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Embedding a Ruthenium-based Structural Mimic of the [Fe]-Hydrogenase Cofactor into Papain

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ABSTRACT: We describe the synthesis of the ruthenacyclic carbamoyl complexes, [Ru(2-NHC(O)C₅H₃NMe)(CO)₂(*o,o*-Me₂-C₆H₃S)(L)] (L = H₂O or MeCN) which have a labile water or acetonitrile ligand at their sixth coordination sites. Steric bulk around the ruthenium centre is essential in preventing isomerization and dimerization, and embedding within papain can be achieved via coordination of its sole free cysteine residue. The observed chemistry parallels that of the natural [Fe]-hydrogenase.

1. INTRODUCTION

Three distinct classes of hydrogenases are known, referred to as the [NiFe]-, [FeFe]-, and [Fe]-hydrogenases, and they all catalyze the reversible oxidation of dihydrogen into protons.¹⁻³ The redox-inactive [Fe]-hydrogenase, or H₂-forming methylene-tetrahydromethanopterin dehydrogenase (Hmd), takes part in the conversion of carbon dioxide to methane in many methanogenic archaea.⁴ In particular, it catalyzes an intermediary step, which is the reversible and stereoselective reduction of a unique biological carbocationic substrate, *N*⁵,*N*¹⁰-methenyltetrahydromethanopterin (methenyl-H₄MPT⁺), with dihydrogen to produce neutral methylenetetrahydromethanopterin (methylene-H₄MPT) and a proton (Scheme 1).⁴⁻⁶

Scheme 1. Reversible reduction of methenyl-H₄MPT⁺.

Crystallographic elucidation of the [Fe]-hydrogenase active site and its apoenzyme have been

achieved, and the former comprises an iron(II) atom with pseudo-octahedral geometry,⁷⁻¹⁰ it is bound to one cysteine residue, two mutually cis CO ligands, and a 2-acetyl-6-hydroxypyridinol ligand. The identity of the sixth ligand, trans to the acyl group, has not been confirmed although it is most probably a water molecule (Figure 1a).¹¹⁻¹³ A number of Fe-based structural models of the active site have been reported over the years, including some which are also functional models.¹⁴ In particular, Hu, *et. al.*, recently showed that reconstitution of the entire [Fe]-hydrogenase using the apoenzyme produced in *E. coli* and an Fe-based model complex could activate dihydrogen.¹⁵ That study points to a role for the protein in [Fe]-hydrogenase for the dihydrogen activation and that the pyridinyl cofactor most likely activates dihydrogen via coordination at the site trans to the acyl group. Such investigations are, nevertheless, hampered by the air-sensitivity of organoiron compounds.

We have recently reported on the first ruthenium and osmium structural mimics for the [Fe]-hydrogenase active site.¹⁶ Ruthenium complexes are generally more air-stable than their iron counterparts, and many are known to activate dihydro-

gen.^{17,18} We therefore hypothesized that a reproduction of the [Fe]-hydrogenase using a more stable ruthenium model complex and an appropriate protein scaffold may serve as a useful model for studying aspects of its chemistry. Such metalloenzymes are also of interest as possibly possessing unique catalytic properties.¹⁹ We chose papain as protein scaffold as it is readily and cheaply available, and its tertiary structure and chemistry are well-understood. In addition, it has a single free sulfhydryl (Cys25) moiety located in a pocket forming the enzyme's active site which may allow binding to the ruthenium centre in an analogous way to that present in native [Fe]-hydrogenase. Our investigations into this are presented here.

Figure 1. Structure of (a) the active site of [Fe]-hydrogenase, (b) ruthenium analogue of the active site of [Fe]-hydrogenase, and (c) ruthenium analogue of the entire enzyme synthesized in this study.

2. RESULTS AND DISCUSSION

Synthesis. The ruthenium-based structural mimics **3** can be synthesized in two steps from the corresponding ionic carbamoyl ruthenacycles **1**, by following a recently reported procedure (Scheme 2).¹⁶

Scheme 2. Synthesis of **2** and **3**.

Unlike the iron analogue, thiolation of **2** requires the addition of a base.^{14d} The mononuclear thiolato complexes **3** could only be isolated when the pyridine ligand is substituted at the 6-position, together with a bulky thiolate (**3bB**); dimerization occurs otherwise (**4aA**, **4aB** and **4bA**). All the complexes **2-4** have been characterized spectroscopically and crystallographically; ORTEP plots of a representative of each are given in Figure 2.

The complexes **4** are sparingly soluble in dimethyl sulphoxide but not in other common organic solvents. All three are symmetrical Ru^{II}-Ru^{II} homodimers bridged by two thiolato ligands but while **4aA** and **4aB** are centrosymmetric, the structure of **4bA** has approximate C₂ symmetry with the two tolyl groups (as are the two pyridyl groups) on the same side of the slightly puckered Ru₂S₂ ring. Their ¹H NMR spectra in d₆-DMSO show only one set of peaks; this is similar to the dimeric iron compound [{Fe(COMe)(CO)₂(μ²-SPyMe)}₂]^{14b}. As in the case of the iron analogues, attempts to cleave the Ru-S bridges with phosphines (PMe₃ and PPh₃) failed.^{14d}

Figure 2. ORTEP plots showing the molecular structures of (a) **2b**, (b) **3bB**, and (c) **4aB** (left) and **4bA** (right). Thermal ellipsoids are drawn at the 50% probability level, and all hydrogen atoms have been omitted for clarity.

Isomerism. Steric effects of the 6-substituent on the pyridine ring are apparent in the bond parameters of **2a** and **2b**. These include an increasing angle between the pyridine ring and the acetonitrile, and a longer Ru-N1 bond in moving from R = H (**2a**) to Me (**2b**); $\angle \text{N1-Ru-N2} = 92.73(8)^\circ$ and $101.6(1)^\circ$, and $d(\text{Ru-N1}) = 2.110(2)$ and $2.163(3)$ Å, respectively. There is also greater distortion of the five-membered metallacycle, which does not lie directly above the Ru-N2 bond, in **2b**; the N(2)-Ru-N(1)-R torsion angles are -2.43° and -11.40° in **2a** and **2b**, respectively. One chemical consequence of this steric interaction is that **2b** is present as one isomer in solution while **2a** exists as a mixture of two isomers; the IR spectrum of **2b** shows only two bands for the CO stretches, while that for **2a** shows a split in the lower frequency band (*vide infra*). Similarly, the ^1H NMR spectrum of **2a** shows two sets of resonances, in a 55:45 ratio, but only one for **2b**. The observation of two isomers in **2a** is also in agreement with an earlier report on an iron analogue.^{14d} A series of VT ^1H NMR spectra of **2a** shows broadening and coalescence at higher temperatures, indicating isomerization at the NMR timescale (Figure 3).

Figure 3. Top: Computed free energies (in kJ/mol) of the two lowest energy isomers of **2a** and **2b** relative to the most stable isomer (isomer 1). Bottom: VT ^1H NMR spectra of **2a** in d_3 -acetonitrile.

The exchange rate and activation energy barrier estimated from the coalescence of the doublet at ~ 8.35 ppm are 12 s^{-1} and 75 kJ mol^{-1} , respectively, at the coalescence temperature of 70°C . In contrast, the VT ^1H NMR spectra of **2b** show no significant broadening of the resonances, except for the NH resonance (Figure S9), consistent with the presence of essentially one isomer.

A computational study carried out on all the seven possible stereoisomers of **2a** and **2b** (Tables S6 and S7) shows that the energy gap between the two lowest energy isomers (Figure 3 and Table S8) is about 1 and 8 kJ mol^{-1} , which corresponds to an isomeric ratio of 60:40 and 96:4 for **2a** and **2b**, respectively. These values are in very good agreement with those from NMR integration. The computed values for the CO vibrations also point to a larger difference ($\sim 10\text{ cm}^{-1}$) in the lower frequency band between the isomers (Table S6), in agreement with the observed splitting of this band. The major isomer for both **2a** and **2b** from the computational study (isomer 1) corresponds to that observed in the solid-state structure, thus the minor isomer in **2a** is presumably isomer 2. The relative stability of the isomers can be rationalized on the basis of the *trans* influence: $\text{NHCO} > \text{CO} > \text{Br}^- > \text{MeCN} > \text{Py}$.²⁰ Thus, the strongly π -accepting CO and the strongly σ -donating carbamoyl prefer to be *trans* to weak π -acceptors or σ -donors such as Br^- , MeCN or Py. The larger isomer energy gap in **2b** may be attributed to steric interaction between the bulky bromide ligand and the methyl substituent of the pyridine ring.

Dissolution in polar solvents. Complex **3bB** is a ruthenium-based structural mimic of the [Fe]-hydrogenase active site, with acetonitrile in place of the unknown sixth ligand.^{14d} The infrared spectrum (in acetonitrile) shows two intense absorption bands at 2047 and 1978 cm^{-1} ($\Delta\nu = 69\text{ cm}^{-1}$) for the *cis*-dicarbonyl ligands which are blue-shifted from that for the previously reported iron analogues ($\sim 20\text{ cm}^{-1}$),^{14d} and the [Fe]-hydrogenase ($\sim 15\text{ cm}^{-1}$),¹¹ consistent with ruthenium being less electron-rich.^{21,22}

Although complexes **2** and **3** are not very soluble in most of the common organic solvents, their solubility in acetone and tetrahydrofuran increased upon the addition of small amounts of water. It is also noted that the CO vibrations of **2b** in acetone,

for example, are blue shifted with increasing amounts of water added (Figure S4), and a similar shift is observed for **3bB** in acetonitrile (Figure S5). These shifts are indicative of changes in composition and in the case of **2b** in acetonitrile, may be attributed to the formation of a new species upon the addition of water, which has tentatively been identified as a cationic species in which both the Br⁻ and MeCN ligands have been replaced by water molecules; the experimental ν_{CO} values at 2086 and 2027 cm⁻¹ are in good agreement with the values of 2089 and 2028 cm⁻¹ calculated for this species. (Figure 4).

Figure 4. (Top) IR spectra of **2b** in acetonitrile (—), and various compositions of acetonitrile/water (v/v): 10:1 (---); 5:2 (·····); 1:2 (-·-·-). (Bottom) Proposed exchange process. Free energies are given in kJ/mol for the forward reactions.

Computational studies on both **2b** and **3bB** also suggest that replacement of the acetonitrile ligand with water will lead to a decrease in the frequency difference between the two CO vibrational modes, as observed in the experimental spectra (Table S5). The weak Ru-OH₂ bond is corroborated by

the mass spectrum; the molecular ion peak for **2b** can be observed at ambient temperature but not at higher temperatures. The resonance at 2.10 ppm for the metal-bound CH₃CN ligand in the ¹H NMR spectrum (in acetone-d₆) shifts to 2.05 ppm, corresponding to free acetonitrile, upon addition of water (Figure S12).²³ A DFT calculation gives a ΔG° of -0.1 kJ mol⁻¹ for the reaction: **2b** + H₂O → **2b.H₂O** + CH₃CN (in acetone). All these indicate that the sixth (acetonitrile) ligand in **2** and **3** is very weakly bound, reminiscent of the situation in the [Fe]-hydrogenase.^{8,10}

Bioconjugation to papain. That complexes **2** can be thiolated suggests that they may also be incorporated into a protein via reaction with a free sulfhydryl moiety. Papain contains a free sulfhydryl moiety borne by Cys25 which, together with His159 are the two essential aminoacids responsible for its proteolytic activity.¹⁹ Conjugation to the sulfhydryl group of Cys25 can thus be readily monitored via a fall in its enzymatic activity. As histidine residues such as His159 may also be involved in coordination owing to their imidazole group, a preliminary assessment of the reactivity of **2b** with imidazole was first carried out.

We found that **2b** reacts with imidazole to give **2b.Im**, and this can in turn react with N-acetyl-L-cysteine-methyl ester (**C**) to afford the dimer **4bC**; the latter can also be obtained directly from the reaction of **2b** and **C** (Scheme 3). The complex **3bB** reacts similarly with imidazole to form **3bB.Im**. Both the imidazole complexes **2b.Im** and **3bB.Im**, however, slowly convert back to **2b** and **3bB**, respectively, on standing in acetonitrile (Figures S7 and S8). These results suggest that conjugation of **2b** to a cysteine side chain should be favoured over histidine.

Scheme 3. Reaction of **2b** and **3bB** with imidazole and N-acetyl-L-cysteine-methyl ester.

Conjugation of **2b** to papain was assessed through measurement of its effect on the hydrolysis rate of N- α -carbobenzoxy-glycine p-nitrophenyl ester (ZGlyONP).²⁴ The activity dropped to almost zero, suggesting that Cys25 has been affected (Figure S16). After a gel filtration step to remove excess complex, bioconjugate **5** was analyzed by several spectroscopic techniques. The UV-visible spectrum of **5** (Figure S17) shows two peaks at 279 and $\sim$ 318 nm ascribable to papain,²⁵ and the ruthenium center, respectively. The IR spectrum of **5** shows that the characteristic carbonyl bands are red-shifted (2064 and 2000 cm^{-1}) from that of **2b** (Figure 5). This red shift is similar to the shift in comparing **2b** with **3bB**, and the smaller frequency gap between the two stretches (64 cm^{-1}) is also consistent with thiolation. Indeed, the frequency gap is the same as that computed for **3bB.H₂O** (corresponding to replacement of the acetonitrile ligand in **3bB** with water), suggesting that **5** contains thiolato and water ligands (Table S5).

Figure 5. ATR spectra of **2b** (gray dotted line) and **5** (black full line).

While the electrospray-high resolution mass spectrum (ESI-HRMS) of papain reveals a charge states distribution from 22+ (m/z 1065.69) to 12+ (m/z 1952.82), centred around 18+ (m/z 1302.28) and 17+ (m/z 1378.82) (Figure 6A), that for **5** shows charge states ranging from 22+ (m/z 1078.81) to 12+ (m/z 1980.15) (Figure 6B). After deconvolution and deisotoping, the mass spectrum of papain showed a major peak at a monoisotopic mass of 23,407.52205, in good agreement with the theoretical value (23,407.52457 $\text{g}\cdot\text{mol}^{-1}$; 0.2 ppm) (Figure S15A). Another species at a mass of 23,437.52205, corresponding to an amino acid

variant with Thr in place of an Ala residue (mass shift of +30.01056) was also observed. In comparison, that for **5** showed two prominent additional species at 23,699.47624 and 23,729.48682, corresponding to an exact mass shift of 291.95417 (Figure S15B). This value matches very well with the conjugation of **2b** to papain by substitution of the bromide ligand by a thiolate (and loss of MeCN). Interestingly, no other species at higher mass was observed, indicating monometallation of the protein.

Figure 6. ESI-HRMS mass spectra of (A) papain and (B) bioconjugate **5**, both at 2.15 μM in water/acetonitrile/formic acid (49:50:1, v/v/v).

3. CONCLUDING REMARKS

In this study, we showed that organoruthenium complexes of the general formula $[\text{Ru}(2\text{-NHC}(\text{O})\text{C}_5\text{H}_3\text{NR})(\text{CO})_2(\text{X})(\text{L})]$ (where $\text{R} = \text{H}, \text{Me}$, $\text{X} = \text{Br}$ or SR' , and $\text{L} = \text{H}_2\text{O}$ or NCCH_3) exhibited chemistry that is relevant to understanding the chemistry of the [Fe]-hydrogenase - the lability of the so-called sixth ligand, and the importance of steric bulk in preventing dimerization via thiolate bridges through this coordination site. We believe that embedding of the metal cofactor in a protein cavity is how nature achieves this, and have shown that this can be readily achieved with the complexes reported here. The extension of this idea to the construction of metalloenzymes is ongoing.

4. EXPERIMENTAL SECTION

General procedures. All experiments were performed under an argon atmosphere using standard Schlenk techniques except for the bioconjugation experiments. Chemicals were obtained from Sigma Aldrich and used as received. Papain was

obtained from Calbiochem and purified by gel filtration on a gel desalting column (Hiprep desalt 26/10, GE Healthcare) using 0.15 M NaCl solution as eluent to remove low molecular weight thiols present in the commercial sample.^{24,26} Compounds **1** were synthesized as previously described.¹⁶ Solvents used for reactions were distilled over the appropriate drying agents under argon before use. ESI-HRMS for compounds were recorded on a Waters UPLC-Q-ToF MS mass spectrometer while that for papain and bioconjugate **5** were performed on a LTQ-Orbitrap XL. ¹H and ¹³C{¹H} NMR spectra were obtained on a JEOL 400 MHz spectrometer at room temperature unless stated otherwise. Chemical shifts for ¹H and ¹³C{¹H} were referenced with respect to the residual resonances of the respective deuterated solvents. Infrared spectra were recorded on a Bruker Alpha FT-IR spectrometer in an IR cell with NaCl windows and a path length of 0.1 mm at a resolution of 2 cm⁻¹; for the bioconjugation experiments, the spectra were recorded after deposition of the solutions onto nitrocellulose membrane and air-dried. Absorbance spectra were recorded using a Varian Cary 50 spectrometer. Elemental analyses were performed on a PerkinElmer 2400 Series II CHN/O Elemental Analyzer in house.

Preparation of [RuBr(2-NHC(O)C₅H₄N)(CO)₂(NCMe)] (2a). A sample of **1a** (200 mg, 0.38 mmol) was dissolved in acetonitrile (50 mL). Slow evaporation of the solvent at room temperature afforded white crystal of **2a**. Yield: 100 mg, 0.26 mmol, 68 %. mp 343–344 °C. IR (acetonitrile, cm⁻¹): ν_{CO} 2061 (vs), 1999 (s), 1989 (s), 1622 (m). ¹H NMR (CD₃CN): δ = 9.10–9.09 (d, 1H, *J* = 5.5 Hz, PyH), 8.71–8.69 (br, 1H, NH), 8.54–8.552 (br, 1H, NH), 8.37–8.36 (d, 1H, *J* = 5.5 Hz, PyH), 7.89–7.80 (m, 2H, PyH), 7.06–7.00 (m, 4H, PyH), 1.96 (s, 6H, Me) ppm. ¹³C{¹H} NMR (CD₃CN): 198.8 (CO), 198.2 (CO), 195.8 (CO), 195.6 (CO), 195.4 (NHCO), 191.2 (NHCO), 158.0, 157.5, 148.6, 147.5, 141.2, 140.6, 116.8, 116.5, 109.3, 109.5 ppm. HRMS. Calcd for [M]⁺: *m/z*: 399.8871. Found: *m/z* 399.8876. Anal. Calcd for C₈H₅N₂O₃BrRu (M-CH₃CN): C 26.83, H 1.41, N 7.82. Found C 26.90, H 1.86, N 7.21. Complex **2b** was similarly prepared from **1b** (213 mg, 0.38 mmol). Yield: 113 mg, 0.27 mmol, 72 %. mp 281–282 °C. IR (acetonitrile, cm⁻¹): ν_{CO} 2061 (vs), 1988 (s), 1619 (m). ¹H NMR (CD₃CN): δ =

8.52–8.50 (br, 1H, NH), 7.67–7.63 (t, 1H, *J* = 7.8 Hz, PyH), 6.88–6.86 (t, 1H, *J* = 7.8 Hz, PyH), 6.82–6.80 (d, 1H, *J* = 8.2 Hz, PyH), 2.65 (s, 3H, Me), 1.98 (s, 3H, Me) ppm. ¹³C{¹H} NMR ((CD₃)₂CO): 195.1 (CO), 191.8 (CO), 159.5 (NHCO), 158.4, 140.2, 117.2, 116.6, 106.9, 25.5 ppm. HRMS. Calcd for [M+H]⁺: *m/z*: 413.9027. Found: *m/z* 413.9021. Anal. Calcd for C₁₁H₁₀N₃O₃BrRu: C 31.98, H 2.44, N 10.17. Found C 31.70, H 2.40, N 10.48.

Preparation of [Ru(2-NHC(O)C₅H₃NMe)(SC₆H₃Me₂)(CO)₂(NCMe)] (3bB). Triethylamine (7 μL, 0.05 mmol) and 2,6-dimethylbenzenethiol (5 μL, 0.05 mmol) were added into a solution of **2b** (21 mg, 0.05 mmol) in acetonitrile (5 mL). The resulting solution was then stirred for 5 min at room temperature. Slow evaporation of the solvent at room temperature afforded **3bB** as yellow crystals. Yield: 13 mg, 0.03 mmol, 55 %. IR (MeCN, cm⁻¹): ν_{CO} 2047 (vs), 1978 (s). ¹H NMR (CD₃CN): δ = 8.45 (br, 1H, NH), 7.64–7.60 (t, 1H, *J* = 7.4 Hz, PyH), 6.92–6.90 (d, 2H, *J* = 7.8 Hz, PyH), 6.83–6.75 (m, 3H, PyH), 2.44 (s, 3H, Me), 2.24 (s, 6H, Me), 1.96 (s, 3H, Me) ppm. ¹³C{¹H} NMR (CD₃CN): 159.5, 143.4, 139.6, 129.9, 123.3, 126.3, 123.5, 116.4, 106.4, 22.2, 20.9 ppm. HRMS. Calcd for [M+H]⁺: *m/z*: 472.0520. Found: *m/z* 472.0539. Anal. Calcd for C₁₉H₁₉N₃O₃SRu: C 48.50, H 4.07, N 8.93, S 6.81. Found C 48.47, H 4.46, N 9.21, S 6.73.

Preparation of [Ru(2-NHC(O)C₅H₄N)(SC₆H₄Me)(CO)₂]₂ (4aA). Triethylamine (5 μL, 0.05 mmol) and 4-methylbenzenethiol (6 mg, 0.05 mmol) were added into a solution of **2a** (20 mg, 0.05 mmol) in acetonitrile (5 mL) at room temperature. A white precipitate was formed immediately, and the solution was left to stir for 2 h. The white precipitate was separated by filtration and washed with acetonitrile (20 mL) to afford analytically pure **4aA**. Yield: 25 mg, 0.03 mmol, 60 %. IR (MeCN, cm⁻¹): ν_{CO} 2053 (m), 2037 (s), 1991 (m), 1970 (s). ¹H NMR ((CD₃)₂SO): δ = 10.70 (br, 1H, NH), 8.35–8.34 (d, 1H, *J* = 5.04 Hz, PyH), 8.00–7.95 (m, 1H, PyH), 7.47–7.45 (d, 2H, *J* = 8.72 Hz, PhH), 7.16–7.14 (d, 1H, *J* = 8.24 Hz, PyH), 7.05–7.03 (d, 2H, *J* = 5.04 Hz, PhH), 6.93–6.89 (m, 1H, *J* = 5.04 Hz, PyH), 2.25 (s, 3H, Me) ppm. ¹³C{¹H} NMR ((CD₃)₂SO): 201.9, 201.4, 193.2, 158.5, 148.1, 141.3, 135.2, 134.6, 132.3, 129.5, 118.6, 110.5, 21.0 ppm.

HRMS. Calcd for $[M+H]^+$: m/z : 804.9312. Found: m/z 804.9329.

Complexes **4aB** and **4bA** were prepared in similar yields following an analogous procedure.

4aB. IR (KBr, cm^{-1}): ν_{CO} 2041 (s), 1979 (s). HRMS. Calcd for $[M+H]^+$: m/z : 832.9625. Found: m/z 832.9624. MS. Calcd for $[M+Na]^+$: m/z : 853.7. Found: m/z 853.8. Anal. Calcd for $\text{C}_{32}\text{H}_{28}\text{N}_4\text{O}_6\text{S}_2\text{Ru}_2$: C 46.26, H 3.40, N 6.74, S 7.72. Found C 46.33, H 3.33, N 6.98, S 7.72.

4bA. IR (KBr, cm^{-1}): ν_{CO} 2051 (m), 2038 (s), 1988 (m), 1975 (s). ^1H NMR (d_6 -DMSO): δ = 10.50 (br, 1H, NH), 7.80-7.77 (t, 1H, J = 8.2 Hz, PyH), 7.52-7.51 (d, 2H, J = 8.2 Hz, PhH), 7.03-7.01 (d, 2H, J = 7.8 Hz, PhH), 6.90-6.91 (d, 1H, J = 4.9 Hz, PyH), 6.82-6.80 (d, 1H, J = 8.2 Hz, PyH), 2.49 (s, 3H, Me), 2.24 (s, 3H, Me) ppm. $^{13}\text{C}\{^1\text{H}\}$ NMR (d_6 -DMSO): 202.3, 199.7, 194.1, 160.7, 159.4, 141.1, 136.8, 134.4, 131.4, 129.5, 118.6, 108.1, 28.3, 20.9 ppm. HRMS. Calcd for $[M+H]^+$: m/z : 832.9625. Found: m/z 832.9626.

Preparation of [RuBr(2-NHC(O)C₅H₃NMe)(CO)₂(C₃H₄N₂)] (2b.Im). A sample of imidazole (1.6 mg, 0.02 mmol) was added into a solution of **2b** (10 mg, 0.02 mmol) in methanol (5 mL). The resulting mixture was then stirred for 30 min at ambient temperature. Slow evaporation of the solvent afforded **2b.Im** as a crystalline solid. Yield: 3 mg, 0.01 mmol, 34 %. IR (MeOH, cm^{-1}): 2062 (s), 1998 (s). ^1H NMR (d_6 -acetone): δ 11.95 (br, 1H, NH), 9.66 (br, 1H, NH), 8.24 (s, 1H, CH), 7.71-7.67 (t, 1H, J = 7.8 Hz, PyH), 7.25-7.20 (d, 2H, J = 10.5 Hz, CH), 6.93-6.87 (m, 2H, PyH), 3.06 (s, 3H, Me) ppm. $^{13}\text{C}\{^1\text{H}\}$ NMR (d_6 -acetone): 202.2, 197.4, 196.1, 162.0, 158.9, 140.4, 139.4, 129.0, 117.7, 117.0, 106.6, 25.0 ppm. HRMS. Calcd for $[M+H]^+$: m/z : 440.0828. Found: m/z 440.0812. Anal. Found C 34.37, H 2.80, N 13.59; Calculated for $\text{C}_{12}\text{H}_{11}\text{N}_4\text{O}_3\text{BrRu.C}_2\text{H}_3\text{N}$: C 34.94, H 2.93, N 14.55.

Preparation of [Ru(2-NHC(O)C₅H₃NMe)(SCH₂CH(NHCOMe)CO₂Me)(CO)₂]₂ (4bC). Triethylamine (3 μL , 0.02 mmol) and N-acetyl-L-cysteine-methyl ester (4.3 mg, 0.02 mmol) were added into a solution of **2b** (10 mg, 0.02 mmol) in methanol (5 mL). The resulting mixture was stirred for 30 min and then the volume reduced under vacuum to approximately 1 mL. After standing for 3 d, complex **4bC** (9.0 mg,

0.01 mmol, 48 %) was obtained as a yellow crystalline solid. IR (MeOH, cm^{-1}): 2052 (s), 2038 (s), 1991 (s), 1978(s). ^1H NMR (d_3 -acetonitrile): δ 8.90 (br, 1H, NH), 7.71-7.66 (m, 1H, PyH), 6.90-6.86 (m, 1H, PyH), 6.73-6.68 (m, 1H, PyH), 4.44-4.39 (m, 1H, CH), 3.59 (s, 3H, Me), 3.27-3.18 (m, 2H, CH₂), 2.16 (s, 3H, Me), 1.88 (s, 3H, Me) ppm. HRMS. Calculated for $[M+H]^+$: m/z : 938.9851. Found: m/z 938.9877.

Purification of papain by affinity chromatography.²⁷ The peptide GGYR (Bachem) was attached to CNBr-activated Sepharose 4B (Sigma) via its terminal NH₂ group. The resin (20 mL) was packed into a column (XK 16, GE Healthcare) and conditioned with EDTA (20 mM), mercaptoethanol (10 mM) pH 4.3, at 1 mL/min. Papain (200 mg solid, Calbiochem) was dissolved in EDTA (25 mM), mercaptoethanol (30 mM) pH 4.3 (20 mL), and the solution was incubated for 30 min. The solution (18 mL) was injected into the column, and the resin was washed successively with conditioning buffer (2 volumes) then with washing buffer (5 mM EDTA, pH 6.0, 2 volumes). Papain was finally eluted with water (2 volumes). Protein concentration was calculated from the absorbance at 280 nm taking an extinction coefficient of 56,000 $\text{M}^{-1}\cdot\text{cm}^{-1}$.

Enzymatic assay. Affinity-purified papain (1.6 mL of a 3.2 μM solution in water; final concentration = 2.56 μM) was mixed with 50 mM phosphate buffer pH 7.8 and **2b** (20 μL of a 10 mM solution; final concentration = 100 μM). 20 μL of the mixture was used to measure the hydrolytic activity of ZGlyONP (50 μL) in 430 μL of phosphate buffer pH 6.5 at various times. The remaining mixture was incubated at RT for 18 h and submitted to gel filtration on a dextran desalting column (10 mL, Thermo scientific). Species were eluted with water. 1-mL fractions were collected and their absorbance at 280 nm measured with a spectrophotometer. The fractions containing the protein were pooled and the solution (3 mL; 1.25 μM) was concentrated by ultrafiltration on Amicon Ultra 4 (10 kDa cutoff, Millipore) to a final volume of 0.22 mL. A control experiment using papain was also carried out to measure the hydrolytic activity of ZGlyONP.

High resolution mass spectrometry of papain samples. ESI-HRMS experiments were carried

out using a LTQ-Orbitrap XL from Thermo Scientific (San Jose, CA, USA) and operated in positive ionization mode, with a spray voltage at 4.5 kV. Native papain or bioconjugate **5** in a water/acetonitrile/formic acid (49:50:1, v/v/v) mixture was continuously infused using a 250 μ L syringe at 5 μ L/min flow. Applied voltages were 40 and 100 V for the ion transfer capillary and the tube lens, respectively. The ion transfer capillary was held at 275 °C. Sheath and auxiliary gas were set at a flow rate of 45 and 15 arbitrary units (a.u.), respectively. Detection was achieved in the Orbitrap with a resolution set to 100,000 (at m/z 400) and a m/z range between 200–3000 in profile mode. Spectrum was analyzed using the acquisition software XCalibur 2.1 (Thermo Fisher Scientific, Courtaboeuf, France) without smoothing and background subtraction. The automatic gain control (AGC) allowed accumulation of up to 2.10^5 ions for FTMS scans. Maximum injection time was set to 500 ms and 1 μ scan were acquired.

Crystallographic studies. Diffraction-quality crystals were grown from acetonitrile (**2a**, **2b**, **3bB**, **4aA**, **4aB** and **4bA**) mounted on quartz fibers. X-ray diffraction data were collected at 103(2) K on a Bruker X8 APEX system, using Mo K α radiation, with the SMART suite of programs.²⁸ Data processing and correction were made for Lorentz and polarization effects with SAINT,²⁹ and for adsorption effects with SADABS.³⁰ Twinning was observed for the crystals of **4bA**; the twin laws and cell parameters were determined using the program CELL_NOW,³¹ and the reflection data processed and corrections applied using the program TWINABS.³² All the structural solution and refinement were carried out with the SHELXTL suite of programs.³³ The structures were solved by direct methods to locate the heavy atoms, followed by successive difference maps for the light, non-hydrogen atoms. Disorder of a carbonyl and the bromide ligand was found for **2b** which were modelled with two sites each, with occupancies summed to unity. All non-hydrogen atoms were refined with anisotropic displacement parameters in the final model. All the organic hydrogen atoms were placed in calculated positions and refined with a riding model. The crystallographic data are summarized in Tables S1 and S2.

Computational studies. The computational studies were carried out using density function theory (DFT), utilizing the hybrid functional of Truhlar and Zhao (M06).³⁴ The basis set used was def2TZVP for ruthenium,³⁵ and 6-311+G(2d,p) for the light atoms. For calculations involving solvated species, the polarized continuum model (PCM) was employed. Spin-restricted calculations were used for structural optimization, harmonic frequency calculations and to evaluate zero-point energy (ZPE) corrections. Optimized geometries were characterized as equilibrium structures with all real frequencies. All calculations were performed using the Gaussian 09 suite of programs.³⁶ The coordinates for parent complexes were taken from the crystal structure and each of the other isomers was modified from this.

ASSOCIATED CONTENT

Supporting Information

CCDC 1820446-7, 1839988 and 1839990-92 contain the supplementary crystallographic data for the structures reported. These are available from the Cambridge Crystallographic Data Centre. NMR spectra showing presence of isomers, crystallographic tables, and optimized geometry of computed structures. Experimental procedure and high-resolution mass spectra of papain and bioconjugate. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Graphic

Synopsis

The ruthenacyclic carbamoyl complexes $[\text{Ru}(2\text{-NHC(O)C}_5\text{H}_3\text{NMe})(\text{CO})_2(o,o\text{-Me}_2\text{-C}_6\text{H}_3\text{S})(\text{L})]$ ($\text{L} = \text{H}_2\text{O}$ or MeCN) have a labile ligand L. Substitution at the 6-position of the pyridine and the 2,6-positions of the thiol ligands are important in preventing dimerization and isomerization of the complexes. These complexes can be embedded into papain via coordination of its only free cysteine residue to produce a structural mimic of the complete [Fe]-hydrogenase.