

Copper complexes as a source of redox active MRI contrast agents.

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Abstract

The study reports an advance in designing copper-based redox sensing MRI contrast agents. Although the data demonstrate that copper(II) complexes are not able to compete with lanthanoids species in terms of contrast, the redox-dependent switch between diamagnetic copper(I) and paramagnetic copper(II) yields a novel redox-sensitive contrast moiety with potential for reversibility.

Introduction

MRI contrast agents capable of detecting redox status in biological tissues would be valuable in the diagnosis of inflammatory diseases that involve an oxidative pathology, such as atherosclerosis or rheumatoid arthritis (Filippin et al. 2008; Profumo et al. 2011). Historically, contrast agents have depended on lanthanide chemistry (De Leon-Rodriguez et al. 2009; Aime et al. 2009) and the development of agents for MRI continues to focus largely on gadolinium owing to its key qualities; i.e. it is a quadrupolar nucleus, it has a large number of unpaired electrons and when coordinatively unsaturated it displays rapid exchange kinetics in aqueous solution (Table 1). However, the synthesis of new, smart imaging agents for measuring biochemical processes (e.g. pH and pO_2) requires that the contrast agent has added capability (Angelovski et al. 2008; Iwaki et al. 2012; Viger et al. 2013; Vologdin et al. 2013). A method of introducing added capability is to modify the ligand that chelates the lanthanoid centre (Angelovski et al. 2008; Iwaki et al. 2012; Vologdin et al. 2013). Alternatively, it is possible to encapsulate a gadolinium species into a polymer matrix which degrades in response to the biological environment (Viger et al. 2013).

- Table 1 in here-

Recent concerns about the nephrotoxicity of gadolinium suggest that there would be advantages in broadening the range of metals used as contrast agents (Badero et al. 2008; Ledneva et al. 2009). The clinical use of transition metals in MRI has been mostly restricted to nanoparticles which, as large multimetallic assemblies, are able to compensate for the weak relaxivity of the parent elements (e.g. iron, copper) (Di Marco et al. 2007; Longmire et al. 2008; Kueny-Stotz et al. 2012; Min et al. 2012). One strategy is to design modified low T_2 nano-particles which assemble in the presence of key biomarkers (e.g. streptavidin-biotin) to form assemblies with a high T_2 (Min et al. 2012).

Certain transition metals (Table 1) have physical properties compatible with their development as contrast agents (Tobe 1972; Weast 1980; Grant et al. 1997; Lejeune 1999; Yam & Lo 1999; Dorazio & Morrow 2012; Drahos et al. 2012; Thunus & Yu 2012; Loving et al. 2013; Gale et al. 2014; Tsitovich et al. 2014). Since transition metals display different coordination chemistry to gadolinium (lower coordination numbers, simpler topologies) and can combine with a wider range of ligand types (e.g. phosphines, thiolates), it should be easier to target them to physiological parameters such as pH and redox status.

Due to its large number of unpaired electrons, manganese(II) has attracted the most attention as an alternative to gadolinium (Grant et al. 1997; Thunus & Lejeune 1999; Yam & Lo 1999; Yu et al. 2012; Loving et al. 2013; Gale et al. 2014). Indeed, there has been some success with manganese in the design of redox sensing contrast agents (Loving et al. 2013; Gale et al. 2014; Tsitovich et al. 2014). Copper(II) also has some attractive properties (Table 1) (Tobe 1972; Weast 1980). Although it only has one unpaired electron, its ligand exchange rate is higher by two orders of magnitude than manganese and commensurate with that of gadolinium (table 1). Furthermore, its chemistry suggests that it should be easier to create an available coordination site for water without the risk of forming insoluble oxides (c.f. MnO_2 , Mn_2O_3). Another advantage of copper over manganese is the availability of a diamagnetic

form, d^{10} Cu(I), which can be sustained under biological conditions. The availability of diamagnetic (Cu(I)) and paramagnetic (Cu(II)) forms allows the design of agents that have redox switch capability. Lauffer briefly discussed the properties of copper as a platform for the design of imaging agents 25 years ago, deeming it unsuitable for use due to its poor relaxivity (Lauffer 1987). However, the advent of better imaging software, access to higher field strengths, more sophisticated pulse sequences and the emergence of chemical exchange saturation transfer (CEST) as a relaxation mechanism, the need for metals with such high relaxivity may be coming to an end (De Leon-Rodriguez et al. 2009; Aime et al. 2009; Hancu et al 2010; Sherry & Wu 2013). The design of MRI contrast agents has developed significantly over the past 25 years and it is not uncommon for discrete sensing moieties to be attached directly to a MRI reporting gadolinium species. Thus, should copper remain an unsuitable metal in its own right, there are opportunities to bolster its poor relaxivity by combining a reporting copper complex with a more responsive gadolinium species. Although the relaxivity of copper will never supersede that of manganese or gadolinium, its potential as a switchable redox sensing contrast moiety makes it an interesting target and, considering all the advances in the field of MRI over the past 25 years, a topic worth re-visiting.

Experimental

Unless otherwise stated, all chemicals were commercially obtained and used without further purification. Gd(DTPA) and $\text{Cu}(\text{BF}_4)_2$ were purchased from the Sigma-Aldrich company UK. $\text{S}_{\text{pr}}\text{N}_{\text{en}}$ and the unmodified copper macrocycles ($[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{en}}]^{2+}$, $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{pr}}]^{2+}$, $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$ and $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{pr}}]^{2+}$) were prepared as previously reported (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al. 2013). Solid reflectance spectra (400 – 900 nm) were recorded on a Photonics CCD array UV-Vis spectrophotometer. Electrospray mass spectra were recorded using a Thermo

Finnigan LCQDuo instrument equipped with anion trap as described previously (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al 2013). The spectrophotometric titrations were carried out on a Unicam UV 300 spectrometer.

X-Ray crystallography: Crystals were coated in mineral oil and mounted on glass fibres. Data were collected at 123 K on a Nonius Kappa CCD diffractometer using graphite monochromated Mo-K α radiation. The heavy atom positions were determined by Patterson methods and the remaining atoms located in the difference electron density maps. Full matrix least squares refinement was based on F₂ with all non-hydrogen atoms anisotropic. While the hydrogen atoms were mostly observed in the difference maps, they were placed in calculated positions riding on the parent atoms. The structure solution and refinement used the programs SHELX-97 and the graphical interface WinGX (Sheldrick 1998; Farrugia 1999)

Preparation of N-substituted macrocycles: S_{pr}N_{en} (1.04 g, 0.003 mol) was dissolved in acetonitrile (100 mL) to which sodium bicarbonate (5.07g, 0.06 mol) was added. The mixture was stirred and heated to reflux. Halo-alkane (2-bromo-ethanol, 3-bromo-propan-1-ol or bromoacetic acid, 2 mol equiv.) in acetonitrile (50 mL) was added dropwise to the refluxing mixture over a 6 hr period. The mixture was then refluxed overnight. The solution was left to cool, before being filtered. The solvent was removed *in vacuo*, resulting in a yellow waxy solid. The products were used as received. Yields were typically in the order of 70%.

cis-S_{pr}N_{en}(-(CH₂)₂OH)₂: Mass Spec. (ESI) m/z 455.27 [M+Na]⁺.

cis-S_{pr}N_{en}(-(CH₂)₃OH)₂: Mass Spec. (ESI) m/z 461.14 [M+H]⁺.

cis-S_{pr}N_{en}(-CH₂COOH)₂: Mass Spec. (ESI) m/z 461.13 [M+H]⁺.

Preparation of N-substituted copper(II) macrocycles: The required N-substituted macrocycle (1.04 g) was dissolved in methanol (150 mL) and treated with an equimolar amount of Cu(BF₄)₂·6H₂O (0.96 g) dissolved in methanol (10 mL). An immediate colour change from pale yellow to dark green was observed. The resulting solution was refluxed gently overnight. The solution was cooled and filtered and the solvent removed *in vacuo* to yield a dark green solid. The crude product was re-dissolved in the minimum amount of methanol, filtered through celite and recrystallised by vapour diffusion using diethyl ether to give a dark green powder.

cis-[Cu(II)S_{pr}N_{en}(-(CH₂)₂OH)₂] 2BF₄: Anal. Calcd. for C₂₃H₃₂ N₂S₂O₂CuB₂F₈: C, 39.30; H, 4.41; N, 4.36 % Found: C, 39.98; H, 4.72; N, 4.57 %. Mass Spec (ESI) m/z 495.08 [M]⁺, 451.13 [M-CH₂CH₂O]⁺. λ_{max} (solid reflectance) 435 nm, 640 nm.

cis-[Cu(II)S_{pr}N_{en}(-(CH₂)₃OH)₂] 2BF₄: Anal. Calcd. for C₂₅H₃₆ N₂S₂O₂CuB₂F₈: C, 43.04; H, 5.16; N, 4.02 % Found: C, 43.83; H, 5.46; N, 3.98 %. Mass Spec (ESI) m/z 523.20 [M]⁺, λ_{max} (solid reflectance) 435 nm, 600 nm. XRD data: Monoclinic C2/c, a = 35.7398(14) Å, b = 9.1810(3) Å, c = 21.0854(7) Å, β = 92.385(3)^o, V (Å³) 6912.7(4), Z = 8, r_{obs} = 0.0438, wR₂ = 0.1104.

cis-[Cu(II)S_{pr}N_{en}(-CH₂COOH)₂] 2BF₄: Anal. Calcd. for C₂₃H₂₈ N₂S₂O₄CuB₂F₈: C, 41.94; H, 4.44; N, 3.86 % Found: C, 42.13; H, 4.42; N, 4.02 %. Mass Spec (ESI) m/z 523.27 [M]⁺. λ_{max} (solid reflectance) 415 nm, 600 nm.

cis-[Cu(II)S_{pr}N_{en}(-(CH₂)₂SPh)₂] 2BF₄: Anal. Calcd. for C₃₅H₄₀ N₂S₄CuB₂F₈: C, 49.22; H, 4.68; N, 3.28 % Found: C, 49.67; H, 4.79; N, 3.11 %. Mass Spec (ESI) m/z 679 [M]⁺, 543 [M-C₉H₁₂S]⁺ λ_{max} (solid reflectance) 425 nm, 610 nm.

Preparation of $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{phen}}\text{R}]^{2+} 2\text{BF}_4^-$: 5-nitro-2-chloro-benzaldehyde (1.86 g, 0.01 mol) was dissolved in iso-propylalcohol to which o-phenylenediamine (0.6 g, 0.005 mol) was added. The solution was stirred at room temperature for an hour and a yellow precipitate formed. The solution was then filtered and the Schiff base, as a yellow solid, collected and dried.

^1H NMR (400 MHz, DMSO; δH) 8.99 (s, 2H, -CH=N-), 8.92 (d, 2H, arom), 8.29 (dd, 4H, arom), 7.87 (d, 2H, arom), 7.17 (d, 2H, arom), Anal. Calcd. for $\text{C}_{20}\text{H}_{12}\text{Cl}_2\text{N}_4\text{O}_4$: C, 54.20; H, 2.73; N, 12.64 % Found: C, 54.34; H, 2.60; N, 12.83 % Mass Spec. (ESI) m/z 443 $[\text{M}+\text{H}]^+$

The Schiff base (10 mmol) was dissolved in dichloromethane to which $\text{Cu}(\text{BF}_4)_2$ (0.34 g, 10 mmol) was added. The compound was heated to reflux and then propanedithiol (0.09g, 5 mmol) added drop-wise. The solution was subsequently refluxed for 48 hours. The solution was then filtered and the solvent removed *in vacuo* to yield a green-black oil. The crude product was dissolved in distilled water and extracted with dichloromethane and dried over anhydrous sodium sulphate. The solvent was removed to yield the copper Schiff base macrocycle as a green-black oil.

Anal. Calcd. for $\text{C}_{23}\text{H}_{18}\text{N}_4\text{O}_4\text{S}_2\text{CuB}_2\text{F}_8$: C, 38.60; H, 2.53; N, 7.83 % Found: C, 40.98; H, 2.60; N, 6.59 % Mass Spec. (ESI) m/z 541 $[\text{M}]^+$

The Schiff base macrocyclic complex (10 mmol) was dissolved in ethanol (100 ml) and an excess of NaBH_4 (1 g) was added portionwise to the solution and stirred

overnight. The excess NaBH₄ was quenched with ammonium acetate. The solvent was then removed *in vacuo* to yield a green oil. This was dissolved in distilled water and extracted with dichloromethane and dried over anhydrous sodium sulphate. The solvent was removed *in vacuo* to yield [Cu(II)S_{pr}N_{phen}R]²⁺ 2BF₄⁻ as a green solid.

Anal. Calcd. for C₂₃H₂₀N₄O₄S₂CuB₂F₈: C, 38.37; H, 4.42; N, 7.67 % Found: C, 39.03; H, 4.26; N, 7.59 %. Mass Spec. (ESI) m/z 543 [M]⁺. λ_{max} (solid reflectance) 425 nm, 610 nm.

Relaxation measurements: Relaxation measurements were carried out in triplicate on a Bruker Avance I NMR spectrometer equipped with an 11.7 T magnet. The measurements were carried out using an inversion recovery pulse sequence under full automation control with a 90° pulse width of 11.5 μs. All spectra were referenced internally to the residual proton resonance of the relevant deuterated solvent. The relaxation delay was varied from 1-100 s, depending on the relaxation characteristics of the sample. Data were transferred for remote data processing using Topspin (version 2.1, Bruker Biospin, Karlsruhe) on a desktop PC running under Windows XP. Following Fourier transformation, all data were baseline corrected. The T₁/T₂ sub-routine within Topspin was used to evaluate the relevant T₁ values from the data.

Samples of the copper macrocycles, GdDTPA and Cu(BF₄)₂ were prepared in 0.1 M NaCl, 100% D₂O in a 10 mL volumetric flask at concentrations of 10 mM, 2.5 mM, 1 mM, 0.1 mM and 0.01 mM. A control sample of 0.1 mM NaCl in 100% D₂O was also included.

Electrochemistry: Cyclic voltammetry in aqueous solution was carried out on a CH Instruments 660A Electrochemical Workstation with iR compensation, using ultra high purity water (Elga: 0.22 μ m filtered, 17 M Ω /cm) and acetonitrile (redistilled from calcium hydride). The electrodes were glassy carbon, platinum wire and silver wire as the working, counter and reference electrodes, respectively. All solutions were degassed with argon and contained sample concentrations 2×10^{-3} M for acetonitrile, together with the supporting electrolyte; tBu₄NBF₄ (0.1 M) respectively. Measurements are referenced against the redox couple ($E_{1/2}$) of ferrocene / ferrocenium.

Chemical redox cycling: Reduction: A solution of the copper macrocyclic (0.3 mM) complex and reducing agent (10 mM ascorbic acid) was prepared in Tris-buffer (5 mM pH 7.4). A 1.0 ml volume of the solution of the complexes was placed in a 1.5 ml quartz cuvette (1 cm pathlength) and the spectrum was recorded (250-800 nm). The solution was titrated with reducing agent in 5 μ l aliquots. The cuvette was allowed to stand for 5 minutes at room temperature to allow the solution to come to equilibrium before the spectrum was recorded. The titrations continued until no further change was observed. Each experiment was carried out in triplicate.

Re-oxidation: A solution of the copper macrocyclic complex (0.3 mM) was prepared in Tris-buffer (5 mM, pH 7.4) and reduced using ascorbic acid from the data above to generate the copper (I) complex *in-situ*. A 1.0 ml volume of the solution of the required copper complexes was placed in a 1.5 ml quartz cuvette (1 cm pathlength) and the spectrum was recorded (250 - 800 nm) against a Tris-buffer reference placed in a matched cuvette. A 100 mM solution of sodium hypochlorite was titrated in 5 μ l aliquots into the solution. The cuvette was allowed to stand for 5 minutes at room temperature to allow the solution to come to equilibrium before the spectrum was

recorded. The titrations continued until no further change was observed in the spectra. Each experiment was carried out in triplicate.

Results and Discussion

As a potential platform for the design of copper-based MRI moieties, we have synthesized and studied three related series of copper complexes (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al 2013). The first series (Figure. 1) comprise four thioether-secondary amine macrocycles where the alkyl chains linking the donor atoms are modified so as to allow the formation of 14, 15 or 16 membered rings. In all of these complexes the macrocycle lies in the meridial plane of the metal. A number of these complexes crystallise with a solvent molecule (MeCN, H₂O) coordinated to the copper atom in an axial position (Taylor et al. 2008; Trotter et al. 2009). This structural feature indicates that a relaxation mechanism similar to that found in Gd(DTPA) should be available with these copper macrocycles. A second series of macrocycles contain tertiary amines have been prepared. Here the pendant arms (X in Figure 1) include a terminal donor atom, which is introduced to increased the denticity of the ligand and the stability of the resulting copper complex. A third system has been synthesised that includes electron withdrawing groups (Figure 1, aryl and -NO₂) in an attempt to support the copper(I) form better.

-Figure 1 in here-

The secondary amine copper macrocycles (figure 1 left) were prepared as previously described (Sowden et al 2013) and their relaxation properties assessed in saline solution (0.10 M NaCl, ²H₂O, 9.4T). The performance of the species were

measured against Gd(DTPA) and $\text{Cu}(\text{BF}_4)_2$. Gd(DTPA) was included as a reference material to demonstrate the relative ability of these copper macrocycles to act as contrast agents. A simple hydrated copper salt ($\text{Cu}(\text{BF}_4)_2$) was also included in the study to gauge the maximum effect one might expect from copper agents and in turn the influence of the macrocyclic ligand. As expected, the relaxation measurements (Table 2) show that hydrated copper(II) ions have poor relaxivity compared to gadolinium (Laufer 1987). The hydrated copper(II) cation has a primary coordination sphere of six waters and a weakly bound secondary coordination sphere of up to eight water molecules (Persson 2010). Consequently, it is unsurprising that once the copper is placed within the macrocycles four of the exchange sites are occupied and the relaxivity falls.

- Table 2 in here-

It has been shown previously that bovine serum albumin (BSA) can efficiently remove the copper from these secondary amine macrocycles (Figure 1, top left) in competitive binding studies (Sowden et al 2013). Thus, although the copper macrocycles have the potential to act as contrast agents, in order to have any real utility the stability of these complexes needs to be increased. Earlier studies have shown that the $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$ motif was marginally more resistant to de-metalation by BSA and this framework was adopted for modification (Sowden et al 2013). Preliminary studies using a tertiary amine complex that incorporates ethanol pendant arms (Figure 1 top right, X = $(\text{CH}_2)_2\text{OH}$) indicated that this modification prevented BSA from removing the metal from the complex (Sowden et al 2013). Mattes et al. have previously synthesised a range of macrocyclic species based on $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{en}}]^{2+}$ and $[\text{Cu}(\text{II})\text{-S}_{\text{en}}\text{N}_{\text{pr}}]^{2+}$ that have

ethanol, acetate and pyridyl groups attached to the secondary amines (Funkemeier & Mattes 1993; Bentfeld et al. 1995; Mattes et al. 2004). In the solid state, these complexes are observed to be cationic, five coordinate species in which the pendant donor oxygen atoms are in the coordination sphere of the copper, lie *cis*- to one another and are found in their protonated form. The two donor oxygen atoms are included in the coordination geometry at the expense of one of the ring sulfur atoms. The $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$ platform has good potential and we have therefore adapted the work of Mattes et al. to introduce ethanol, propanol, acetate and arylthioether pendant arms into this motif (Figure. 1). Although all four complexes were isolated and characterised, only $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-\text{CH}_2)_3\text{OH})_2]^{2+}$ provided X-ray quality crystals. In contrast to the structures reported previously, this complex was found to adopt a five coordinate square based pyramidal structure (Figure 2). Although both alcohol groups are found above the plane of the ligand and are protonated, only one of the hydroxyl groups coordinates. The structures of the corresponding ethanol, acetate thioether adducts remain unclear at this time but we would predict that they are also five coordinate in the solid state (Funkemeier & Mattes 1993; Bentfeld et al. 1995; Mattes et al. 2004).

-Figure 2 in here-

The fifth donor atom was introduced into the ligand to increase the stability of the complexes. Typically the stability of a complex is framed using its formation constants, which are then used to hypothesise about the potential biochemistry. We have developed a more direct method of analysis whereby the copper complex is titrated with BSA (Sowden et al 2013). Using this approach, we circumvented the need to calculate each individual formation constant and directly observed if the

protein can sequester the metal (copper). Using this approach we can show that copper in the five coordinate species are stable to metal sequestration by BSA.[†] Although the introduction of a fifth donor into the motif has generated a suite of complexes that are suitably bio-tolerant, the increase in stability requires that a fifth water exchange site (c.f. $\text{Cu}(\text{BF}_4)_2$) is occupied and consequently the relaxivity of the complexes falls (Table 2).

The ability of the compounds to act as redox sensing contrast agents requires that they have redox reversibility between the copper(II) and copper(I) forms. The issue of redox reversibility has been interrogated in two ways: electrochemically and chemically. Electrochemical analysis of the compounds (Table 1) shows that the addition of the pendant arms has raised the E^0 of the complexes significantly. Surprisingly, the addition of electron withdrawing groups (i.e. $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{phen}}\text{R}]^{2+}$) and soft donor moieties ($[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}((-\text{CH}_2)_2\text{SPh})_2]^{2+}$) into the motif has little impact on redox potential of the copper centre and the values obtained for these species (1.2-1.5 V) consistently come to the higher end of the desired range (table 2). The voltammograms of compounds indicate that the addition of pendant arm donor moieties renders the complexes irreversible except in one case i.e. ($[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}((-\text{CH}_2)_2\text{SPh})_2]^{2+}$). This compound shows good reversibility at a range of scan rates (Figure 3.). This indicates that an increase in the sulphur density around the copper centre is having a beneficial effect and that the additional soft donors is able to support the formation of the copper(I) species during redox cycling better.

- Figure 3 in here-

-figure 4 in here-

In our previous studies we investigated the redox chemistry of our compounds by subjecting them to chemical redox cycling using ascorbic acid reduction followed by hypochlorite oxidation (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al 2013). Repeating this approach here, we found behaviour which paralleled the electrochemical study above. While all the compounds could be stoichiometrically reduced, only $[\text{Cu(II)}\text{S}_{\text{pr}}\text{N}_{\text{en}}((\text{CH}_2)_2\text{SPh})_2]^{2+}$ could be efficiently re-oxidised (~85%) back to the divalent state (Fig 4.). The ability to re-oxidise $[\text{Cu(II)}\text{S}_{\text{pr}}\text{N}_{\text{en}}((\text{CH}_2)_2\text{SPh})_2]^{2+}$ is a significant improvement over the previous compounds studied (table 1) (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al 2013). However, whereas reduction remains stoichiometric, re-oxidation still required an excess of oxidant to effect the redox switch.

Conclusions

Here we present the first major effort to explore copper's ability to be used as a redox-sensitive MRI moiety. As predicted by Lauffer (1987), the relaxivity of these copper based species is poor compared with gadolinium and considerable further development is needed before copper will be able to perform independently as a redox sensitive contrast agent. The study does indicate that it should be possible to generate a compound which is stable in both the +1 and +2 oxidation states under biological conditions with an appropriate oxidation potential. However, it is clear that for these compounds to have any use in MRI they will have to be coupled with a higher relaxivity metal such as gadolinium. Finally, it would be better if the sensing agent was administered in its low valent form and the challenge now is to produce copper(I) complexes of this type that respond to oxidants rather than copper(II) complexes which respond to reductants.

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Electronic Supplementary Information (ESI) available:

Details of the X-ray crystal structure determination of $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2] \cdot 2\text{BF}_4$ may be obtained from the CCDC; e-mail deposit@ccdc.cam.ac.uk or [www:http://ccdc.cam.ac.uk](http://www.ccdc.cam.ac.uk)) on request quoting the depository numbers [ccdc 953160](https://doi.org/10.1039/C9CC00095G). †The supporting information contains details on the competitive binding studies of the macrocycles with BSA.

Notes and references

It should be noted that Kupka et al. (1992) made a brief comment on the potential use of copper(II) D-penicillamine as a contrast agent in plants. To our knowledge this work did not develop further.

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Table Captions

Table 1. The properties of gadolinium and certain first row transition metal species key to their ability to perform as contrast agents (Tobe 1972, Handbook of Chemistry 1980). Unlike the lanthanoids, the exchange kinetics and number of unpaired electrons vary more widely and are governed by the prevailing oxidation state and ligand field considerations. Oh denotes an octahedral geometry

Table 2. Relaxivity and electrochemical data: The relaxivity of GdDTPA, $\text{Cu}(\text{BF}_4)_2$, $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{en}}]^{2+}$; $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$; $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{pr}}]^{2+}$; $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{pr}}]^{2+}$; $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(\text{X})_2]^{2+}$, where $\text{X} = -(\text{CH}_2)_2\text{OH}$, $-(\text{CH}_2)_3\text{OH}$, $-\text{CH}_2\text{COOH}$, $-(\text{CH}_2)_2\text{SPh}$ and $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{phen}}\text{R}]^{2+}$ ($n = 3$) in 0.1 M NaCl in $^2\text{H}_2\text{O}$ at 9.4T. . Electrochemical data for the copper macrocycles in acetonitrile (0.1M $^t\text{Bu}_4\text{NBF}_4$: The values for the complexes are expressed relative to the normal hydrogen electrode (HNE) (Taylor et al. 2008)

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Figure 1 The copper macrocyclic frameworks studied (Taylor et al. 2008; Trotter et al. 2009; Trotter et al. 2010; Sowden et al. 2013). (a) Secondary amine complexes, where the polymethylene chains between the nitrogen and sulfur donor atoms are ethylene (en) or propylene (pr) giving rise to four compounds: $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{en}}]^{2+}$, $[\text{Cu}(\text{II})\text{S}_{\text{en}}\text{N}_{\text{pr}}]^{2+}$, $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$ and $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{pr}}]^{2+}$. (b) Tertiary amine complexes $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(\text{X})_2]^{2+}$, where $\text{X} = -(\text{CH}_2)_2\text{OH}$, $-(\text{CH}_2)_3\text{OH}$, $-\text{CH}_2\text{COOH}$ or $-(\text{CH}_2)_2\text{SPh}$. (c) $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{phen}}\text{R}]^{2+}$ where additional electron withdrawing groups (aryl, $-\text{NO}_2$) are included in the motif.

Figure 2. The X-ray crystal structure of $[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2]^{2+}$. The counter ions (BF_4^-) are omitted for clarity. The copper sits in a distorted square based pyramidal geometry. The hydrogens on the oxygens were found during refinement. The thermal ellipsoids are shown at 50% probability. Selected metrical parameters (d/Å, °) Cu-S 2.3450(7), 2.3487(7); Cu-N 2.045(2), 2.052(2); Cu-O 2.1725(18); $\angle\text{S-Cu-N}$ (°), 163.00(6), 93.48(6), 176.82(6), 93.49(6).

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Table 1

	No. unpaired electrons and relevant oxidation state	Nuclear spin (abundance %)		Exchange rate (s ⁻¹)
Gd	7	3/2 ¹⁵⁵ Gd	(14.7)	10 ⁹
		3/2 ¹⁵⁷ Gd	(15.7)	
V	2 (III)	7/2 ⁵¹ V	(99.8)	10 ³
Cr	4 (II)	3/2 ⁵³ Cr	(9.6)	10 ⁹
	3 (III)			5 x 10 ⁻⁵
Mn	7 (II)	5/2 ⁵⁵ Mn	(100)	5 x 10 ⁷
Co	3 (II) high spin O _h	7/2 ⁵⁹ Co	(100)	5 x 10 ⁶
	0 (III) low spin O _h			5 x 10 ⁻⁵
Cu	1 (II)	3/2 ⁶³ Cu	(69.1)	10 ⁹
	0 (I)	3/2 ⁶⁵ Cu	(30.9)	

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Table 2

	Compound	Relaxivity $s^{-1}mM^{-1}$	E^0 V
Reference compounds	Gd(DTPA)	4.73	
	$Cu^{II}(BF_4)_2$	0.67	
Simple macrocycles (Fig 1a)	$[Cu(II)S_{en}N_{pr}]^{2+}$,	0.27 ± 0.03	0.65^{*c}
	$[Cu(II)S_{pr}N_{en}]^{2+}$	0.27 ± 0.01	0.85 (57%)
	$[Cu(II)S_{pr}N_{pr}]^{2+}$	0.26 ± 0.03	0.85^{*c}
Pendant arm macrocycles $[Cu(II)S_{pr}N_{en}(X)_2]^{2+}$ (Fig 1b)	X= $-(CH_2)_2OH$	0.42 ± 0.02	1.04 (22%)
	X= $-(CH_2)_3OH$	0.16 ± 0.02	1.23 (21%)
	X= $-CH_2COOH$	0.11 ± 0.01	1.44 (24%)
	X= $-(CH_2)_2SPh$	0.12 ± 0.02	1.28 (79%)
Macrocycle-EWG (Fig 1c)	$[Cu(II)S_{pr}N_{phen}R]^{2+}$	0.11 ± 0.02	1.42 (9%)

Relaxivity and electrochemical data: The relaxivity of GdDTPA, $Cu(BF_4)_2$, $[Cu(II)S_{en}N_{en}]^{2+}$; $[Cu(II)S_{pr}N_{en}]^{2+}$; $[Cu(II)S_{en}N_{pr}]^{2+}$; $[Cu(II)S_{pr}N_{pr}]^{2+}$; $[Cu(II)S_{pr}N_{en}(X)_2]^{2+}$, where X = $-(CH_2)_2OH$, $-(CH_2)_3OH$, $-CH_2COOH$, $-(CH_2)_2SPh$ and $[Cu(II)S_{pr}N_{phen}R]^{2+}$ (n = 3) in 0.1 M NaCl in 2H_2O at 9.4T. . Electrochemical data for the copper macrocycles in acetonitrile (0.1M tBu_4NBF_4 : The values for the complexes are expressed relative to the normal hydrogen electrode (HNE). * (Taylor 2008)

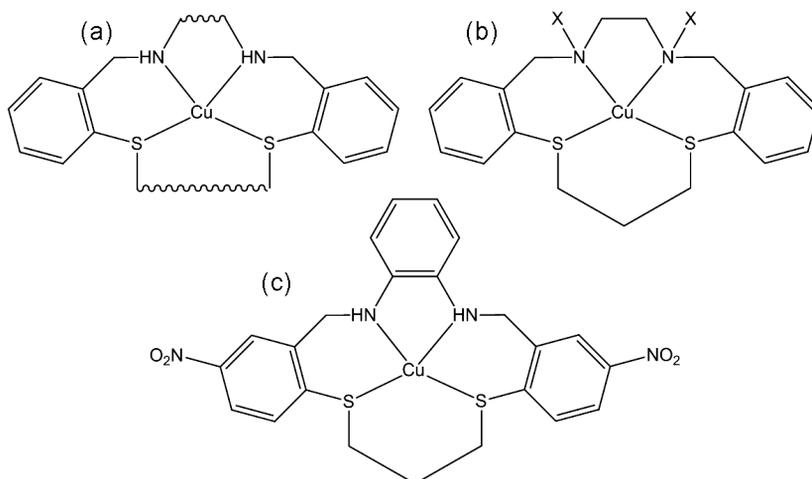


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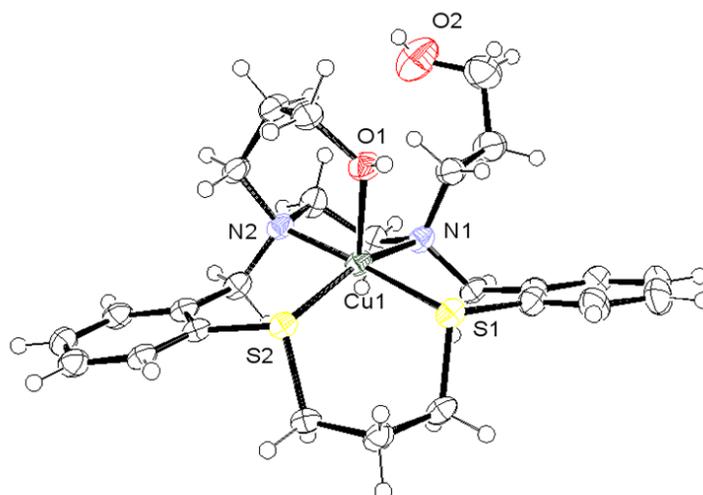


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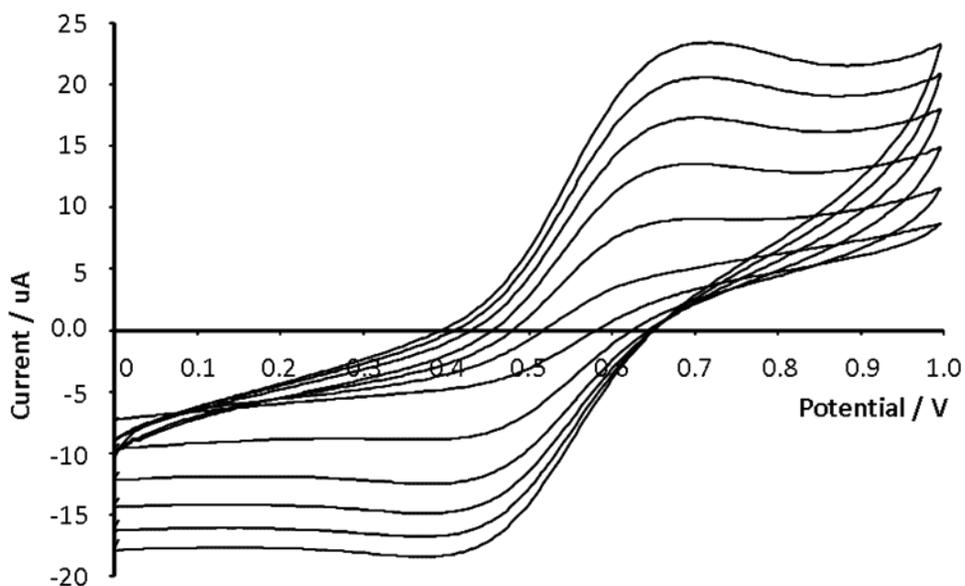


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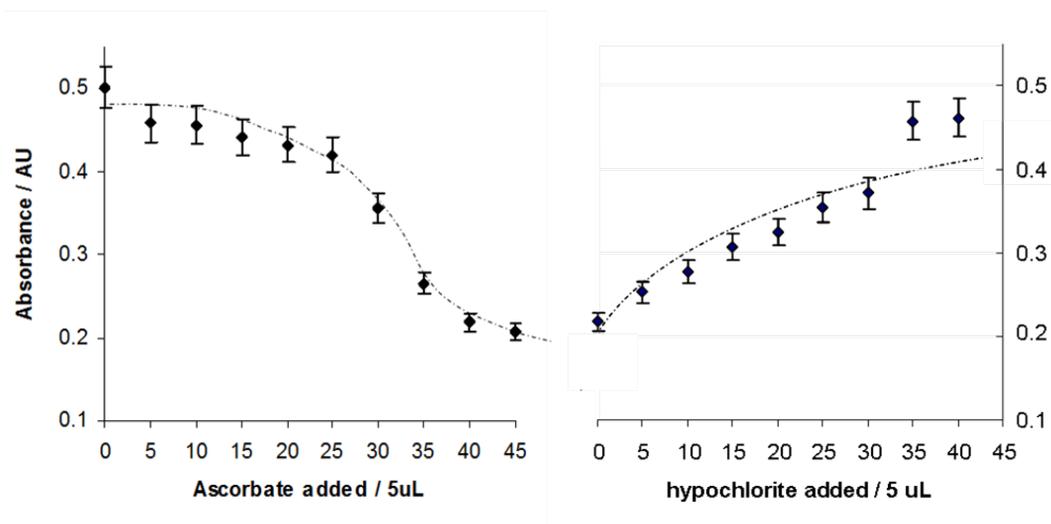


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Copper complexes as a source of redox active MRI contrast agents.

Lynsey Dunbar, Rebecca J. Sowden, Katherine D. Trotter, Michelle K. Taylor, David Smith, Alan R. Kennedy, John Reglinski and Corinne M. Spickett

Supporting Information

Competitive binding studies: Rather than measure the formation constants to arrive at a value which has limited biological impact, we elected to challenge our copper complexes directly with bovine serum albumin (BSA). This simple experiment is designed to probe stability of our copper complexes directly in competition with BSA, which is a major metal scavenger present in the blood stream. This simple, direct experiment more closely models the sequestration of metals *in-vivo*.

Reaction of the copper(II) complexes with bovine serum albumin (BSA): 0.3 mM solutions of each of the copper macrocycles and BSA (199 mg, 1 mL) were prepared in Tris-HCl buffer (5 mM, pH 7.4). A 1.0 mL volume of the solution of each of the required copper complexes was placed in a 1.5 mL quartz cuvette (1.0 cm path length) and the spectrum was recorded between 200 and 800 nm against a Tris-buffer reference placed in a matched cuvette. BSA solution was titrated in aliquots of 20 μ L into the cuvette.

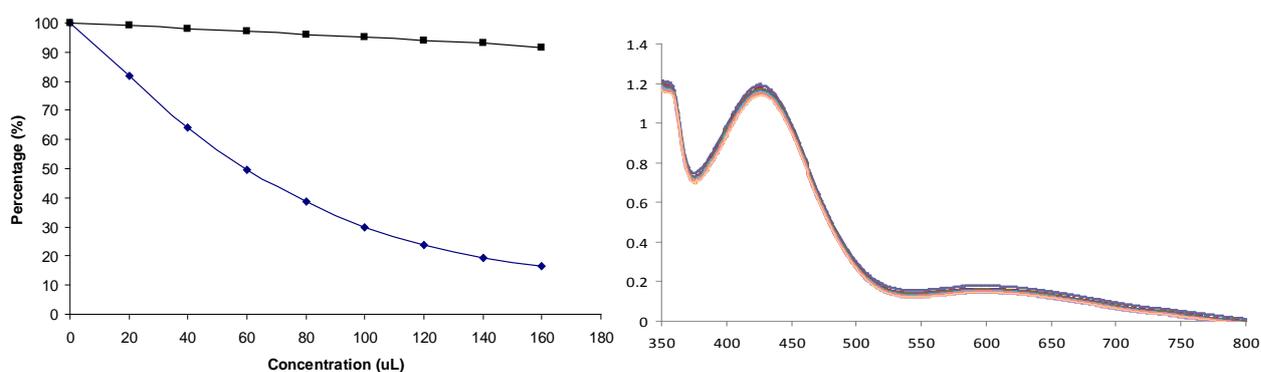


Figure ESI1: Left: The spectrophotometric titrations (412 nm) of $\text{cis}[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}]^{2+}$ (adapted from Sowden et al.) and $\text{cis}[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2]^{2+}$. Right: The spectrophotometric titrations (350-800) of $\text{cis}[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2]^{2+}$: (0.3 mM in 5 mM Tris buffer at pH 7.4) with BSA solution (199 mg, 1 mL). A small diminution in the spectrum of $\text{cis}[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2]^{2+}$ of the absorbance occurs owing to dilution of the sample as the titrant is added. The integrity of the complex ($\text{cis}[\text{Cu}(\text{II})\text{S}_{\text{pr}}\text{N}_{\text{en}}(-(\text{CH}_2)_3\text{OH})_2]^{2+}$) is maintained throughout and the copper is not being sequestered from the ligand cavity.

Sowden RJ et al (2013) *Biometals* 26: 85-95.