, 1396.
- P. J. Dyson, *Chimia*, 2007, **61**, 698.
- Z. Adhireksan, G. E. Davey, P. Campomanes, M. Groessl, C. M. Clavel, H. Yu, A. A. Nazarov, C. H. Yeo, W. H. Ang, P. Droge, U. Rothlisberger, P. J. Dyson and C. A. Davey, *Nat. Commun.*, 2014, **5**, 3462.
- S. M. Meier, D. Kreutz, L. Winter, M. H. M. Klose, K. Cseh, T. Weiss, A. Bileck, B. Alte, J. C. Mader, S. Jana, A. Chatterjee, A. Bhattacharyya, M. Hejl, M. A. Jakupec, P. Heffeter, W. Berger, C. G. Hartinger, B. K. Keppler, G. Wiche and C. Gerner, *Angew. Chem., Int. Ed.*, 2017, **56**, 8267.
- M. Patra, T. Joshi, V. Pierroz, K. Ingram, M. Kaiser, S. Ferrari, B. Spingler, J. Keiser and G. Gasser, *Chem. – Eur. J.*, 2013, **19**, 14768.
- B. Bernhard, *Mini-Rev. Med. Chem.*, 2016, **16**, 804.
- F. E. Hahn and M. C. Jahnke, *Angew. Chem., Int. Ed. Engl.*, 2008, **47**, 3122.
- M. N. Hopkinson, C. Richter, M. Schedler and F. Glorius, *Nature*, 2014, **510**, 485.
- R. Rubbiani, E. Schuh, A. Meyer, J. Lemke, J. Wimberg, N. Metzler-Nolte, F. Meyer, F. Mohr and I. Ott, *MedChemComm*, 2013, **4**, 942.
- A. Pratesi, C. Gabbiani, E. Michelucci, M. Ginanneschi, A. M. Papini, R. Rubbiani, I. Ott and L. Messori, *J. Inorg. Biochem.*, 2014, **136**, 161.
- G. Ferraro, C. Gabbiani and A. Merlino, *Bioconjugate Chem.*, 2016, **27**, 1584.
- A. Casini, C. Gabbiani, F. Sorrentino, M. P. Rigobello, A. Bindoli, T. J. Geldbach, A. Marrone, N. Re, C. G. Hartinger, P. J. Dyson and L. Messori, *J. Med. Chem.*, 2008, **51**, 6773.
- L. Oehninger, M. Stefanopoulou, H. Alborzina, J. Schur, S. Ludewig, K. Namikawa, A. Munoz-Castro, R. W. Koster, K. Baumann, S. Wolf, W. S. Sheldrick and I. Ott, *Dalton Trans.*, 2013, **42**, 1657.
- I. W. McNaie, K. Fishburne, A. Habtemariam, T. M. Hunter, M. Melchart, F. Wang, M. D. Walkinshaw and P. J. Sadler, *Chem. Commun.*, 2004, 1786.
- A. Casini, G. Mastrobuoni, C. Temperini, C. Gabbiani, S. Francese, G. Moneti, C. T. Supuran, A. Scozzafava and L. Messori, *Chem. Commun.*, 2007, 156.
- O. Pinato, C. Musetti, N. P. Farrell and C. Sissi, *J. Inorg. Biochem.*, 2013, **122**, 27.
- C. Muegge, T. Marzo, L. Massai, J. Hildebrandt, G. Ferraro, P. Rivera-Fuentes, N. Metzler-Nolte, A. Merlino, L. Messori and W. Weigand, *Inorg. Chem.*, 2015, **54**, 8560.
- Y. Gothe, T. Marzo, L. Messori and N. Metzler-Nolte, *Chem. – Eur. J.*, 2016, **22**, 12487.
- M. Hanif, S. Moon, M. P. Sullivan, S. Movassaghi, M. Kubanik, D. C. Goldstone, T. Sohnle, S. M. F. Jamieson and C. G. Hartinger, *J. Inorg. Biochem.*, 2016, **165**, 100.
- G. Ferraro, A. Pica, I. Russo Krauss, F. Pane, A. Amoresano and A. Merlino, *J. Biol. Inorg. Chem.*, 2016, **21**, 433.
- L. Messori and A. Merlino, *Coord. Chem. Rev.*, 2016, **315**, 67.
- M. P. Sullivan, M. Groessl, S. M. Meier, R. L. Kingston, D. C. Goldstone and C. G. Hartinger, *Chem. Commun.*, 2017, **53**, 4246.
- F. Zobi and B. Spingler, *Inorg. Chem.*, 2012, **51**, 1210.
- G. Strahs and J. Kraut, *J. Mol. Biol.*, 1968, **35**, 503.
- A. Bijelic, S. Theiner, B. K. Keppler and A. Rompel, *J. Med. Chem.*, 2016, **59**, 5894.
- M. Hollering, M. Albrecht and F. E. Kühn, *Organometallics*, 2016, **35**, 2980.
- N. Cetinbas, M. I. Webb, J. A. Dubland and C. J. Walsby, *J. Biol. Inorg. Chem.*, 2010, **15**, 131.
- B. M. Blunden, D. S. Thomas and M. H. Stenzel, *Polym. Chem.*, 2012, **3**, 2964.
- S. J. Dugan, M. Melchart, A. Habtemariam, S. Parsons and P. J. Sadler, *Inorg. Chem.*, 2006, **45**, 10882.

