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New Water-Soluble Copper(II) Complexes with Morpholine-Thiosemicarbazone Hybrids: Insights into the Anticancer and Antibacterial Mode of Action

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Abstract

Six morpholine-(iso)thiosemicarbazone hybrids **HL**¹–**HL**⁶ and their Cu(II) complexes with good-to-moderate solubility and stability in water were synthesized and characterized. Cu(II) complexes [Cu(**L**^{1–6})Cl] (**1**–**6**) formed weak dimeric associates in the solid state, which did not remain intact in solution as evidenced by ESI-MS. The lead proligands and Cu(II) complexes displayed higher antiproliferative activity in cancer cells than Triapine. In addition, complexes **2**–**5** were found to specifically inhibit the growth of Gram-positive bacteria *S. aureus* with MIC₅₀ values at 2 to 5 µg/mL. Insights into the processes controlling intracellular accumulation and mechanism of action were investigated for **2** and **5**, including the role of ribonucleotide reductase (RNR) inhibition, endoplasmic reticulum (ER) stress induction and regulation of other cancer signaling pathways. Their ability to moderately inhibit R2 RNR protein in the presence of DTT is likely related to Fe chelating properties of the proligands liberated upon reduction.

Introduction

Progress in modern anticancer therapy has led to significant improvements in survival rates of cancer patients. However, even though more patients achieve remission nowadays, their life-span often remains short.¹ One of the major causes of cancer death is the malignant process itself, which is associated with extensive formation of metastases, but it is not commonly mentioned that a significant number of immunocompromised cancer patients die due to infections, such as pneumonia and peritonitis.² Despite that cancer patients are very prone to develop infections during chemotherapy, their preventive antibiotic treatment is hampered by additional adverse effects.³ However, recent clinical evidence demonstrated that benefits of antibiotic prophylaxis of cancer patients outweighed its risks.⁴ The simultaneous suppression of pathogenic microorganisms during anticancer chemotherapy could not only interrupt or abolish tumor growth but eventually protect cancer patients from infection. Hence, the development of novel drugs which exhibit dual anticancer and antibacterial properties in comparable concentration range would affect the malignant process and simultaneously decrease the risk of patients' death due to infection, febrile neutropenia and bacteraemia.

Both cancer and bacterial cells share similar properties, such as high rate of proliferation, rapid adjustment and quick spreading within the host and aggressive disease progression.⁵ In order to sustain such rapid proliferation, cancer and bacterial cells have to support DNA replication and production of RNA by increasing *de novo* nucleotide synthesis. Ribonucleotide reductase (RNR) is the key enzyme that catalyzes the reduction of ribonucleotides to their corresponding deoxyribonucleotides, thereby initiating DNA synthesis or repair, making it an important biomolecular target for drugs with anticancer and antibacterial properties.^{6,7} RNR consists of a large subunit (NrdA or R1), which contains the allosteric site that regulates and catalyzes substrate reduction, and a small subunit (NrdB or R2) with the diferric-tyrosyl radical cofactor, essential for the catalytic activity of the RNR enzyme. Small molecules,

that can sequester Fe(III) from the dinuclear metal center and/or scavenge the tyrosyl radical, inhibit R2 activity, thereby preventing *de novo* DNA synthesis in cancer cells and bacteria.

α -N-heterocyclic thiosemicarbazones (TSCs) are excellent transition metal chelators and have a broad range of activities, including anticancer and antibacterial properties, which are believed to be at least partially due to their RNR inhibition.⁸ To date, several TSC compounds, namely 3-amino-2-pyridinecarboxaldehyde thiosemicarbazone (Triapine),^{9,10,11} di-2-pyridylketone 4-cyclohexyl-4-methyl-3-thiosemicarbazone (DpC)^{12,13} and (*E*)-*N'*-(6,7-dihydroquinolin-8(5*H*)-ylidene)-4-(pyridine-2-yl)piperazine-1-carbothiohydrazide (COTI-2)¹⁴ are undergoing phase I and II clinical trials against various types of cancer. The main caveat in the use of TSCs is their high toxicity which is reflected in a number of side-effects in clinical trials involving Triapine.^{9,10,11} Many studies have shown that Cu(II)-TSC complexes often display better selectivity than their TSC proligands, which can be attributed to the induction of different intracellular signaling pathways.¹⁵

Despite advances in the design and synthesis of new TSCs and Cu(II)-TSC complexes over the years, these compounds are facing the problem of low aqueous solubility, thereby hampering their further development. Therefore, the synthesis of water-soluble and cytotoxic TSCs and Cu(II)-TSC complexes requires a thoughtful selection of the functional groups that can be attached at the TSC backbone without reducing their biological activity.^{16,17,18,19,20} With the aim to reach the optimal combination of aqueous solubility and high cytotoxicity, we designed new (iso)thiosemicarbazone-morpholine hybrids and their Cu(II) complexes. Notably, morpholine moiety was chosen since it confers excellent water solubility, which typically translates into an improved pharmacological effect. Additionally, morpholine derivatives possess a broad spectrum of biological activities, including anticancer and antibacterial therapeutic potential.²¹ For example, commercial anticancer drugs Aprepitant and Gefinitib, as well as antibacterial drugs Finafloxacin and Levofloxacin include a morpholine fragment in their structures.

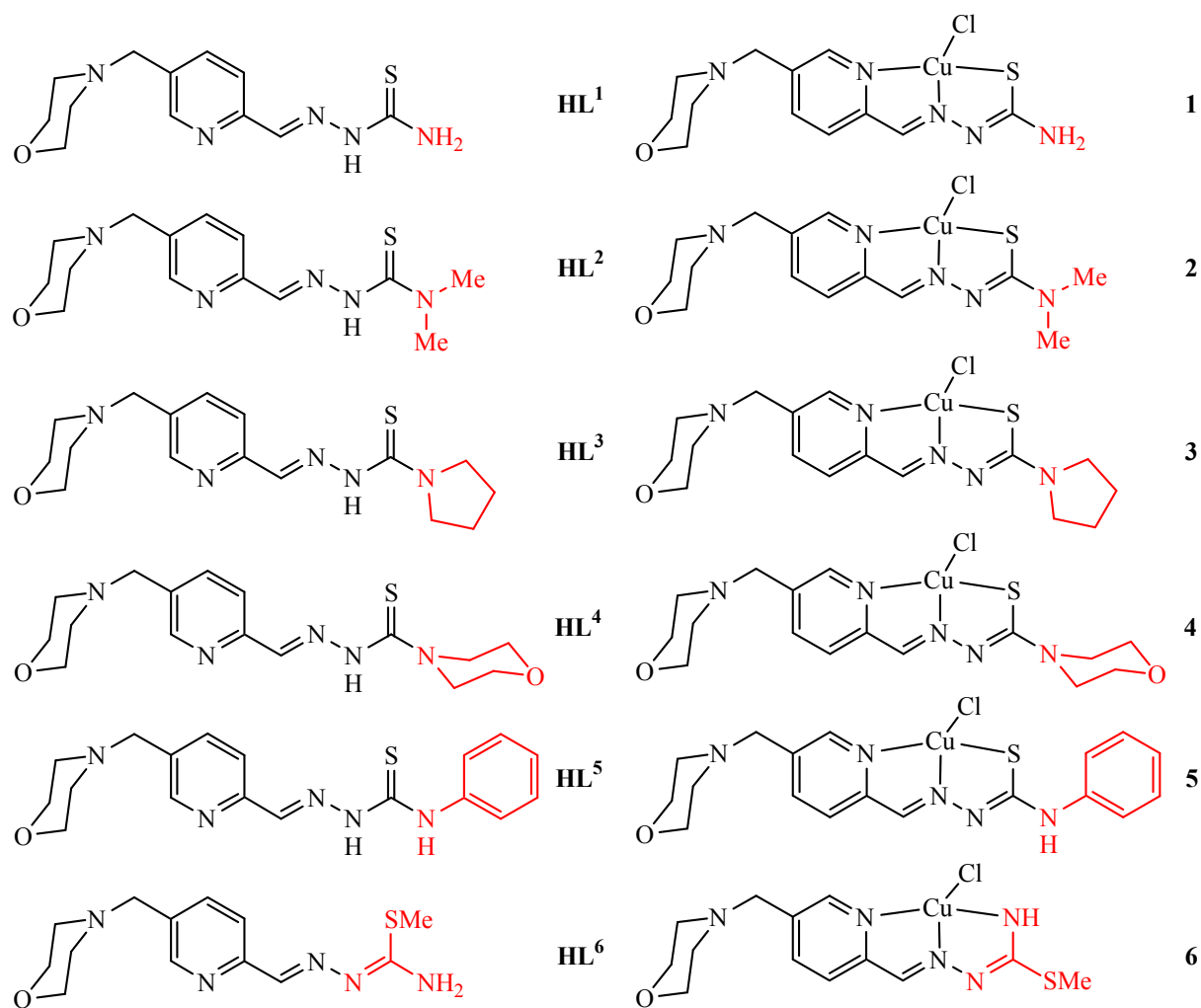


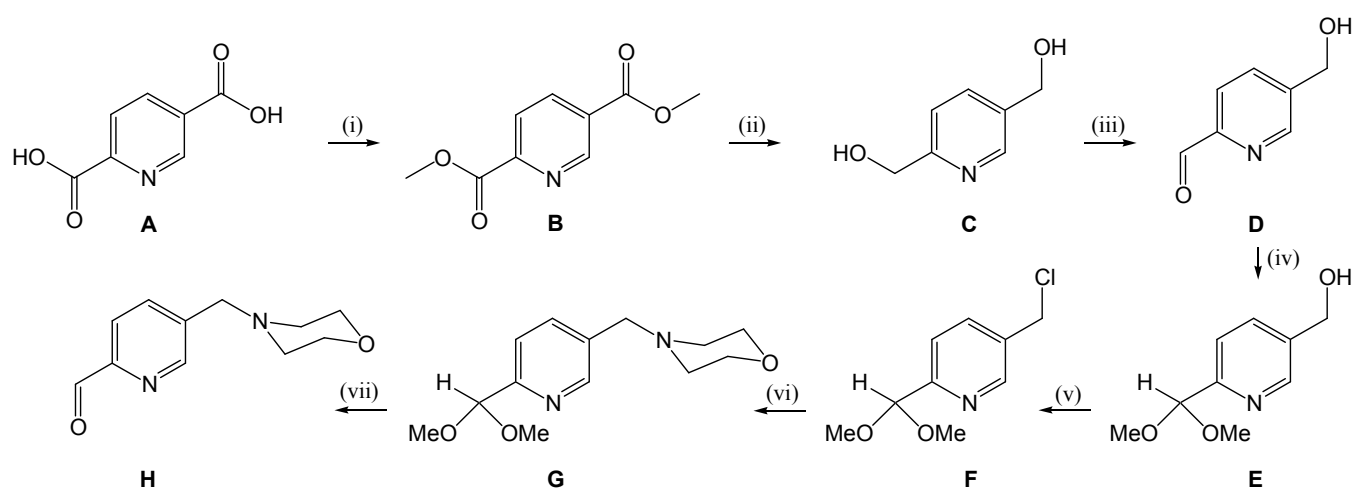
Chart 1. Line drawings of proligands **HL**^{1–6} and Cu(II) complexes [**Cu**(**L**^{1–6})**Cl**] (**1–6**) reported in this work. All proligands and Cu(II) complexes, but **HL**³ and **2**, were investigated by single crystal X-ray crystallography.

Herein we report on the synthesis of water-soluble proligands **HL**^{1–6} and Cu(II) complexes **1–6** (Chart 1), which were characterized by analytical and various spectroscopic techniques. X-ray diffraction structures of **HL**¹, **HL**², **HL**³–**HL**⁶ and **1**, **3–6** were established and solution equilibria studies for **HL**¹ and Cu(II) complex formation with **HL**¹ by pH-potentiometry, UV–vis, EPR and ¹H NMR spectroscopy, as well as electrochemistry were performed. Antiproliferative activity against human ovarian cancer A2780 and cisplatin resistant A2780cis cell lines, non-cancerous HEK293 cell line, as well as antibacterial activity against Gram-positive (*S. aureus*) compared to Gram-negative bacteria (*P. aeruginosa*) were investigated and structure-activity relationships discussed. The processes controlling cellular accumulation of **2** were investigated. Preliminary insights into the mode of action of **2** and **5**,

including mouse R2 RNR inhibitory potential, studied by molecular modeling and tyrosyl radical quenching monitored by EPR spectroscopy, as well as by Western Blotting and flow cytometry are also presented.

Results

Synthesis and characterization of (iso)thiosemicarbazone-morpholine hybrids and their Cu(II) complexes. The 5-methylmorpholine-pyridine-2-carboxaldehyde **H** was prepared in seven steps as shown in Scheme 1 and the detailed synthesis is described in the Supporting Information. First pyridine-2,5-dicarboxylic acid **A** was converted into diester **B** that was further reduced to the diol **C**. The latter was selectively oxidized with SeO₂ to the aldehyde **D**. After protection of the aldehyde group the alcohol **E** was converted into the chloride **F** that was further reacted with morpholine to give species **G**. Finally, the hydrolysis of the methyl ester function in acidic conditions afforded the required aldehyde **H** for condensation reactions with thiosemicarbazide, 4*N*-dimethyl-3-thiosemicarbazide, 4*N*-pyrrolydiny-3-thiosemicarbazide, 4*N*-morpholinyl-3-thiosemicarbazide, 4*N*-phenyl-3-thiosemicarbazide and *S*-methylisothiosemicarbazide hydroiodide to give the hybrids **HL**¹⁻⁶ (Scheme 1), respectively. One- and two-dimensional NMR spectra were in agreement with the proposed structures for **HL**¹⁻⁶, enabling the assignment of all ¹H and ¹³C resonances. The ESI mass spectra recorded in a positive ion mode showed strong peaks corresponding to [M+H]⁺ and [M+Na]⁺ ions, respectively. The proligands **HL**¹⁻⁶ were reacted with Cu(II) chloride dihydrate and triethylamine in 1:1:1 mole ratio in methanol to give [Cu(L¹⁻⁶)Cl] (**1-6**) in 35–91% yields. Positive ion ESI mass spectra showed strong peaks attributed to [Cu(L¹⁻⁶)]⁺ ions. The structures of **1**, **3-6** were also established by single crystal X-ray diffraction (*vide infra*).



Scheme 1. Synthesis of 5-methylmorpholine-pyridine-2-carboxaldehyde. Reagents and conditions: (i) thionyl chloride, methanol, 0 °C → room temperature, 12 h;²² (ii) sodium borohydride, ethanol, acetone, potassium carbonate, chloroform, 0 °C, 1 h → reflux, overnight;²³ (iii) selenium dioxide, dioxane, water, 100 °C, 3 h;²⁴ (iv) trimethyl orthoformate, methanesulfonic acid, methanol, reflux, 48 h; (v) thionyl chloride, dichloromethane, −80 °C → room temperature, overnight; (vi) morpholine, triethylamine, THF/CH₂Cl₂ 1:1, 50 °C, overnight, purification by column chromatography; (vii) HCl, water, 60 °C, overnight.

X-ray Crystallography. The results of X-ray diffraction studies of **HL**¹, **HL**² and **HL**⁴⁻⁶ are shown in Figure S1, while those of [Cu(**L**¹)Cl] (**1**), [Cu(**L**³)Cl] (**3**), [Cu(**L**⁴)Cl] (**4**), [Cu(**L**⁵)Cl] (**5**) and [Cu(**L**¹)Cl(H₂O)] (**1'**), [Cu(**L**⁶)Cl] (**6**) in Figures 1 and 2, respectively. Selected bond distances and bond angles are quoted in Tables S1 and S2. The proligands adopted different isomeric configurations in the solid state depending on substituents at the terminal nitrogen atom of the thiosemicarbazide moiety. Complexes **1**, **3**, **4** and **5** form dimeric associates as shown in Figure 1 (co-crystallized solvent was omitted for clarity). Each Cu(II) ion has a distorted square-planar coordination geometry. Intermolecular contacts supporting the dimeric associates in the crystals are of different nature in **1** and **3–5**, respectively. The presence of long intermolecular contacts Cu⋯Cl or Cu⋯Cl and Cu⋯S (see legend to Figure 1) provides evidence of weak association of complexes in dimers, which most probably dissociate in solution with formation of monomeric species. There was no evidence from ESI mass spectra on the presence of dimeric species in solution.

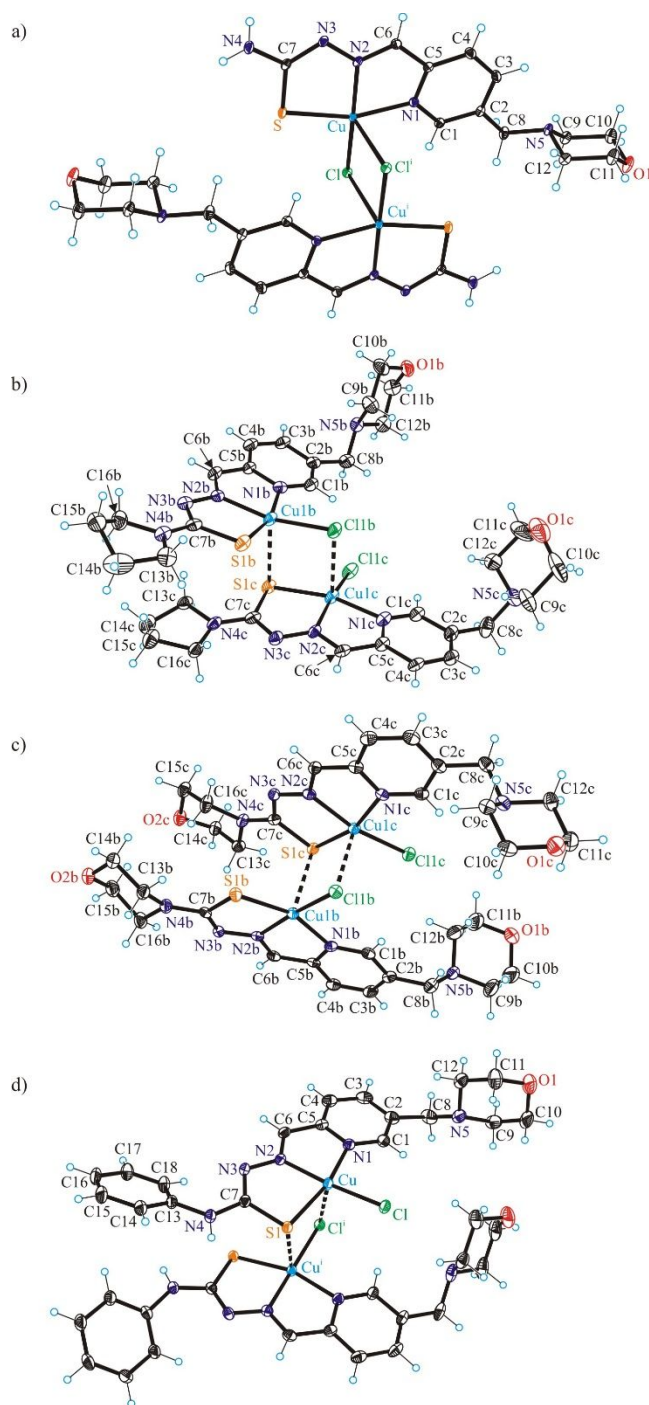


Figure 1. ORTEP views of weak dimeric associates of (a) [Cu(L¹)Cl] (**1**), (b), [Cu(L³)Cl] (**3**), (c) [Cu(L⁴)Cl]₂ (**4**) and (d) [Cu(L⁵)Cl] (**5**).

Moreover, **HL**¹ reacts with CuCl₂ in aqueous solution with formation of five-coordinate complex [Cu(L¹)Cl(H₂O)] (**1'**·2H₂O) (Figure 2b). The monomeric square-planar complex [Cu(L⁶)Cl] forms an

infinite chain via weak coordination of the morpholine oxygen atom of one complex to the Cu(II) atom of the next molecule as shown in Figure S2. The intermolecular Cu \cdots O contact is of 2.603(3) Å.

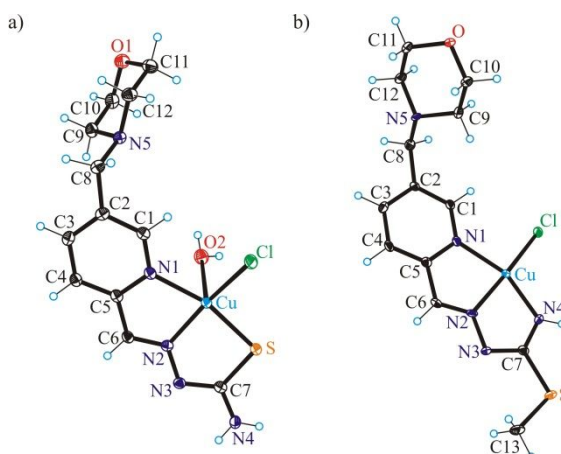


Figure 2. ORTEP views of (a) [Cu(L¹)Cl(H₂O)] (**1**⁺) and (b) [Cu(L⁶)Cl] (**6**).

Solution Chemistry. In order to establish the presence of isomers of the proligands in aqueous solution and proton dissociation processes, in which these can be involved solution equilibrium studies have been performed. Likewise, the solution speciation of the copper(II) complexes, as well as their stability have been investigated to elucidate the species, which is the most stable and abundant at physiological pH. The proligand **HL**¹ (Chart 1) was chosen for the detailed solution equilibrium studies since it has the simplest structure and the best aqueous solubility among the proligands prepared. The presence of different isomers in solution was excluded by measurements of ¹H NMR spectra in 10% D₂O/90% H₂O (Figure S3), which showed only one set of signals for the proligand in accordance to its low (*C*₁) molecular symmetry. The proligand most probably adopts the *E* configuration found in the solid state (Figure S1). The same configuration was reported previously for the reference compound 2-formylpyridine thiosemicarbazone in polar solvents.²⁵ Proton dissociation processes were monitored in aqueous solution by pH-potentiometric and ¹H NMR titrations and three p*K*_a values were determined by both methods (Table S3). According to the obtained p*K*_a values **HL**¹ is mainly neutral (97% HL, 3% H₂L⁺) at physiological pH (Figure S4). Inspection of ¹H NMR spectra revealed stepwise deprotonation

of three functional groups in the following order: $N_{\text{pyridinium}}H^+ \rightarrow N_{\text{morpholinium}}H^+ \rightarrow N_{\text{hydrazine}}H$ (Figure S3).

The solution speciation of the Cu(II) complexes with **HL**¹ was characterized by the combined use of pH-potentiometry, UV-vis spectrophotometry (via charge transfer (CT) and d-d bands) and EPR spectroscopy. The spectral changes (Table S4, Figures 3a and S5) in the UV and visible regions measured at 1:1 metal-to-ligand ratio show the high-extent formation of a Cu(II) complex already at strongly acidic pH values (e.g. pH 1) and its stepwise deprotonation by increasing the pH. At pH 7.4 the dominant species is $[CuL]^+$ (Figure 3b).

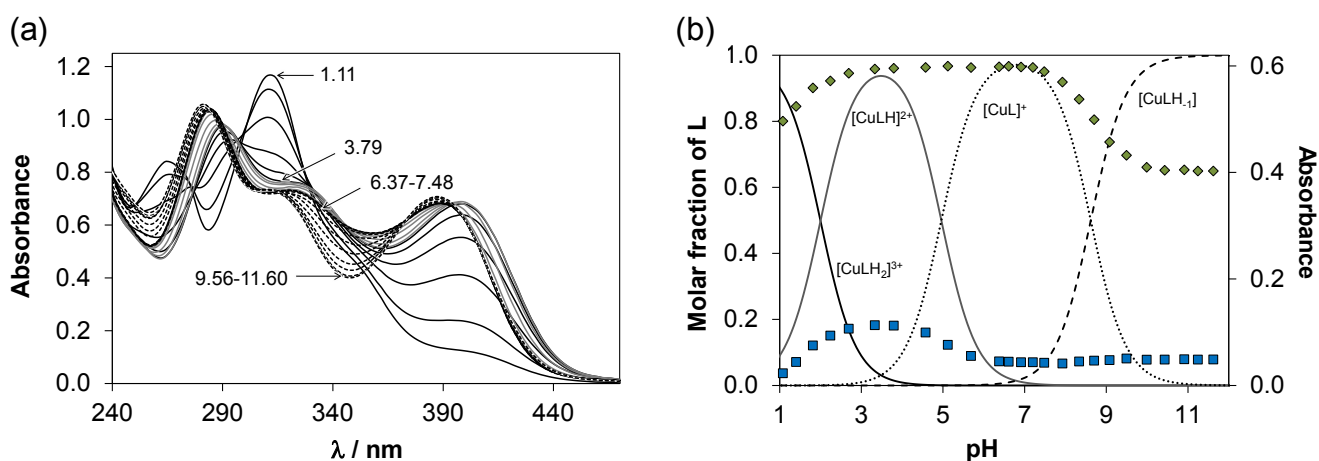


Figure 3. UV-vis spectra recorded for the Cu(II)-**HL**¹ (1:1) system at various pH values (a), and concentration distribution curves and measured absorbance values at 346 nm (♦) and 440 nm (■) for the same system (b). $\{c_{Cu(II)} = c_{lig} = 121 \mu M; T = 298 K; I = 0.10 M (KCl); l = 0.5 cm\}$.

Due to the high stability of the Cu(II) complexes formed, the cumulative constant ($\log\beta$) for the $[CuL]^+$ species (Table S4) was determined via EDTA displacement studies (Figure S6). Then the pK_a and the $\log\beta$ values for the other type of complexes ($[CuLH_2]^{3+}$, $[CuLH]^{2+}$, $[CuLH_{-1}]$) were computed using each method and data were in a fairly good agreement (Table S4). It is worth noting that formation of a bis-ligand complex $[CuL_2]$ at ligand excess was also confirmed by the UV-vis CT and EPR titrations. The

d-d bands (Figure S5) recorded from pH 3.4 to 7.5 at 1:1 metal-to-ligand ratio revealed that the proton dissociation process of $[\text{CuLH}]^{2+} \rightarrow [\text{CuL}]^+$ is accompanied by rather weak spectral changes as most probably the non-coordinating morpholinium NH^+ is deprotonating. In order to further characterize the coordination modes of the various complexes in solution EPR parameters (Table S5) obtained by the deconvolution of the recorded spectra (Figure S7) were analyzed. The isotropic g and A values of $[\text{CuLH}]^{2+}$ and $[\text{CuL}]^+$ are quite similar indicating the like coordination mode.

Comparison of the EPR parameters of the Cu(II) complexes of **HL**¹ with those of 2-formylpyridine thiosemicarbazone²⁵ and Triapine²⁶ permitted to conclude that in the $[\text{CuLH}]^{2+}$ and $[\text{CuL}]^+$ complexes a typical coordination mode via the $\text{N}_{\text{pyridyl}}, \text{N}, \text{S}^-$ binding site is realized in accordance with the X-ray diffraction data. $[\text{CuLH}_2]^{3+}$ contains the diprotonated ligand in which the protons are attributed to the non-coordinating N^2H and morpholinium NH^+ moieties, while $[\text{CuLH}_1]$ is most probably a mixed hydroxido complex $[\text{CuL}(\text{OH})]$ formed by the deprotonation of water molecule coordinated in the fourth equatorial position. At ligand excess besides the bis-ligand complex a minor dinuclear species $[\text{Cu}_2\text{L}_3\text{H}]^{2+}$ was detected resulting in the weak exchange coupling between neighboring Cu(II) centers smearing out the expected hyperfine structure.

Cu(II)-TSC complexes undergo quasi-reversible one-electron reduction at biologically accessible potentials. In order to assess the redox properties of **1–6**, detailed electrochemical and spectroscopic studies in various solvents were performed using cyclic voltammetry as well as EPR–spectroelectrochemistry. Additionally, the UV–vis spectra of **1–5** were measured (Figure S8). Electrochemical and spectroscopic data are summarized in Table S6. Almost reversible one-electron cathodic reduction was observed for **1–5** in DMSO with half-wave redox potentials from -0.77 V to -0.81 V vs Fc^+/Fc (for comparison from -0.13 V to -0.17 V vs NHE) (Table S6 and Figure S9). In general, the values of redox potentials of **1–6** for the first reduction step decreased in the following rank

order ($E_{1/2}(\mathbf{5}) > E_{1/2}(\mathbf{2}) = E_{1/2}(\mathbf{3}) = E_{1/2}(\mathbf{4}) > E_{1/2}(\mathbf{1}) > E_{1/2}(\mathbf{6})$). The second reduction step occurred at -1.8 V vs Fc^+/Fc (-1.16 V vs NHE) and was less reversible indicating a ligand based reduction. This was confirmed by the CVs of the corresponding proligands which exhibited the first reduction step at around -1.9 V vs Fc^+/Fc (not shown). In aqueous solutions the reduction potentials shifted to the less negative values and reduction was less reversible as shown for **2** in Figure S10. Complex **6** exhibited different redox behavior with the lowest electrochemical reversibility and the most negative reduction potential of -0.86 V vs Fc^+/Fc (-0.26 V vs NHE) (Figure S9c). Unlike the cathodic reduction, the anodic oxidation of **1–6** was irreversible with potential values in the region from 0.4 to 0.8 V vs Fc^+/Fc (0.68 V to 1.44 V vs NHE). Similar response was observed for the corresponding proligand **HL**² (Figure S11), indicating ligand based oxidation in **2**.

The biologically accessible reduction of Cu(II)-TSC complexes is Cu-centered. To investigate whether the biologically accessible reduction is metal-centered, the reversible one-electron reduction of **5** was further studied by *in situ* UV–vis-spectroelectrochemistry (Figure 4a). Upon cathodic reduction at the first reduction peak two isosbestic points at 389 and 325 nm were detected. The spectral changes of the $\text{S} \rightarrow \text{Cu(II)}$ charge transfer bands (~ 425 nm) clearly confirmed the reduction of Cu(II) to Cu(I) . Additionally, upon voltammetric reverse scan, reoxidation and a nearly full recovery of the initial optical bands were observed, attesting the chemical reversibility of the cathodic reduction even at low scan rates (Figure 4). To confirm the involvement of Cu(II) in the reduction processes, *in situ* EPR electrochemistry of **1–6** in $n\text{Bu}_4\text{NPF}_6/\text{DMSO}$ and water was performed, since metal-based reduction would result in the formation of EPR-silent Cu(I) species (Figure 4b). As can be seen for electrochemical reduction of **2** in $n\text{Bu}_4\text{NPF}_6/\text{DMSO}$ in the region of the first one-electron reduction step (see inset in Figure 4b), a significant decrease of EPR signal was observed in accord with the formation of diamagnetic $\text{Cu(I)} d^{10}$ complex. For aqueous solutions the reversibility was significantly reduced, implying a more complex mechanism involving the release of the proligand. However, by decreasing the scan rate and going to

the more positive potentials upon reverse scan, a partial recovery of the initial optical bands was also observed in aqueous solutions as shown for **2** in Figure S12. Thus, electrochemical data indicated a likely reduction of Cu(II)-TSC complexes to Cu(I) species with the subsequent release of the proligands.

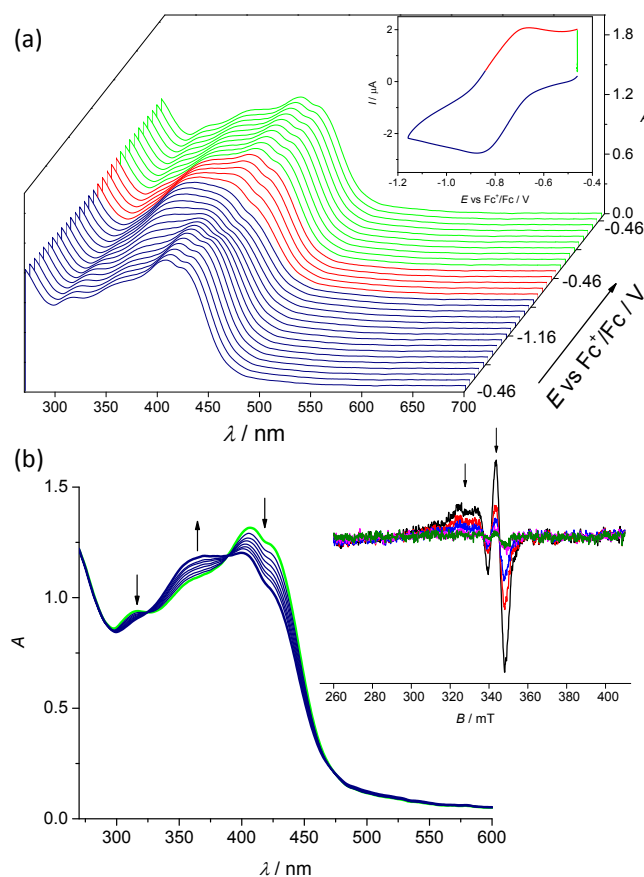


Figure 4. Spectroelectrochemistry of **2** in nBu_4NPF_6 /DMSO in the region of the first cathodic peak. (a) Potential dependence of UV-vis spectra with respective cyclic voltammogram (Pt-microstructured honeycomb working electrode, scan rate $\nu = 5$ $mV\ s^{-1}$); (b) evolution of UV-vis spectra in 2D projection in forward scan (Inset: EPR spectra measured at the first reduction peak using Pt mesh working electrode).

Lead TSCs and their Cu(II) complexes exhibited marked antiproliferative activity in a nanomolar concentration range. The *in vitro* anticancer activity of **1–6** and their respective TSCs was determined in ovarian carcinoma cells (A2780 and A2780cisR) and non-cancerous human embryonic kidney cells (HEK293) by the colorimetric MTT assay with an exposure time of 72 h. The IC_{50} values for **HL¹–HL⁶**

and **1–6** in comparison with Triapine, CuCl₂, Cu-Triapine and cisplatin are listed in Tables 1 and 2, respectively, and concentration-effect curves are depicted in Figure S13.

Table 1. Cytotoxicity of proligands **HL**^{1–6} and their *n*-octanol/water distribution coefficients (logD_{7.4})

Compound	IC ₅₀ [μM] ^a					logD _{7.4} ^d
	A2780	A2780cis	RF ^b	HEK293	SF ^c	
HL ¹	7.6 ± 1.9	13 ± 1	1.7	9.1 ± 1.7	1.2	+0.46 ± 0.01
HL ²	0.010 ± 0.001	0.035 ± 0.006	3.5	0.030 ± 0.005	3.0	+0.51 ± 0.02
HL ³	0.008 ± 0.002	0.028 ± 0.006	3.5	0.022 ± 0.005	2.8	+1.0 ± 0.1
HL ⁴	0.07 ± 0.01	0.76 ± 0.06	10.9	0.35 ± 0.05	5.0	+0.37 ± 0.04
HL ⁵	0.31 ± 0.08	1.1 ± 0.2	3.5	0.67 ± 0.06	2.2	≥2
HL ⁶	157 ± 21	426 ± 44	2.7	89 ± 2	0.6	+0.84 ± 0.01
Triapine	0.67 ± 0.22	1.1 ± 0.1	4.6	0.39 ± 0.05	0.6	n.d. ^e

^a 50% inhibitory concentrations (IC₅₀) in human ovarian carcinoma cell lines A2780 and A2780cisR and human embryonic kidney cell line HEK293, determined by means of the MTT assay after 72 h exposure. Values are means ± standard deviations obtained from at least three independent experiments, ^bResistance Factor (RF) is determined as IC₅₀ (A2780cisR)/IC₅₀ (A2780), ^c Selectivity Factor (SF) is determined as IC₅₀ (HEK293)/IC₅₀ (A2780), ^d Distribution coefficients between *n*-octanol and buffered aqueous solution determined at physiological pH by UV–vis spectroscopy. Values are means ± standard deviations obtained from at least three independent experiments, ^e n.d. – not determined.

With the exception of **HL**¹ and **HL**⁶, the proligands demonstrated marked antiproliferative activity in a submicromolar to nanomolar concentration range in both A2780 and A2780cisR cells. The efficacy of compounds **HL**², **HL**³ and **HL**⁵ in cisplatin-resistant A2780cisR cells decreased by factor 3.5 in comparison to that in sensitive A2780 cells, whereas a 4.6- and 11-fold drop of activity was observed for Triapine and **HL**⁴, respectively. The activity of proligands **HL**^{1–6} and Triapine increased in the following rank order **HL**⁶ < **HL**¹ < **Triapine** < **HL**⁵ < **HL**⁴ << **HL**² ≈ **HL**³ (Table 1).

Table 2. Cytotoxicity of Cu(II)-TSC complexes **1–6**, CuCl₂ and cisplatin and their *n*-octanol/water distribution coefficients (logD_{7.4})

Compound	IC ₅₀ [μM] ^a					Cellular accumulation, ^d nmol Cu/ mg protein	logD _{7.4} ^e
	A2780	A2780cis	HEK293	RF ^b	SF ^c	A2780	
1	2.2 ± 0.1	13 ± 4	16 ± 2	5.9	7.3	0.63 ± 0.09	−0.81 ± 0.04
2	0.012 ± 0.002	0.030 ± 0.003	0.032 ± 0.008	2.5	2.7	3.9 ± 0.6	+0.30 ± 0.01
3	0.26 ± 0.02	0.60 ± 0.05	0.78 ± 0.20	2.3	3.0	2.2 ± 0.6	+0.86 ± 0.02
4	0.08 ± 0.01	0.26 ± 0.04	0.24 ± 0.02	3.3	3.0	3.2 ± 0.7	+0.12 ± 0.01
5	0.009 ± 0.002	0.017 ± 0.002	0.020 ± 0.003	1.9	2.2	4.1 ± 0.9	+1.49 ± 0.06
6	43 ± 3	62 ± 6	72 ± 0	1.4	1.7	0.47 ± 0.08	−0.23 ± 0.01
CuCl₂	83 ± 12	82 ± 3	187 ± 37	1.0	2.3	0.17 ± 0.05 ^f	n.d. ^g
Cu-Triapine^h	1.3 ± 0.1	29 ± 0.3	n.d.	22.3	-	n.d.	n.d.
Cisplatin	0.44 ± 0.13	4.6 ± 0.3	n.d.	10.5	-	-	−2.30 ± 0.79 ⁱ

^a 50% inhibitory concentrations (IC₅₀) in human ovarian carcinoma cell lines A2780 and A2780cisR and human embryonic kidney cell line HEK293, determined by means of the MTT assay after exposure for 72 h. Values are means ± standard deviations obtained from at least three independent experiments, ^bResistance Factor (RF) is determined as IC₅₀ (A2780cisR)/IC₅₀ (A2780), ^c Selectivity Factor (SF) is determined as IC₅₀ (HEK293)/IC₅₀ (A2780), ^d Cellular accumulation in A2780 cells, determined by ICP-MS after 24 h exposure at concentration of 1 μM. Values are means ± standard deviations obtained from at least three independent experiments, ^e Distribution coefficients between *n*-octanol and buffered aqueous solution determined at physiological pH by UV–vis spectroscopy. Values are means ± standard deviations obtained from at least three independent experiments, ^f Cellular accumulation of CuCl₂ was detected at concentration of 2.5 μM, ^g n.d. – not determined, ^h the IC₅₀ values (exposure for 72 h) were taken from the ref.: 27, ⁱ Log P_{o/w} value was taken from the ref. 28.

As can be seen in Table 2, with the exception of **3**, all Cu(II)-TSC complexes were equally or more cytotoxic than the respective proligands. The activity of complexes **1–6** increased in the following order **6** < **1** < **3** < **4** < **2** < **5**, similar to the trend in electrochemical redox potentials for Cu-based reduction

step (Table S6). In order to compare the cytotoxicity of **1–6** with that of $[\text{Cu}(\text{H}_2\text{O})_6]^{2+}$, A2780 cells were treated with aqueous solution of CuCl_2 . In agreement with the literature,²⁹ CuCl_2 revealed antiproliferative activity in the high micromolar range, in contrast to high activity of **1–6**, which should be regarded as individual entities with their own biological, pharmacokinetic and metabolic profiles. Complexes **1–6** were up to ~50 times more cytotoxic than cisplatin in A2780 cells and up to ~270 times more cytotoxic in A2780cisR cells. Additionally, the differences in cytotoxicity of **1–6** in A2780 and A2780cisR cells were significantly lower than for **HL¹–HL⁶** and cisplatin, indicating high potential of Cu(II) complexes for the treatment of cisplatin-resistant tumors. The most active TSC proligands and their respective Cu(II) complexes were more selective towards A2780 cells over HEK293 cells; however, they did not show any selectivity towards A2780cisR cells over HEK293 cells. On the contrary, Triapine was significantly more cytotoxic towards non-cancerous HEK cells than A2780 or A2780cis cells.

Antiproliferative activity of Cu(II)-TSC complexes correlates with their cellular accumulation and lipophilicity. The ability of anticancer drugs to penetrate biological membranes, tissues and barriers underlies their biological, pharmacokinetic and metabolic properties. These properties can be predicted by calculating physicochemical characteristics of drug candidates using molecular descriptors.³⁰ In order to estimate if the proligands **HL¹–HL⁶** and corresponding Cu(II) complexes demonstrate drug-like properties, they were evaluated with main stream molecular descriptors, such as molecular weight (MW), number of hydrogen bond donors and acceptors (HBD/HBA), *n*-octanol-water partition coefficient (log P), polar surface area (PSA) and rotatable bonds (RB). The results are shown in Table S7 in the Supporting Information. Based on the calculated physicochemical parameters, we determined if novel compounds belonged to the *drug-like* chemical space, which is commonly defined by the well-known Lipinski rule, or to the known drug space (KDS), which includes all small compounds in medical use.³¹ As can be seen from Table S7, all compounds fell within the boundaries of *drug-like* chemical

space with the exception of **HL**⁴ which demonstrated higher HBA value, referring to KDS space. Based on these results, novel compounds are expected to be sufficiently cell permeable. Following the calculations, we determined the lipophilicity of all compounds experimentally ($\log D_{7.4}$, Tables 2 and 3). It is known that cancer cells accumulate hydrophilic compounds to a smaller extent relative to hydrophobic compounds, thereby affecting their activity. Hence, the differences in the lipophilicity of compounds might be related to the observed trends in cytotoxicity of **1–6**. In general, with the exception of morpholine derivative **HL**⁴, proligands which were obtained by systematic substitution at the terminal thioamide nitrogen, demonstrated higher lipophilicity when compared to that of **HL**¹. As expected, proligands were more lipophilic than the corresponding Cu(II) species, which are positively charged at physiological pH (Figure 3b). Among Cu(II)-TSC complexes, only the least cytotoxic complexes **1** and **6**, as well as cisplatin, demonstrated negative $\log D_{7.4}$ values, indicating their higher hydrophilicity. The lipophilicity of **2–5** increased in the following order $4 < 2 < 3 < 5$, and correlated well with their cytotoxicity in cancer cells, with the exception of **3**. Subsequently, we determined the total cellular accumulation of Cu in A2780 cells by ICP-MS upon 24 h exposure to **1–6** in comparison with CuCl₂ (Table 4). The IC₅₀ values from MTT assays with 72 h drug exposure varied from nanomolar to high micromolar concentrations; therefore, in the cellular accumulation experiment cells were treated with 1 μ M of compounds of interest for 24 h to minimize cell detachment. The cellular accumulation of all Cu(II)-TSC complexes was significantly higher than for CuCl₂, indicating the role of lipophilic TSC ligands in the delivery of the complexes into the cells. The accumulation increased in the order $6 < 1 < 3 < 4 < 2 < 5$.

Cellular accumulation and efflux of Cu(II)-TSC complexes are energy- and/or temperature-dependent processes. Dependence of cytotoxicity on lipophilicity of **1–6** indicates that their cellular accumulation at least partially occurs via passive diffusion. In order to gain additional insights into the mechanisms controlling the accumulation of Cu(II)-TSC complexes, we investigated the effects of

temperature and various inhibitors on total cellular accumulation of **2** with the results shown in Figure 5. A2780 cells were treated with **2** at 3 μM for 30, 60 and 120 min at 37 and 4 $^{\circ}\text{C}$ and intracellular Cu content was measured by ICP-MS. Prolonged incubation at low temperatures may result in the changes in membrane fluidity, decreasing membrane permeability and restricting drug uptake;³² therefore, the shorter time point of 10 min was also included. The cellular accumulation of **2** dramatically decreased at low temperature even after 10 min treatment, indicating the involvement of active carrier-mediated transport or facilitated diffusion. Likewise, the decrease of cellular accumulation at low temperatures has been reported for both Cu(II)-TSC complexes and metal-free TSCs.^{33,34} To further clarify whether the uptake of **2** requires energy, the cellular ATP production was blocked by incubating cells in a saline solution (HBSS), resulting in total starvation and rapid reduction of intracellular ATP content, and/or by addition of oligomycin, leading to the inhibition of oxidative phosphorylation.

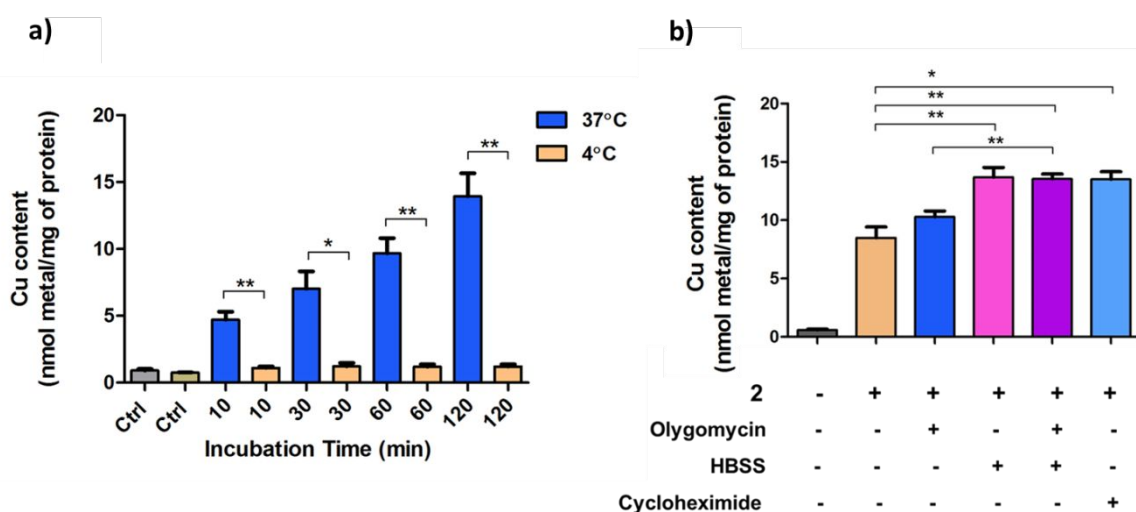


Figure 5. Effects of temperature and inhibitors on the accumulation of **2** in A2780 cells. Intracellular content was determined by ICP-MS: a) A2780 cells were treated with **2** (3 μM) at 37 or 4 $^{\circ}\text{C}$ for the indicated time periods. b) A2780 cells were pre-treated with oligomycin (5 μM), and/or pre-incubated in HBSS for 1 h, or pre-treated with cycloheximide (100 μM) for 4 h and subsequently co-treated with **2** (3 μM) for 1 h. The presence of inhibitors did not show any effects on the intracellular Cu level of untreated A2780 cells (data not shown). Statistical analysis was performed by two-tailed T-test using GraphPad Prism software (GraphPad Software Inc., CA) with $p < 0.05$ considered as significant (* $p < 0.05$, ** $p < 0.01$).

When A2780 cells were pre-treated with 5 μ M of oligomycin for 1 or 4 h and subsequently co-treated with 3 μ M of **2**, the intracellular Cu levels were not affected, even though the mitochondrial respiration of the cancer cells was successfully inhibited (data not shown). However, when A2780 cells were starved in HBSS for 1 h and subsequently treated with 3 μ M of **2** for 1 h, an increase in intracellular Cu content was observed. Co-treatment of **2** with 5 μ M of oligomycin in HBSS did not result in a further increase of intracellular Cu levels. The increase of Cu content in HBSS-starved cells might stem from the inhibition of energy-dependent efflux processes. In order to assess the role of proteins in both cellular accumulation and efflux of **2**, A2780 cells were pre-treated with cycloheximide, which is a known inhibitor of protein synthesis. Significant increase of intracellular Cu content was observed, similar to HBSS starvation, providing further evidence for the involvement of protein-dependent efflux processes. Similar effects have been reported for related Cu(II)-TSC complexes.³³ Accumulated data provide evidence that cellular accumulation of **2** occurs via carrier-facilitated diffusion, which might co-exist with passive permeation through the lipid bilayer.³⁵ The efflux processes, however, are actively mediated by proteins and upon inhibition of protein synthesis or energy production a strong increase of intracellular Cu concentration was observed.

Cu(II)-TSC complexes demonstrated high antibacterial activity against *S. aureus* in a comparable concentration range to ciprofloxacin. The antibacterial activity of **HL**¹–**HL**⁶ and **1**–**6** on planktonic cells of *P. aeruginosa* and *S. aureus* was investigated by determination of MIC₅₀ and MIC₁₀₀ values (Table 3). Overall, compounds showed a higher antimicrobial activity against Gram-positive (*S. aureus*) compared to Gram-negative bacteria (*P. aeruginosa*). The highest activity against *S. aureus* was observed for **2**–**5**, which specifically inhibited its growth with MIC₅₀ values ranging between 2 and 5 μ g/mL and completely abolished its growth at 10 μ g/mL (MIC₁₀₀). This activity is comparable but slightly lower than the antibacterial activity of a well-known benchmarked antibiotics ciprofloxacin

(CPX). On the contrary, *P. aeruginosa* exhibited high resistance to most of the compounds tested with the exception of **HL**² and **2** with MIC₅₀ values at 70 and 50 µg/mL, respectively.

Table 3. Antibacterial activity of **HL**^{1–6} and **1–6**^a

	<i>S. aureus</i> (µg/mL)		<i>P. aeruginosa</i> (µg/mL)	
	MIC ₅₀	MIC ₁₀₀	MIC ₅₀	MIC ₁₀₀
HL ¹	100	>400	NA	NA
HL ²	10	400	50	100
HL ³	10	300	100	NA
HL ⁴	100	>400	NA	NA
HL ⁵	10	300	NA	NA
HL ⁶	NA	NA	NA	NA
1	100	>500	NA	NA
2	2	10	70	200
3	3	10	100	400
4	5	10	100	400
5	3	10	200	450
6	NA	NA	200	NA
Ciprofloxacin	0.5	1	0.5	2

^aSee experimental section for details. NA denotes no antibacterial activity detected (>500 µg/mL).

Compounds **HL**⁶ and **6** based on S-methylisothiosemicarbazide were completely devoid of antibacterial activity and no inhibition of bacterial growth was observed at the whole concentration range tested. Interestingly, metal-free thiosemicarbazones **HL**^{1–HL}⁵ displayed relatively low antibacterial activity (MIC₁₀₀ > 300 µg/mL), which was significantly enhanced upon coordination to Cu(II). Subsequently, the mode of action of the most active compounds **2–5** was evaluated in *S. aureus* and *P. aeruginosa* using the Live/Dead viability assay. As shown in Figure 6, compounds **2–5** greatly reduced cell

replication, reflected by the lower number of cells compared to the untreated sample and high percentage of PI-stained cells, indicating compromised cell membrane allowing the penetration of propidium iodide and a bacteriostatic action of these compounds.

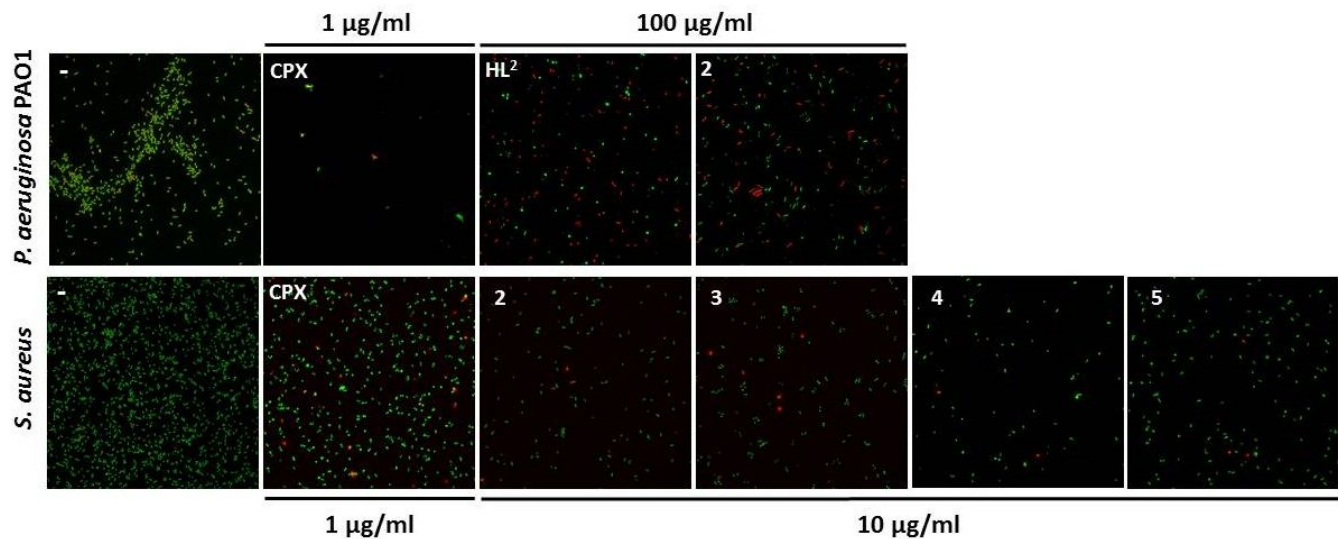


Figure 6. Live/dead bacterial cells are shown by staining with green fluorescence indicating live cells and red fluorescence indicating dead cells.

TSCs and their Cu(II) complexes are located in a close proximity to the active site of mouse R2 RNR protein. Next, the RNR inhibitory potential of the TSC proligands **HL¹–HL⁶** and their corresponding Cu(II) complexes **1–6** was investigated. The proligands **HL¹–HL⁶** and Cu(II) complexes **1–6** were docked into the crystal structure of mouse R2 RNR subunit (PDBid:1W68)³⁶ using GOLD software.³⁷ To predict the binding of **1–6**, the GoldScore function was used and its parameters were modified to include Cu since they are not in GOLD’s database.³⁸ The scoring functions are presented in Table S8. Overall, similar docking results were observed for all proligands and their Cu(II) complexes, suggesting a plausible binding to the mouse R2 protein. The compounds were deeply embedded into the pocket of the R2 subunit, close to the diferric center (Fe₂O), and the enzymatically essential tyrosyl residue (Tyr177). The docked configuration of **5** in comparison with Triapine into the binding site is

shown in Figure 7. An overlap with Triapine was predicted, resulting in a partial reproduction of a hydrogen-bonding pattern (Figure S14, Table S8).

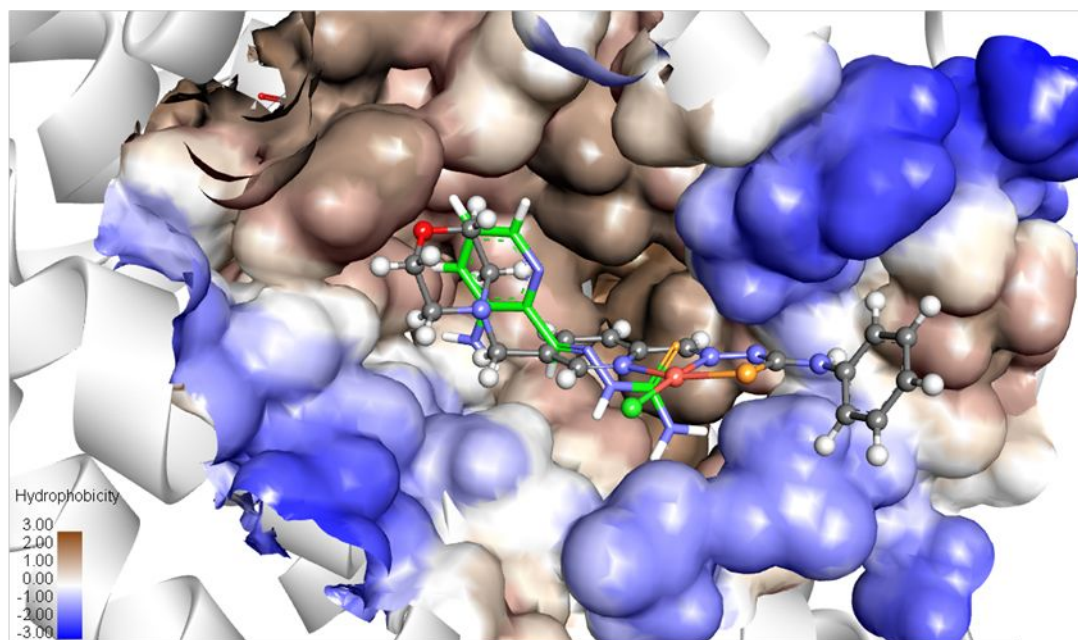


Figure 7. The overlay of docked configurations of Cu(II) complex **5** (grey) and Triapine (green) in the binding site of the mouse R2 RNR protein (PDB ID: 1W68). The surface is rendered. Blue and brown depict hydrophilic and hydrophobic areas, respectively.

Lead TSCs and their Cu(II) complexes cause moderate inhibition of a mouse R2 RNR protein in a reducing environment as a result of iron chelation. The effects of the most cytotoxic complexes **2** and **5**, and the corresponding proligands **HL**² and **HL**⁵, on the mouse R2 tyrosyl radical were investigated by EPR spectroscopy at 30 K. The proligands and their Cu(II) complexes were incubated with the mouse R2 RNR protein at 1:1 protein-to-compound mole ratio at 298 K in the presence or absence of the reducing agent dithiothreitol (DTT), and the time-dependent reduction of the tyrosyl radical was monitored.³⁹ Tyrosyl radical reduction by the investigated compounds was observed only in the presence of DTT. Upon incubation with **HL**², **HL**⁵ and **2**, **5**, 50% of tyrosyl radical was reduced within 90 s, after which no further reduction was observed (Figure S15). Since the diferric center of the mouse R2 protein is expected to be reduced upon addition of DTT,^{40,41} the ability of proligands **HL**² and **HL**⁵ to form

stable Fe(II) complexes was investigated by monitoring these reactions by UV–vis measurements in buffered aqueous solutions at pH 7.4. **HL**² and **HL**⁵ ($\lambda_{\text{max}} = 312$ and 320 nm, respectively) readily reacted with $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ affording Fe(II) bis-ligand complexes ($\lambda_{\text{max}} = 373, 610$ nm and $\lambda_{\text{max}} = 385, 645$ nm, respectively) at both 1:1 and 1:2 Fe-to-ligand mole ratios. Additionally, the formation of Fe(III) complexes with the subsequent reduction by DTT was investigated (Figure S15). **HL**² and **HL**⁵ reacted with $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ (at 1:1 or 1:2 Fe-to-ligand mole ratios) to give $[\text{Fe}^{\text{III}}(\text{L}^2)_2]^+$ and $[\text{Fe}^{\text{III}}(\text{L}^5)_2]^+$ complexes ($\lambda_{\text{max}} = 378$ nm, 612 nm and $\lambda_{\text{max}} = 386$ nm, 635 nm, respectively). The formation of $[\text{Fe}^{\text{III}}(\text{L}^5)_2]^+$ species was confirmed by ESI-MS (Figure S16). The absorption spectra of reduced forms were almost identical to those of Fe(II) bis-ligand species, and to the previously reported spectra of other Fe^{II}-TSC complexes.⁴²

HL² and **2** induced unfolded protein response (UPR), antioxidant defense and cell cycle perturbations. In order to get further insight into the mechanism of action of **2**, Western Blotting of ER stress-related protein markers, namely IRE1 α , CHOP, BiP/GRP78 and p-ERK, was performed (Figure 8). When A2780 cells were treated with increasing concentrations of **2** for 24 h, the dose-dependent upregulation of IRE1 α and CHOP proteins, decrease of ER stress chaperone BiP/GRP78, as well as phosphorylation of ERK were observed as a response to ER stress. To assess the effects of **2** on the cell cycle, we examined the expression of cyclin D1 and cyclin B1 which serve as cell cycle regulatory switches in actively proliferating cells and are required for cell cycle progression in G₁ and G₂ phases, respectively. Complex **2** displayed significant inhibition of cyclin D1 expression, indicating cell cycle arrest at G₁/S phase, whereas no changes in cyclin B1 expression were detected. Subsequently, we compared the effects of **2** and the respective proligand **HL**² on the expression of p-ERK1/2 and the marker of antioxidant defence NRF2. Both **HL**² and **2** demonstrated phosphorylation of ERK1/2; however, the effects on NRF2 were markedly different. Whereas no increase of NRF2 expression was observed, when cells were treated with **HL**², complex **2** induced increased expression of NRF2 even at

low concentration, indicating activated antioxidant defense, possibly as a result of ROS insult caused by Cu(II) reduction.

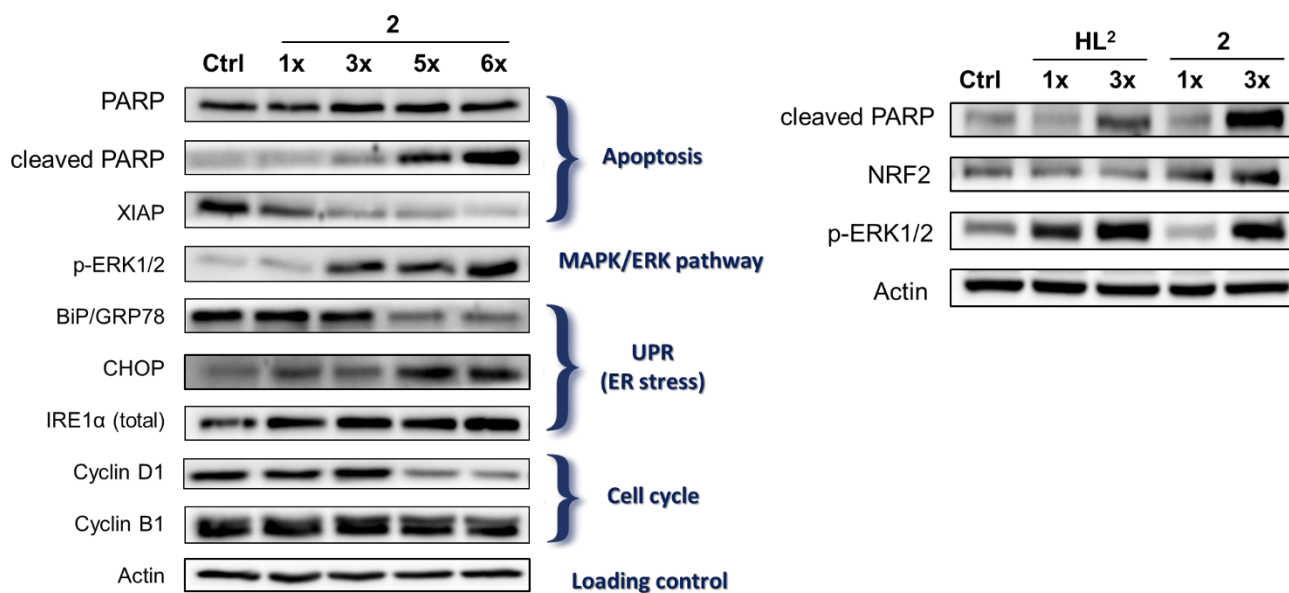


Figure 8. Western Blot analysis of various proteins. A2780 cells were treated with increasing concentrations of proligand **HL**² and complex **2** (corresponding to IC₅₀ value from MTT experiment with exposure time of 72 h) for 24 h. Total lysates were isolated and examined by Western Blot. Actin was used as a loading control.

HL² and **2** induced dose-dependent apoptosis accompanied by PARP cleavage and XIAP inhibition. To elucidate if observed cytotoxicity of new compounds described in this study was a result of apoptosis induction, we conducted Annexin V/PI apoptosis assay. In brief, fluorochrome-labeled Annexin V reagent is used for detection of membrane phosphatidylserine (PS), which translocates to the cellular surface from the inner side of plasma membrane in the event of early apoptosis. Simultaneously, propidium iodide (PI) is used for detection of later stages of apoptosis or necrosis, which are characterized by loss of membrane integrity and accumulation of PI inside the cells. A2780 cells were treated with 2IC₅₀ and 6IC₅₀ concentrations of **HL**² and **2** (IC₅₀ values were obtained from MTT experiment in A2780 cells with the exposure time of 72 h). Results are illustrated in Figure 9. Upon incubation of A2780 cells with increasing concentrations of **HL**² and **2**, the dose-dependent increase of apoptotic cells was observed.

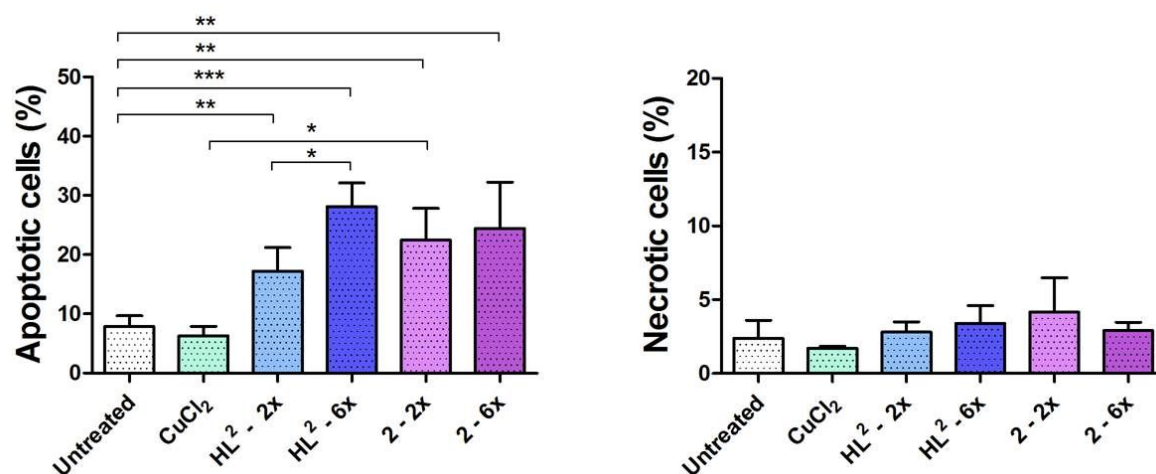


Figure 9. Bar graphs showing the percentage of cell death due to apoptosis (left) and necrosis (right) in A2780 cells treated with increasing concentrations of **HL²**, **2** (2IC₅₀ and 6IC₅₀) and CuCl₂ and detected by Annexin V/PI apoptosis assay. Statistical analysis was performed by two-tailed T-test using GraphPad Prism software (GraphPad Software Inc., CA) with $p < 0.05$ considered as significant (* $p < 0.05$, ** $p < 0.01$).

At low concentrations both the proligand and Cu(II) complex induced a 2- to 3-fold increase of apoptotic cells in comparison with untreated cells (7.8 ± 1.8 %, 17.2 ± 3.9 % and 22.5 ± 5.3 % for untreated cells, **HL²** (2IC₅₀) and **2** (2IC₅₀), respectively). At high concentrations a further increase of apoptotic cells was observed (28.1 ± 4.0 % and 24.4 ± 7.8 % for **HL²** (6IC₅₀) and **2** (6IC₅₀)). No significant differences between the apoptosis-inducing properties of the proligand **HL²** and **2** were noticed. When CuCl₂ was incubated with the cells at the concentration, corresponding to Cu content in **2** at 2IC₅₀, no induction of apoptosis was detected, which can be explained by poor cellular accumulation of CuCl₂. The proportion of necrotic cells was negligible and no significant deviation from untreated cells was detected; therefore, the induction of necrosis by **HL²** and **2** was ruled out. Subsequently, the effects of **HL²** and **2** on apoptotic signaling in cells were investigated by Western Blotting. A2780 cells were treated with increasing concentrations of the drug for 24 h, and the cleavage of poly (ADP-ribose) polymerase-1 (PARP) was investigated (Figure 8). Cleavage of poly (ADP-ribose) polymerase-1 (PARP) through suicidal proteases such as caspases has been widely accepted as an indicator of apoptosis.⁴³ As can be seen in Figure 8,

both **HL²** and **2** revealed a dose-dependent PARP cleavage, indicating apoptosis induction in agreement with the results of Annexin V/PI assay. Additionally, expression of X-linked inhibitor of apoptosis (XIAP) was markedly reduced in a dose-dependent manner upon treatment with increasing concentrations of **2**.

Discussion

The diverse biological and chemical properties of Triapine and other TSCs, as well as their transition metal complexes, have been extensively studied for decades.^{44,45,46,47,48,41,49,50,51} Their antiproliferative and antibacterial activity varies greatly, from low nanomolar to high micromolar range, depending on the substituents at the TSC backbone.^{16,20,52} Even though clinical trials demonstrated some benefits of patient treatment with Triapine, its low aqueous solubility and high toxicity prevent it from further clinical progress. In a continuous effort to improve the bioavailability and therapeutic profile of TSCs and Cu(II)-TSC complexes, novel thiosemicarbazone hybrids with bioactive L-proline, homoproline, piperazine or iminodiacetate moieties, using the molecular hybridization approach, have been designed and synthesized.^{16,17,18,19,20} However, the improvement of the aqueous solubility was often accompanied by a significant decrease in cytotoxicity. In this study, the attachment of N-substituted morpholine moiety at the TSC backbone resulted in a simultaneous improvement of aqueous solubility and cytotoxicity (up to ~50 times in comparison with Triapine). To ensure the selectivity of novel TSCs towards cancer cells over healthy cells, their cytotoxicity against A2780 cancer cells and noncancerous HEK293 cells was compared. It was shown that new compounds were 2–5 fold more toxic towards cancer cells than healthy cells (Tables 1 and 2), whereas Triapine was more toxic towards noncancerous cells, which is in agreement with its known high toxicity. The coordination of TSCs to Cu(II) resulted in Cu(II)-TSC complexes with similar or improved antiproliferative activity. In general, when TSCs are coordinated to Cu(II), an increase in cytotoxicity is observed;^{17,18,53} however, Cu(II)-Triapine complex was shown to be significantly less cytotoxic than Triapine itself.²⁷ The activity of the Cu(II) complexes

1–6 was in a good correlation with their lipophilicity, cellular accumulation, as well as their electrochemical redox potentials (Tables S6 and 2). Interestingly, metal-free TSCs exhibited only low antibacterial properties against *S. aureus*, whereas Cu(II)-TSC complexes revealed significant antibacterial activity similar to that of the commonly used antibiotic ciprofloxacin (Table 4). These results indicate that coordination of TSCs to Cu(II) improved their therapeutic potential. It should be noted that the replacement of N,N,S- with N,N,N-chelating moiety in **6** resulted in a drop of cytotoxicity and antibacterial activity, which is in agreement with the previously reported data.²⁰

The antiproliferative activity of Triapine and other TSC proligands is often related to their Fe-chelating properties.⁵⁴ To sustain their high proliferation rate, cancer cells rely on increased uptake of Fe from Fe-transporting proteins; therefore, Fe chelation is a valuable therapeutic strategy for cancer treatment. Based on UV-vis and ESI-MS experiments, it was demonstrated that irrespective of metal-to-ligand ratio, proligands **HL**² and **HL**⁵ readily reacted with Fe(III) to form $[\text{Fe}^{\text{III}}(\text{L}^2)_2]^+$ and $[\text{Fe}^{\text{III}}(\text{L}^5)_2]^+$ species, which could be reduced to Fe(II) bis-ligand complexes by the reducing agent DTT. These results indicate that morpholine-TSC hybrids, investigated in this study, may act as Fe chelators. Additionally, spectroelectrochemical experiments revealed that under reducing conditions, the Cu(II)-TSC complexes could undergo reversible reduction to Cu(I) species, with simultaneous release of proligands, suggesting that these complexes may act as Fe chelators, also in cancer cells. In addition, the presence of the Cu(II) ion in the cells may lead to the production of ROS *in vivo*, due to Cu(II)/Cu(I) redox cycling.⁵⁵

It is well known that one of the mechanisms contributing to the cytotoxic effects of Fe chelators is inhibition of the RNR enzyme, which is highly expressed in cancer cells, and essential for their DNA synthesis. The mechanism of RNR inhibition by Triapine has been extensively studied by several research groups.^{41,46,47,51,56} It has been proposed that Triapine interferes with the assembly of the differic-tyrosyl radical cofactor in R2, necessary for the catalytic activity of RNR.^{46,56} Moreover, it has

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2 been shown that the presence of Fe is required for effective R2 inhibition,^{44,45} and that it is actually the
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4 Fe(II)-Triapine species that is responsible for the R2-specific RNR inhibitory effect of
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6 Triapine.^{41,46,47,51-58} The Fe(II)-Triapine complex may be formed by chelation of Fe from the differic
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8 cofactor in R2 *in vitro*,^{44,41} and/or from intracellular iron pools *in vivo*.^{46,56} The potent inhibition of
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10 human,⁴⁷ and mouse⁴¹ R2 RNR *in vitro* by catalytic amounts of Fe(II)-Triapine has been proposed to
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12 involve ROS. Namely, ROS had been spin-trapped, and detected by EPR spectroscopy, in the aerobic
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14 reaction between human R2 RNR and Fe(II)-Triapine, which had implicated that O₂ is important in
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16 tyrosyl radical destruction, and that ROS may ultimately be responsible for the pharmacologic effects of
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18 Triapine *in vivo*.⁴⁷ In another study of time-dependent tyrosyl radical reduction in mouse R2 by Triapine,
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20 and its Zn, Ga, Cu, and Fe complexes, the requirement of O₂ in R2 inhibition was also suggested.⁴¹ This
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22 was based on the observation that substoichiometric amounts of Fe(III)-Triapine relative to R2 (protein-
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24 to-complex mole ratio, 5:1) reduced only 20% tyrosyl radical in anaerobic conditions, compared to
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26 100% in aerobic conditions. However, more recently, in a different experimental setup, namely in the
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28 presence of a 10-fold excess of Fe(II)-Triapine over R2 protein, the role of ROS (and O₂) in human R2
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30 inhibition was excluded.⁵¹ This study showed that the principle mechanism of human R2 RNR inhibition
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32 by Triapine, based on kinetic measurements of tyrosyl radical, and ⁵⁵Fe loss, is direct radical quenching,
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34 in an iron-loaded protein. Furthermore, it implied that Fe(II)-Triapine can rapidly reduce the tyrosyl
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36 radical, while leaving the protein in the met-state.
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46 In order to investigate if R2 RNR could be the potential biomolecular target for the novel morpholine-
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48 TSC hybrids, and their Cu(II) complexes, molecular docking studies, and R2 RNR tyrosyl radical
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50 reduction kinetic experiments, were performed. It has been suggested that the efficiency of R2 inhibition
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52 is likely to depend on the access of the inhibitors to the differic center of R2;^{41,47} therefore, we estimated
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54 the likelihood of the binding of TSC proligands **HL**¹–**HL**⁶ and their corresponding Cu(II) complexes
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56 **1**–**6** to the surface of the R2 protein. It was demonstrated that new TSCs, and their Cu(II) complexes
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were deeply embedded into the pocket of the R2 protein and exhibited hydrogen bonding patterns similar to Triapine (Figure 7). Based on the results of the molecular docking studies suggesting plausible binding to the mouse R2 protein, the time-dependent tyrosyl radical reduction in the mouse R2 protein upon incubation with the most cytotoxic complexes, **2** and **5**, and the corresponding proligands **HL**² and **HL**⁵, at a 1:1 protein-to-compound ratio, was measured by EPR spectroscopy. Only in the presence of DTT (which can reduce Fe(III) to Fe(II) in the R2 differic cofactor),⁴⁰ **HL**² and **HL**⁵ caused 50% tyrosyl radical reduction (Figure S15a). These results show that R2 inactivation by **HL**² and **HL**⁵ is not as efficient as with Triapine, which causes 100% tyrosyl radical loss in mouse R2 after 5 min (in the same experimental conditions).⁴¹ The most plausible explanation for this observation is that different mechanisms are likely to be involved in R2 inhibition by **HL**², **HL**⁵ and Triapine. The R2 inhibitory activity of the corresponding Cu(II) complexes **2** and **5** was found to be similar to that of the proligands (Figure S15a). This is likely due to the fact that in the presence of DTT, the reduction of Cu(II) to Cu(I) leads to the release of the proligands, which is in agreement with the results of spectroelectrochemical experiments (*vide supra*). Since the UV-vis, and ESI-MS experiments showed that **HL**² and **HL**⁵ act as tridentate ligands, and form stable Fe(II)L₂ complexes (Figure S15), it is possible to propose that the mechanism of R2 inhibition by the TSC proligands involves Fe chelation from the differic cofactor. Based on the extent of tyrosyl radical reduction, it is tempting to speculate that the subsequently formed Fe(II) complexes are not potent inhibitors like the Fe(II)-Triapine complex. However, as this was not the aim of this paper, it is reasonable to refrain from any conclusions prior to the detailed mechanistic studies of mouse, human, and p53 R2 RNR inhibition by the morpholine-TSC hybrids. Moreover, a separate study should be dedicated only to the comparison of the inhibitory potentials of selected TSCs, and their Fe(II) complexes, in order to comprehensively understand the structure-activity relationship, which in turn may explain the mechanism of R2 RNR inhibition. Finally, even though R2 inhibition can be partially attributed to the marked anticancer activity of **2** and **5**, other biomolecular targets are likely to be involved (Figure 10).

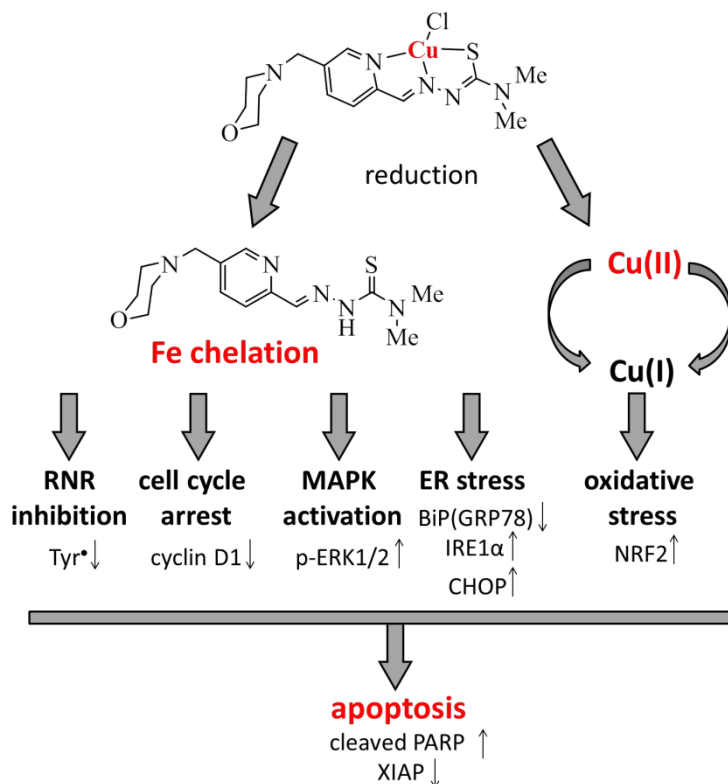


Figure 10. Proposed molecular mechanism of complex **2**.

Fe homeostasis is an intrinsically complex process; hence, Fe sequestration results in the alteration of various biomolecular pathways. Besides RNR, Fe depletion affects multiple molecular targets including those regulating the cell cycle.⁵⁷ It has been reported that the expression of cyclin D1, which ensures the progression through the G₁/S phase, was strongly dependent on the cellular Fe status.⁵⁸ It is believed that in Fe-depleted conditions, down-regulation of cyclin D1 prevents cells from entering the S phase, where Fe is needed for the activity of the RNR enzyme and DNA synthesis.⁵⁹ Fe chelation by TSCs often results in the down-regulation of cyclin D1 and subsequent cell cycle arrest at the G₁/S phase.^{60,61} As expected, complex **2** was shown to induce strong dose-dependent down-regulation of cyclin D1, whereas the protein expression of cyclin B1, which is responsible for G₂/M progression, remained unchanged. In order to further investigate the potential biomolecular targets of Fe-chelating compounds of interest, we studied the phosphorylation of ERK protein, which belongs to the MAPK family. The MAP kinases occupy the central position in cell growth and apoptosis and each MAP kinase member plays a different

role in cellular responses.⁶² It was reported that some Fe chelators induced cancer cell death associated with the activation of p38 MAPK and ERK.⁶³ Moreover, the MAPK-mediated cell death was characterized by the decreased expression of cyclin D1.⁶⁴ Complex **2**, as well as the corresponding proligand **HL**² caused dose-dependent phosphorylation ERK in line with its activation, suggesting that MAPK pathway might be a potential mediator of the cell death, induced by this complex.

Recently, it was discovered that chelation and depletion of intracellular Fe leads to the activation of certain ER stress-response proteins.^{65,66} Previously, several TSCs and their Cu(II) complexes were reported to induce ER stress, characterized by UPR activation,^{67,68} ER expansion and release of intracellular calcium.⁶⁹ Since under reducing conditions, the novel TSCs and their Cu(II) complexes demonstrated formation of Fe(II)-TSC complexes, it may be suggested that they could possibly induce sequestration of cellular Fe resulting in the ER stress. It occurs due to the accumulation of misfolded and/or unfolded proteins in ER, resulting in the disruption of cellular homeostasis. In order to overcome stress, cells activate a rescue program known as UPR, which operates via several signaling pathways, namely ATF6, PERK and IRE1 α . However, in case of severe ER stress, the pro-survival program is switched off and pro-apoptotic signaling takes place, characterized by the enhanced expression of CHOP protein. Complex **2** was shown to induce dose-dependent increase in the expression of IRE1 α and CHOP proteins, indicating the induction of UPR signaling as a result of ER stress. Cell treatment with **2** resulted in a concentration-dependent decrease of BiP/GRP78 protein marker. The ER chaperon BiP/GRP78 plays a central role in the survival machinery and its overexpression is associated with tumor progression and metastases.⁷⁰ The inhibition of BiP/GRP78 indicates severe ER stress, resulting in the suppression of pro-survival chaperones. The antibacterial properties of novel Cu(II)-TSC complexes can be also related to ER stress induction, since it was recently demonstrated *in vitro* and *in vivo* experiments that chemical modulation of ER stress might result in the overload of its machinery that can be effective in elimination of bacterial infection.⁷¹ Additionally, it was reported that efficient killing of methicillin-

resistant *S. aureus* was dependent on the induction of IRE1 α UPR pathway and required sustained generation of ROS.⁷² Based on the observations that **2**, but not proligand **HL**² induced antioxidant defence in cancer cells characterized by NRF2 induction, it may be speculated that the ability of **2**, but not proligand **HL**², to induce oxidative and ER stress, may be related to the beneficial antibacterial activity of Cu(II)-TSC complexes over metal-free TSCs. However, it is not clear yet why new complexes **1–6** did not inhibit the growth of *P. aeruginosa* and their mechanism of action should be investigated in more detail.

In order to cope with the severe perturbations imposed by drug treatment, cancer cells develop adaptive responses to provide them with survival advantage. However, novel TSCs and their Cu(II) complexes significantly disturbed cellular life-death balance towards cell death resulting in the induction of apoptosis. Both proligand **HL**², and complex **2** were demonstrated to induce apoptosis characterized by PARP cleavage, decreased expression of XIAP and increased PS translocation. It should be noted that XIAP is a determining factor of cisplatin chemoresistance in ovarian cancer cells as it effectively suppresses apoptosis via caspase-3 and caspase-7 inhibition.⁷³ Therefore, the ability of **2** to decrease XIAP expression is highly important and to the best of our knowledge has not been previously reported for TSCs and metal thiosemicarbazones, with the exception of Triapine.⁷⁴ Some TSCs and their respective Cu(II) complexes have been reported to simultaneously induce various types of cell death including apoptosis,^{75,76} autophagy^{76,77} and methuosis, which is a very special form of non-apoptotic cell death involving massive cytoplasmic vacuolization.⁷⁸ Therefore, induction of other forms of cell death by new TSCs and their Cu(II) complexes presented in this study cannot be excluded. The results show that the attachment of the morpholine moiety at the TSC backbone and the subsequent coordination to Cu(II), resulted in new drug candidates, likely with better therapeutic profile than Triapine and a promising potential for further clinical development.

Conclusions

One of the reasons hindering the clinical development of Cu(II) thiosemicarbazone complexes is their low aqueous solubility. The decoration of the TSC backbone with functional groups that increase the aqueous solubility and bioavailability often results in a significant decrease of the compound activity. In this work, attachment of a morpholine moiety to 2-formylpyridine (iso)thiosemicarbazone in position 5 of 2-formylpyridine moiety yielded new water-soluble TSC-morpholine hybrids and their Cu(II) complexes, which demonstrated high antiproliferative activity against cisplatin-sensitive and cisplatin-resistant ovarian carcinoma cells, in a nanomolar to submicromolar concentration range. Sufficient aqueous solubility of the new compounds enabled the study of their solution behavior and electrochemical properties under conditions similar to the intracellular environment. Solution speciation studies of **1** revealed that the dissociation of the complex does not occur at the physiological pH, even at micromolar concentrations, indicating its high stability in aqueous media. However, under reducing conditions associated with the cancer cell environment, the reversible reduction of Cu(II) with subsequent release of the proligand within the biologically accessible electrochemical window is plausible. Since the R2 RNR protein is believed to be one of the biomolecular targets of TSCs, and the fact that the proligands form 2:1 complexes with Fe(II), the chelation of Fe(II) from R2 was expected. The Cu(II)-TSCs and their corresponding proligands showed only moderate Tyrosyl radical reduction in mouse R2. Therefore, the marked anticancer activity of the compounds investigated in this work was not solely related to their RNR inhibitory potential, but also to the induction of apoptosis as a result of ER stress, MAPK activation and cell cycle perturbations. Studies of the mechanisms controlling cellular accumulation of the lead drug **2** have shown that it accumulates in cancer cells as a result of passive and/or facilitated diffusion. Finally, it was discovered that coordination of TSC ligands to Cu(II) resulted in a significant enhancement of their antibacterial properties against *S. aureus*. The combination of excellent antiproliferative and antibacterial activity, as well as water-solubility, is a sound basis for further development of this class of Cu(II) thiosemicarbazones as dual pharmaceutical agents.

Experimental

1. Chemicals. 2,5-Pyridinecarboxylic acid was purchased from Alfa Aesar. Ethylenediaminetetraacetic acid (EDTA), KOH and 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES) were purchased from Sigma-Aldrich in puriss quality. KCl, CuCl₂, hydrochloric acid and *n*-octanol are products of Molar Chemicals. CuCl₂ stock solution was prepared from anhydrous CuCl₂ and water and its exact concentration was determined by complexometry through the EDTA complex. All solvents were of analytical grade and used without further purification. Milli-Q water was used for sample preparation. The synthesis of the 5-methylmorpholine-pyridine-2-carboxaldehyde **H** (Scheme 1) is described in detail in Supporting Information. Chemical grade cisplatin (1 mg/mL) was purchased from Hospira Pty Ltd (Melbourne, Australia). IGEPAL CA-630, DL-dithiothreitol (DTT), tetramethylethylenediamine (TEMED), sodium deoxycholate, non-fat dried milk bovine, TWEEN® 20, ponceau S, propidium iodide (PI) were purchased from Sigma-Aldrich (St Louis, MO, USA). Thiazolyl Blue tetrazolium bromide (MTT) was purchased from Alfa Aesar. Tris was purchased from Vivantis Technologies. Glycine, Hyclone™ Trypsin Protease 2.5% (10×) solution, RPMI 1640, DMEM medium, Fetal bovine serum (FBS), Bovine Serum Albumin (BSA), Hank's Balanced Salt Solution (HBSS) and Pierce™ Protease and Phosphatase Inhibitor Mini Tablets were purchased from Thermo Fisher Scientific. Hyclone™ Dulbecco's Phosphate-Buffered Saline (10×), were purchased from GE Healthcare Life Sciences. Biorad protein assay dye reagent concentrate, 30% Acrylamide/Bis solution, 5× Laemmli Sample Buffer, Nitrocellulose Membrane (0.2 μm) were purchased from Bio-rad Laboratories. Luminata™ Classico and Crescendo Western HRP substrate were purchased from Merck Millipore Corporation. Milli-Q-grade purified water was obtained from a Milli-Q UV purification system (Sartorius Stedim Biotech S.A., Aubagne Cedex, France). All antibodies were obtained from Cell Signaling Technologies (Beverly, MA, USA). Nitric acid (65% to 71%, TraceSELECT Ultra) for ICP-MS analysis and *n*Bu₄NPF₆ for cyclic voltammetry experiments were obtained from Fluka (Sigma Aldrich) and used without further purification. Cu and In standards for ICP-MS measurements were obtained from CPI

international (Amsterdam, The Netherlands). Cycloheximide, oligomycin, annexin V-FITC apoptosis detection reagent (500X) were purchased from Abcam (Cambridge, UK).

2. Synthesis of proligands.

The yields, m.p. and analytical data for **HL**¹–**HL**⁶ and **1**–**6** are presented in Tables S9 and S10. The experimental CHNS contents were within ± 0.4 with those calculated, providing evidence for $\geq 95\%$ purity.

5-(morpholinomethyl)pyridine-2-carboxaldehyde thiosemicarbazone (HL¹). 5-morpholinomethylpyridine-2-carboxaldehyde (0.29 g, 1.41 mmol), thiosemicarbazide (0.13 g, 1.41 mmol) and EtOH (5 mL) were mixed in a 25 mL Shlenk tube. The reaction mixture was stirred at 78 °C overnight. The next day the reaction mixture was cooled to room temperature and stored at 4 °C for 4 h. The white crystalline precipitate was filtered off, washed with cold EtOH and dried in vacuo. Yield: 0.34 g. ¹H NMR 500 MHz (DMSO-d₆, δ , ppm): ¹H NMR (500 MHz, DMSO) δ 11.62 (s, 1H, NH¹⁶), 8.48 (d, J = 1.3 Hz, 1H, H⁶), 8.34 (s, 1H, NH¹⁹), 8.24 (d, J = 8.1 Hz, 1H, H³), 8.15 (s, 1H, NH¹⁹), 8.09 (s, 1H, H¹⁴), 7.74 (dd, J = 8.2, 1.7 Hz, 1H, H⁴), 3.59 – 3.54 (m, 4H, H¹⁰, H¹²), 3.51 (s, 2H, H⁷), 2.36 (s, 4H, H⁹, H¹³). ¹³C NMR (126 MHz, DMSO) δ 178.74 (C_q, C¹⁷), 152.73 (C_q, C²), 150.12 (CH, C⁶), 142.85 (CH, C¹⁴), 137.51 (CH, C⁴), 134.23 (C_q, C⁵), 120.23 (CH, C³), 66.57 (2CH₂, C¹⁰, C¹²), 59.74 (CH₂, C⁷), 53.49 (2CH₂, C⁹, C¹³). IR (ATR, selected bands, $\tilde{\nu}_{\text{max}}$): 1612, 1527, 1468, 1400, 1339, 1274, 1102, 1004, 938, 859, 626 cm⁻¹.

5-(morpholinomethyl)pyridine-2-carboxaldehyde 4,4-dimethyl-3-thiosemicarbazone (HL²). 5-(morpholinomethyl)pyridine-2-carboxaldehyde (0.29 g, 1.41 mmol) and 4,4-dimethyl-3-thiosemicarbazide (0.17 g, 1.41 mmol) were dissolved in EtOH (5 mL) and stirred in a Shlenk tube at 78 °C for 2 h until the solution became clear. The reaction mixture was left to stand at 4 °C for 6 h. The yellow crystalline precipitate was filtered off, washed with cold EtOH and dried in vacuo. Yield: 0.25 g. ¹H NMR (500 MHz, DMSO) δ 15.08 (s, 1H, NH¹⁶), 8.71 (s, 1H, H³), 8.49 (s, 1H, H⁶), 8.23 (d, 1H, H¹⁴), 7.76 (dd, J = 13.9, 8.1 Hz 1H, H⁴), 3.59 (s, 4H, H¹⁰, H¹²), 3.52 (s, 2H, H⁷), 3.38 (s, 4H, H²⁰, H²¹),

2.37 (s, 4H, H⁹, H¹³). ¹³C NMR (126 MHz, DMSO) δ 180.02 (C_q, C¹⁷), 153.06 (C_q, C²), 150.27 (CH, C⁶), 148.84 (CH, C³), 144.29 (CH, C¹⁴), 137.75 (CH, C⁴), 134.06 (C_q, C⁵), 66.61 (2CH₂, C¹⁰, C¹²), 59.77 (CH₂, C⁷), 53.53 (2CH₂, C⁹, C¹³), 42.72 (2CH₃, C²⁰, C²¹). IR (ATR, selected bands, $\tilde{\nu}_{\text{max}}$): 1497, 1443, 1362, 1303, 1209, 1158, 1108, 1064, 1003, 901, 862, 716, 627 cm⁻¹.

5-(morpholinomethyl)pyridine-2-carboxaldehyde *N*-pyrrolidinylthiosemicarbazone (HL³). 5-(morpholinomethyl)pyridine-2-carboxaldehyde (0.29 g, 1.41 mmol) and *N*-pyrrolidine-3-thiosemicarbazide (0.20 g, 1.41 mmol) were dissolved in EtOH (5 mL) and stirred in a Shlenk tube at 78 °C for 2 h until the solution became clear. The mixture was left to stand at 4 °C for 12 h. The yellow crystalline precipitate was filtered off, washed with cold EtOH and dried in vacuo. Yield: 0.36 g. ¹H NMR (500 MHz, DMSO) δ 11.23 (s, 1H, NH¹⁶), 8.49 (d, *J* = 1.4 Hz, 1H, H³), 8.20 (s, 1H, H⁶), 8.01 (dd, *J* = 8.0, 2.1 Hz, 4H, H²¹, H²²), 7.84 (s, 1H, H¹⁴), 7.77 (d, *J* = 1.9 Hz, 1H, H⁴), 3.76 (s, 4H, H²⁰, H²³), 3.59 (dd, *J* = 8.1, 5.3 Hz, 4H, H⁹, H¹³), 3.52 (s, 2H, H⁷), 2.42 – 2.32 (m, 4H, H¹⁰, H¹²). ¹³C NMR (126 MHz, DMSO) δ 177.02 (C, C¹⁷), 153.11 (C, C²), 150.17 (CH, C⁶), 149.02 (CH, C³), 143.48 (CH, C¹⁴), 139.44, (2CH₂, C²¹, C²²), 138.08 (CH, C¹⁶), 134.05 (C, C⁵), 119.43 (CH, C³), 66.62 (2CH₂, C⁹, C¹³), 59.91 (CH₂, C⁷), 53.53 (2CH₂, C²⁰, C²³), 40.00 (2CH₂, C¹⁰, C¹², overlapped with residual DMSO signal). IR (ATR, selected bands, $\tilde{\nu}_{\text{max}}$): 1585, 1529, 1419, 1347, 1287, 1108, 1001, 855, 796, 741 cm⁻¹.

5-(morpholinomethyl)pyridine-2-carboxaldehyde *N*-morpholinylthiosemicarbazone (HL⁴). 5-(morpholinomethyl)pyridine-2-carboxaldehyde (0.29 g, 1.41 mmol) and *N*-morpholinyl-3-thiosemicarbazide (0.23 g, 1.41 mmol) were dissolved in EtOH (5 mL) and stirred in a Shlenk tube at 78 °C for 2.5 h until the solution became clear. Light-yellow crystals formed within 3 weeks at 4 °C after concentration of the reaction mixture. These were filtered off, washed with cold EtOH and dried in vacuo. Yield: 0.12 g. ¹H NMR (500 MHz, DMSO) δ 11.43 (s, 1H, NH¹⁶), 8.50 (d, *J* = 1.3 Hz, 1H, H⁶), 8.18 (s, 1H, H¹⁴), 7.83 (d, *J* = 8.1 Hz, 1H, H⁴), 7.78 (dd, *J* = 8.2, 1.9 Hz, 1H, H³), 3.98 – 3.91 (m, 4H, H²⁰, H²⁴), 3.72 – 3.66 (m, 4H, H²¹, H²³), 3.60 – 3.55 (m, 4H, H⁹, H¹³), 3.52 (s, 2H, H⁷), 2.36 (br. s, 4H, H¹⁰, H¹²). ¹³C NMR (126 MHz, DMSO) δ 181.27 (C_q, C¹⁷), 152.75 (C_q, C⁴), 150.33 (CH, C⁶), 144.44

(CH, C¹⁴), 137.83 (CH, C³), 134.27 (CH, C²), 119.60 (C_q, C⁴), 66.60 (2CH₂, C²¹, C²³), 66.48 (CH₂, C⁷), 59.73 (2CH₂, C⁹, C¹³).

5-(morpholinomethyl)pyridine-2-carboxaldehyde 4-phenylthiosemicarbazone (HL⁵). 5-

(morpholinomethyl)pyridine-2-carboxaldehyde (0.29 g, 1.41 mmol) and 4-phenyl-3-thiosemicarbazide (0.24 g, 1.41 mmol) were dissolved in EtOH (5 mL) and stirred in a Shlenk tube at 78 °C for 3 h until the solution became clear. Then the reaction mixture was allowed to stand at 4 °C for 12 h. The yellow crystalline precipitate was filtered off, washed with cold EtOH and dried in vacuo. Yield: 0.32 g. ¹H NMR (500 MHz, DMSO) δ 12.02 (s, 1H, NH¹⁶), 10.24 (s, 1H, NH¹⁸), 8.52 (s, 1H, H⁶), 8.41 (d, *J* = 8.1 Hz, 1H, H³), 8.21 (s, 1H, H¹⁴), 7.78 (d, *J* = 10.1 Hz, 1H, H⁴), 7.56 (d, *J* = 8.6 Hz, 2H, H²¹, H²⁵), 7.40 (t, *J* = 7.9 Hz, 2H, H²², H²⁴), 7.24 (t, *J* = 7.4 Hz, 1H, H²³), 3.59 (s, 4H, H¹⁰, H¹²), 3.54 (s, 2H, H⁷), 2.38 (s, 4H, H⁹, H¹³). ¹³C NMR (126 MHz, DMSO) δ 176.84 (C_q, C¹⁷), 152.62 (C_q, C²), 150.25 (CH, C⁶), 143.45 (C_q, C¹⁴), 139.44 (C_q, C²⁰), 137.51 (CH, C⁴), 134.46 (C_q, C⁵), 128.59 (2CH, C²², C²⁴), 126.56 (2CH, C²¹, C²⁵), 120.68 (CH, C³), 66.62 (2CH₂, C¹⁰, C¹²), 53.54 (2CH₂, C⁹, C¹³), 40.00 (CH₂, C⁷). IR (ATR, selected bands, $\tilde{\nu}_{\text{max}}$): 1592, 1505, 1466, 1338, 1291, 1188, 1105, 1060, 1000, 944, 909, 863, 791, 751, 689, 634 cm⁻¹.

5-(morpholinomethyl)pyridine-2-carboxaldehyde S-methylisothiosemicarbazone (HL⁶). To a solution of 5-(morpholinomethyl)pyridine-2-carboxaldehyde (0.29 g, 1.41 mmol) in EtOH (6 mL) was added a solution of S-methylisothiosemicarbazide hydroiodide (0.33 g, 1.41 mmol) in water (2 mL). Then the reaction mixture was heated to 60 °C and a solution of NaHCO₃ (0.118 g, 1.41 mmol) in water (4 mL) was added dropwise. When bubbles of CO₂ disappeared, the mixture was allowed to cool to room temperature. The solvent was removed under reduced pressure and the residue dissolved in water. The product was extracted with chloroform. After removal of the solvent the residue was dissolved in a small amount of EtOH and allowed to stand at -20 °C overnight. Light-brown crystalline product was obtained the next day. Yield: 0.14 g. ¹H NMR (500 MHz, DMSO) δ 8.49 (d, *J* = 1.4 Hz, 1H, H⁴), 8.21 (d, *J* = 8.1 Hz, 1H, H¹⁴), 8.17 (s, 1H, H⁶), 7.84 (d, *J* = 8.1 Hz, 1H, H³), 7.16 (s, 2H, H¹⁸), 3.60-3.55 (m,

8H, H⁹, H¹³), 3.51 (s, 2H, H⁷), 2.40 (br. s, 3H, H²⁰), 2.36 (d, J = 6.0 Hz, 8H, H¹⁰, H¹²). ¹³C NMR (126 MHz, DMSO) δ 164.19 (C, C¹⁷), 154.02 (C, C⁵), 151.84 (CH, C⁶), 150.12 (CH, C⁴), 137.35 (CH, C³), 133.89, (C, C²), 120.68 (CH, C¹⁴), 66.61 (2CH₂, C⁹, C¹³), 59.84 (CH₂, C⁷), 53.53 (2CH₂, C¹⁰, C¹²), 12.52 (CH₃, C²⁰). IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1600, 1497, 1472, 1335, 1290, 1111, 1069, 1015, 931, 870, 790, 752, 714, 626 cm⁻¹.

3. Synthesis of Cu(II) complexes.

[Cu(L¹)Cl] (1). To **HL¹** (100 mg, 0.36 mmol) in MeOH (20 mL) was added triethylamine (50 μ l, 0.36 mmol) and then CuCl₂·2H₂O (60 mg, 0.36 mmol) in MeOH (5 mL) at 60 °C. The solution was stirred for 5 min and left to stand at room temperature overnight. The precipitate formed was filtered off, washed with cold EtOH and dried in air. Yield: 123 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1631, 1426, 1310, 1158, 1107, 1060, 907, 722, 688 cm⁻¹.

[Cu(L²)Cl] (2). To **HL²** (100 mg, 0.32 mmol) in MeOH (20 mL) was added triethylamine (44.5 μ L, 0.32 mmol) and then CuCl₂·2H₂O (54 mg, 0.32 mmol) in MeOH (5 mL) at 60 °C. The solution was stirred for 5 min and left to stand at room temperature overnight. The precipitate formed was filtered off, washed with cold EtOH and dried in air. Yield: 104 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1505, 1372, 1310, 1252, 1111, 1000, 913, 872, 749, 626 cm⁻¹.

[Cu(L³)Cl] (3). To **HL³** (100 mg, 0.3 mmol) in MeOH (20 mL) was added triethylamine (42 μ l, 0.28 mmol) and then CuCl₂·2H₂O (50 mg, 0.3 mmol) in MeOH (5 mL) at 60 °C. The solution was stirred for 5 min and left to stand at room temperature overnight. The precipitate formed was filtered off, washed with cold EtOH and dried in air. Yield: 74 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1444, 1375, 1281, 1242, 1111, 1004, 913, 885, 623 cm⁻¹.

[Cu(L⁴)Cl] (4). To **HL⁴** (50 mg, 0.28 mmol) in MeOH (20 mL) was added triethylamine (20 μ l, 0.28 mmol) and then CuCl₂·2H₂O (25 mg, 0.28 mmol) in MeOH (5 mL) at 60 °C. The solution was stirred for 5 min and left to stand at room temperature overnight. The precipitate formed was filtered off, washed

with cold EtOH and dried in air. Yield: 22 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1465, 1429, 1381, 1313, 1265, 1229, 1109, 1028, 932, 888, 860 cm^{-1} .

[Cu(L⁵)Cl] (5). To **HL⁵** (100 mg, 0.28 mmol) in MeOH (20 mL) was added triethylamine (39 μl , 0.28 mmol) and then $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$ (47 mg, 0.82 mmol) in MeOH (5 mL) on heating. The solution was stirred for 5 min and left to stand at room temperature overnight. The precipitate formed was filtered off, washed with cold EtOH and dried in air. Yield: 67 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1600, 1550, 1493, 1417, 1182, 1111, 1000, 899, 863, 799, 752, 691 cm^{-1} .

[Cu(L⁶)Cl] (6). To **HL⁶** (100 mg, 0.34 mmol) in MeOH (20 mL) was added triethylamine (47 μl , 0.34 mmol). Then the reaction was heated to 40 °C and $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$ (58 mg, 0.34 mmol) in MeOH (5 mL) was added. The solution was stirred for 5 min at 50 °C and left to stand at –20 °C overnight. The precipitate formed was filtered off, washed with ethanol, ether and dried in air. Yield: 75 mg. IR (ATR, selected bands, $\tilde{\nu}_{\max}$): 1599, 1491, 1431, 1320, 1115, 1035, 1004, 973, 911, 796, 724 cm^{-1} .

4. Solution equilibrium studies in aqueous phase

pH-potentiometric measurements. These measurements were performed as described previously.¹⁶ The initial volume of the samples was 10.0 mL. The ligand concentration was 1 mM and metal ion-to-ligand ratios of 1:1 – 1:3 were used. The exact concentration of the ligand stock solutions together with the proton dissociation constants were determined by pH-potentiometric titrations with the use of the computer program HYPERQUAD.⁷⁹ HYPERQUAD was also utilised to establish the stoichiometry of the complexes and to calculate the stability constants ($\log \beta(\text{M}_p\text{L}_q\text{H}_r)$). $\beta(\text{M}_p\text{L}_q\text{H}_r)$ is defined for the general equilibrium $p\text{M} + q\text{L} + r\text{H} \rightleftharpoons \text{M}_p\text{L}_q\text{H}_r$ as $\beta(\text{M}_p\text{L}_q\text{H}_r) = [\text{M}_p\text{L}_q\text{H}_r]/[\text{M}]^p[\text{L}]^q[\text{H}]^r$, where M denotes the metal ion and L the completely deprotonated ligand. The uncertainties (standard deviations, SD) of the equilibrium constants are shown in parentheses for the species determined in the present work.

5. Instrumentation. ICP-OES determination of Cu content was performed in Chemical, Molecular and Analysis Centre, National University of Singapore with Optima ICP-OES (Perkin Elmer, Waltham, MA, USA). The absorbance of thiazolyl blue tetrazolium bromide (MTT) was measured by synergy H1 hybrid multimode microplate reader (Bio-Tek, Winoosky, VT, USA). Cu and Re contents in cells were determined by Agilent 7700 Series ICP-MS (Agilent Technologies, Santa Clara, CA, USA). Flow cytometry was performed on BD LSRFortessa Cell Analyzer (BD Biosciences, Franklin Lakes, NJ, USA). Western blot images were generated from G:Box (Syngene, Cambridge, UK). The UV-vis spectrophotometric measurements were performed on a Hewlett Packard 8452A diode array spectrophotometer and a Thermo Scientific Evolution 220 spectrophotometer. CW-EPR spectra were recorded with a BRUKER EleXsys E500 spectrometer. *In situ* ultraviolet-visible-near-infrared (UV-vis-NIR) spectroelectrochemical measurements were performed on a spectrometer (Avantes, Model AvaSpec-2048x14-USB2).

6. UV-vis spectrophotometric, ¹H NMR, EPR and lipophilicity measurements. UV-vis spectra were recorded in the ranges 200–800 nm and 450–1050 nm, respectively. The path length was 0.5, 1 or 2 cm. Stability constants of the complexes and the molar absorbance spectra of the individual species were calculated with the computer program PSEQUAD.⁸⁰ The spectrophotometric titrations were performed on samples containing the proligand with or without Cu(II) ions and the concentration of the proligand was 120 μM – 1.4 mM. The metal-to-proligand ratios were 1:1 and 1:2 in the pH range from 1.0 to 11.5 at 25.0 ± 0.1 °C at an ionic strength of 0.10 M (KCl). pH values in the range 1.0–2.0, were calculated from the strong acid and strong base content. The conditional stability constant of [CuL] at pH 6.0 (50 mM MES) for **1** was determined from competition titrations of the Cu(II) complex of EDTA with the proligand **HL**¹. Samples contained 34 μM Cu(II) ion and 34 μM **HL**¹, and the concentration of the EDTA was varied in the range of 0–83 μM. Absorbance data were recorded after 0.5 h incubation. ¹H NMR studies for **HL**¹ were carried out on a Bruker Avance III HD Ascend 500 Plus instrument in a 10% (v/v) D₂O/H₂O mixture at ionic strength of 0.10 M (KCl). All CW-EPR spectra were recorded with a

BRUKER EleXsys E500 spectrometer (microwave frequency 9.85 GHz, microwave power 10 mW, modulation amplitude 5 G, modulation frequency 100 kHz. The pH dependent series of isotropic EPR spectra were recorded in a circulating system, at room temperature. A Heidolph Pumpdrive 5101 peristaltic pump was used for circulate the solution from the titration pot through a capillary tube into a Bruker flat cell placed in the cavity of the instrument. The titrations were carried out under nitrogen atmosphere. EPR spectra were recorded at 1.00 mM Cu(II) and 0.75 mM ligand concentration, and at 1.00 mM CuCl₂ and 1.50 mM ligand concentration, both between pH = 1 – 11.5. The ionic strength of 0.1 M was adjusted with KCl. Before the simulation of the room temperature spectra, the measured spectra were corrected by subtracting the spectra of aqua solution measured in the same circulating system. The series of pH-dependent isotropic EPR spectra recorded in the equimolar solution were simulated by the „two-dimensional” method using the 2D_EPR program⁸¹ and EPR parameters were computed as published previously.^{16,25} Distribution coefficients ($D_{7.4}$) values of complexes **1–6** and **HL¹–HL⁶** were determined by the traditional shake-flask method in *n*-octanol/buffered aqueous solution at pH 7.40 (20 mM HEPES, 0.10 M KCl) at 25.0 ± 0.2 °C as described previously.⁸²

7. Crystallographic Structure Determination. X-ray diffraction quality single crystals of **HL¹**, **HL²**, **HL⁴–HL⁶** were obtained by recrystallization in ethanol, while **1** and **3–6** by slow diffusion of diethyl ether into the DMF and methanolic solution of the complexes, respectively. The measurements were performed on a Bruker X8 APEX-II CCD (**HL¹**, **HL²**, **HL⁴**, **HL⁵**, **1'**, **3–6**), Bruker D8 Venture (**1**) or Gemini (**HL⁶**) diffractometer. Single crystals were positioned at 35, 35, 35, 35 and 55 mm from the detector, and 388, 1110, 946, 1108 and 2935 frames were measured, each for 40, 3, 60, 3 and 32s over 0.5, 0.5, 0.5, 0.5 and 1.0° scan width for **HL¹–HL⁶**, respectively. For Cu(II) complexes **1**, **1'**, **3–6** the single crystals were placed at 24, 35, 35, 33, 35 and 24 mm from the detector, and 8088, 2692, 794, 1212, 1261 and 567 frames were measured, each for 3, 2, 60, 60, 60 and 3s over 0.6, 0.5, 0.5, 0.5, 0.4 and 0.5° scan width, respectively. The data were processed using SAINT or CrysAlis software.^{83,84} Crystal data, data collection parameters, and structure refinement details are given in Tables S11 and S12. The

structures were solved by direct methods and refined by full-matrix least-squares techniques. Non-H atoms were refined with anisotropic displacement parameters. H atoms were inserted in calculated positions and refined with a riding model. The morpholine group and the N-piperidinyl unit in one of three crystallographically independent molecules in the asymmetric unit were found to be disordered over two positions with s.o.f. 0.5:0.5 and 0.6:0.4 in **3·0.25MeOH**. The morpholine moiety attached to pyridine unit in one of the three crystallographically independent molecules of **4·0.58MeOH** was found to be disordered over two positions with s.o.f. 0.7:0.3, while one molecule of methanol over two positions with s.o.f. 0.75:0.25. The positional parameters of disordered atoms were refined by using PART, DFIX and SADI tools implemented in SHELX. The following computer programs and hardware were used: structure solution, *SHELXS-2014* and refinement, *SHELXL-2014*;⁸⁵ molecular diagrams, ORTEP;⁸⁶ computer, Intel CoreDuo. CCDC 1850567–1850577.

8. Electrochemistry and spectroelectrochemistry. Cyclic voltammetric experiments were performed as described previously.⁵³ The cyclic voltammograms were measured in the cathodic region in different solvents (DMSO, methanol, DMSO/H₂O). The analytical purity grade LiClO₄ (Sigma-Aldrich) and distilled and deionized water were used for preparation of 1 mM aqueous solutions of the investigated complexes **1–6**. EPR spectra were recorded with the EMX plus. *In situ* ultraviolet-visible-near-infrared (UV–vis–NIR) spectroelectrochemical measurements were performed on a spectrometer (Avantes, Model AvaSpec-2048x14-USB2 in the spectroelectrochemical cell kit (AKSTCKIT3) with the Pt-microstructured honeycomb working electrode, purchased from Pine Research Instrumentation. The cell was positioned in the CUV–UV Cuvette Holder (Ocean Optics) connected to the diode-array UV–vis–NIR spectrometer by optical fibers. UV–vis–NIR spectra were processed using the AvaSoft 7.7 software package. Halogen and deuterium lamps were used as light sources (Avantes, Model AvaLight-DH-S-BAL). The *in situ* EPR spectroelectrochemical experiments were carried out under an argon atmosphere in the EPR flat cell equipped with a large platinum mesh working electrode. The freshly prepared solutions were carefully purged with argon and the electrolytic cell was polarized in the

galvanostatic mode directly in the cylindrical EPR cavity TM-110 (ER 4103 TM) and the EPR spectra were measured in situ.

9. Cell lines and culture conditions. Human ovarian carcinoma cells A2780 and A2780cisR, and human embryonic kidney HEK293 were obtained from ATCC. A2780 and A2780cisR cells were cultured in RPMI 1640 medium containing 10% fetal bovine serum (FBS). HEK293 were cultured in DMEM medium containing 10% FBS. All cells were grown in tissue culture 25 cm² flasks (BD Biosciences, Singapore) at 37 °C in a humidified atmosphere of 95% air and 5% CO₂. All drug stock solutions were prepared in sterile water. The amount of Cu was determined by ICP-OES. All compounds were soluble in water; more specifically, the solubility of **1**, **2** and **5** was moderate and did not exceed 2 mM, whereas **3** was less soluble (0.3 mM) in water.

10. Inhibition of cell viability assay. The cytotoxicity of the compounds was determined by colorimetric microculture assay (MTT assay, MTT = 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide) as described previously.⁵³

11. Cellular accumulation. Cellular accumulation of **1–6** was determined in A2780 cells. Cells were seeded into Cellstar 6-well plates (Greiner Bio-one) at a density of 60×10^4 cells/well (2 mL per well). After the cells were allowed to resume exponential growth for 24 h, they were exposed to **1–6** at 1 μM for 24 h at 37 °C. The cells were washed twice with 1 mL of PBS and lysed with RIPA lysis buffer for 5–10 min at 4 °C. The cell lysates were scraped from the wells and transferred to separate 1.5 mL microtubes. The supernatant was then collected after centrifugation (13 000 rpm, 4 °C for 15 min) and total protein content of each sample was quantified via Bradford's assay. Cell lysates were transferred to 2 mL glass vials and then digested with ultrapure 65% HNO₃ at 100 °C for 24 h. The resulting solution was diluted to 1 mL (2–4% v/v HNO₃) with ultrapure Milli-Q water. Cu content of each sample was quantified by ICP-MS. In was used as an internal standard. Cu and In were measured at *m/z* 64 and *m/z* 115, respectively. Metal standards for calibration curve (0, 0.5, 1, 2, 5, 10, 20, 40 ppb) were freshly prepared before each measurement. All readings were made in triplicates in He mode. For temperature-

dependent cellular accumulation experiments, A2780 cells were seeded into 35×10 mm tissue culture plates at a density of 60×10⁴ cells/plate. After the cells were allowed to resume exponential growth for 24 h, they were exposed to **2** at 3 μM for 10, 30, 60 and 120 min at 37 and 4 °C. The subsequent analysis was performed as described above. For energy-dependent experiments, A2780 cells were seeded into 35×10 mm tissue culture plates at a density of 1×10⁶ cells/plate. After the cells were allowed to resume exponential growth for 24 h, they were pre-incubated with oligomycin (5 μM) for 1 h or 4 h or with cycloheximide (100 μM) for 4 h and further co-incubated with 3 μM of **2** at 37 °C. To induce total starvation, after the cells were allowed to resume exponential growth for 24 h, RPMI media was replaced with Hanks' Balanced Salt Solution (HBSS) and cells were pre-incubated in HBSS for 1 h and further incubated with 3 μM of **2** for 1 h at 37 °C. The subsequent analysis was performed as described above.

12. Western blot analysis. A2780 cells were seeded into Cellstar 6-well plates (Greiner Bio-One) at a density of 60 × 10⁴ cells/well (2 mL per well). After the cells were allowed to resume exponential growth for 24 h, they were exposed to **HL**² and **2** at different concentrations for 24 h. The experiment was performed essentially as described previously.⁵³ The membranes were blocked in 5% BSA (w/v) in TBST wash buffer for 1 h and subsequently incubated with the appropriate primary antibodies in 5% BSA (w/v) in TBST wash buffer (actin antibody) at 4 °C overnight. The membranes were washed with a wash buffer 3 times for 5 min. After incubation with horseradish peroxidase-conjugated secondary antibodies (room temperature, 1.5 h), the membranes were washed with a wash buffer 4 times for 5 min. Immune complexes were detected with Luminata HRP substrates and analyzed using enhanced chemiluminescence imaging. Actin was used as a loading control. The following antibodies were used: NRF2 (sc13032) from Santa Cruz Biotechnologies, ECL Antirabbit IgG (NA934 V) and ECL Antimouse IgG (NA931) from GE Healthcare Life Sciences, cleaved PARP (Asp214) (D64E10), PARP, CHOP (D46F1), BiP (C50B12), IRE1α (14C10), β-actin (13E5), phospho-p44/p42 MAPK (Erk1/2) (Thr202/Tyr204) (D13.14.4E), cyclin D1 (92G2), cyclin B1 (D5C10), XIAP antibodies from Cell

Signalling Technologies. All antibodies were used at 1 : 500 dilutions except for actin (1 : 10 000), anti-mouse and anti-rabbit (1 : 5000).

13. Annexin V/PI apoptosis assay. A2780 cells were seeded into Cellstar 12-well plates (Greiner Bio-One) at a density of 20×10^4 cells/well (1 mL per well). The cells were allowed to resume exponential growth for 24 h and subsequently they were exposed to **HL**² and **2** at different concentrations for 24 h. After the supernatant solution was collected in 1.5 mL microtubes, the cells were washed with 100 μ L of trypsin, which was combined with the supernatant. Subsequently, cells were trypsinized with 200 μ L of trypsin for 5 min at 37 °C, 5% CO₂, washed with 200 μ L of PBS and combined with the supernatant. The cells were centrifuged at 2.5×10^3 rpm for 5 min and the pellets were washed once with PBS and resuspended in 500 μ L of Annexin V binding buffer and stained with Annexin V-FITC and PI reagents. The fluorescence was immediately analyzed by flow cytometry. The resulting dot blots were acquired from 10 000 events and quantified using Flowjo software (Flowjo LLC, Ashland, OR, USA).

14. Tyrosyl radical reduction in mouse R2 ribonucleotide reductase protein. The 9.4 GHz EPR spectra were recorded at 30 K on a Bruker EleXsys II E540 EPR spectrometer with an Oxford Instruments ESR900 helium cryostat, essentially as described previously.^{Error! Bookmark not defined.} The small mouse R2 subunit without diferric-tyrosyl cofactor was produced from *E.coli* carrying a mouse R2 cDNA plasmid. The cofactor was further regenerated, resulting in the formation of the cluster with 0.38 tyrosyl radical/Fe(III) ratio, which is in agreement with the literature.⁴¹

15. Molecular docking calculations. The calculations were performed as described previously.²⁰ The centre of the binding pocket was defined ($x = 102.276$, $y = 87.568$, $z = 80.588$)⁸⁷ with 10 Å radius. The basic amino acids lysine and arginine were defined as protonated. Furthermore, aspartic and glutamic acids were assumed to be deprotonated. The GoldScore (GS),³⁷ ChemScore (CS),^{88,89} Chem Piecewise Linear Potential (ChemPLP)⁹⁰ and Astex Statistical Potential (ASP)⁹⁰ scoring functions were implemented to validate the predicted binding modes and relative energies of the ligands using the GOLD v5.4 software suite. The parameter file for GS was augmented for Cu according to Sciortino et

al.³⁸ The QikProp 4.6⁹¹ and Marvin software package⁹² was used to calculate the molecular descriptors of the compounds. The reliability of QikProp is established for the molecular descriptors.⁹³

16. Bacterial Strains and antibacterial activity. *Wild-type Pseudomonas aeruginosa PAO1 strain CECT 4122 (ATCC 15692) and Staphylococcus aureus CECT 86 (ATCC 12600)* were obtained from the Spanish Type Culture Collection (CECT). All strains were routinely cultivated in TSB medium (Sharlab, Spain) at 37 °C. Minimal inhibitory concentration (MIC) assays were determined by the microdilution method using TSB broth following the method described by the Clinical and Laboratory standards Institute.⁹⁴ In brief, compounds were diluted in a 96-well microtiter plate (tissue culture-treated polystyrene; Costar 3595, Corning Inc., Corning, NY) to a final concentration ranging from 0.1 to 100 µg/mL. A 100-µl aliquot of the bacterial suspension (around 5×10^5 colony forming units/mL (CFU/mL)) was inoculated, incubated at 37 °C for 8 h at 150 rpm, and absorbance at 550 nm was read every 15 min in an Infinity 200 Pro microplate reader (Tecan). The MIC₁₀₀ was determined as the lowest concentration that completely inhibited bacterial growth, and MIC₅₀ in which bacterial growth was inhibited at 50%.

17. Bacterial viability test analysis. Cultures of *S. aureus* and *P. aeruginosa* were diluted in fresh TSB medium and grown overnight to the beginning of exponential phase (A₅₅₀ 0.3) and different compounds were added. After 3 h of incubation at 37 °C in shaking conditions, cells were harvested and stained using the LIVE/DEAD BactLight Bacterial Viability Kit (Thermofisher) for 30 min. Fluorescent bacteria were visualized by a Nikon inverted fluorescent microscope ELIPSE Ti-S/L100 (Nikon) coupled with a DS-Qi2 Nikon camera.

ASSOCIATED CONTENT

Supporting Information Available: Synthetic details, atom numbering scheme for NMR spectra (Scheme 1), ORTEP views of **HL**¹, **HL**², **HL**⁴⁻⁶ (Figure S1), portion of the crystal structure of **6** (Figure S2), low field region of the ¹H NMR spectra of **HL**¹ recorded at pH 0.75–12.50 (Figure S3),

concentration distribution curves of **HL**¹ (Figure S4), UV–vis and EPR spectra for Cu(II) containing systems (Figures S5–S7), cyclic voltammograms and UV–vis spectra of **1–6** and one of the proligands (Figures S8–S11), plot with potential dependence of UV–vis spectra of **2** (Figure S12), concentration–effect curves (Figure S13), hydrogen binding of **5** to R2 protein (Figure S14), EPR spectrum of R2 RNR tyrosyl radical, complex formation reactions of Fe^{III} and Fe^{II} with **HL**² and **HL**⁵ monitored by UV–vis spectroscopy (Figure S15), ESI MS spectrum of [Fe^{III}(**HL**⁵)]⁺ (Figure S16), selected bond distances and angles in Cu(II) complexes (Tables S1 and S2), proton dissociation constants for **HL**¹ (Table S3), cumulative stability and proton dissociation constants of Cu(II) complexes of **HL**¹ (Table S4), formation constants for Cu(II)–**HL**¹ complexes (Table S5), electrochemical data for **1–6** (Table S6), Molecular descriptors for **HL**¹–**HL**⁶ and **1–6** (Table S7), details of molecular docking calculations (Table S8), analytical data (Tables S9 and S10), details of data collection and refinement of proligands (Table S11) and Cu(II) complexes (Table S12). Molecular formula strings (CSV). This material is available free of charge via the Internet at <http://pubs.acs.org>.

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ABBREVIATIONS

TSC, thiosemicarbazone, COTI-2, (*E*)-*N'*-(6,7-dihydroquinolin-8(5*H*)-ylidene)-4-(pyridine-2-yl)piperazine-1-carbothiohydrazide, DpC, di-2-pyridylketone 4-cycloheptyl-4-methyl-3-thiosemicarbazone, ROS, reactive oxygen species, GSH, glutathione, UPR, unfolded protein response, TEMED, tetramethylethylenediamine, PI, propidium iodide, FBS, Foetal Bovine Serum, BSA, Bovine Serum Albumine, HBSS, Hank's Balanced Salt Solution, PS, phosphatidylserine, KDS, known drug space.

Accession Codes

PDB ID of the complexes of Triapine and **5** docked in mouse R2 protein: 1w68-5-T. Authors will release the atomic coordinates upon article publication.

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Table of Contents graphic

