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## An enolato-bridged dinuclear Cu(II) complex with a coumarin-assisted precursor: a spectral, magnetic and biological study<sup>†</sup>

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A new, phenoxy-bridged Cu<sup>II</sup> dinuclear complex  $\text{Cu}_2[(\text{L})_2(\text{DMF})_2]$  (**1**) has been obtained by employing the coumarin-assisted tridentate precursor,  $\text{H}_2\text{L}$ , [benzoic acid(7-hydroxy-4-methyl-2-oxo-2H-chromen-8-ylmethylene)-hydrazide]. Complex **1** has been systematically characterized by FTIR, UV-Vis, fluorescence and EPR spectrometry. The single crystal X-ray diffraction analysis of **1** shows that the geometry around each copper ion is square pyramidal, comprising two enolato oxygen atoms belonging to different ligands (which assemble the dimer bridging the two metal centers), one imine-N and one phenolic-O atoms of the Schiff base and one oxygen atom from the DMF molecule. The temperature dependent magnetic interpretation agrees with the existence of weak ferromagnetic interactions between the bridging dinuclear Cu(II) ions. Both the ligand and complex **1** exhibit anti-mycobacterial activity and considerable efficacy towards *M. tuberculosis* H37Rv ATCC 27294 and *M. tuberculosis* H37Ra ATCC 25177 strains. The cytotoxicity study on human adenocarcinoma cell lines (MCF7) suggests that the ligand and complex **1** have potential anticancer properties. Molecular docking of  $\text{H}_2\text{L}$  with the enoyl acyl carrier protein reductase of *M. tuberculosis* H37Rv (PDB ID: 4U0K) is examined and the best docked pose of  $\text{H}_2\text{L}$  shows one hydrogen bond with Thr196 (1.99 Å).

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## Introduction

Recent research trends reveal that there has been considerable interest among researchers in studying the chemistry of coumarin functionalised compounds due to their broad spectrum of activities ranging from photophysical or photochemical properties<sup>1</sup> to biological properties.<sup>2</sup> Many products that contain

a coumarin subunit exhibit diverse biological activities such as molluscicidal<sup>3a</sup> and insecticidal activities,<sup>3b</sup> cytotoxicity,<sup>3c</sup> and anti-HIV activity,<sup>3d</sup> some serve as antioxidant<sup>3c</sup> and anticoagulant agents.<sup>4</sup> Basically, the coumarin nucleus is one of the most interesting chromophores with high fluorescence efficiency,<sup>5</sup> which opens up a new way for recognition and determination of metal cations or anions.<sup>6</sup> There are several coumarin appended fluorescent sensors reported in the literature which are potentially used to detect cations *e.g.*  $\text{Ca}^{2+}$ <sup>6a</sup>  $\text{Zn}^{2+}$ <sup>6b</sup> and  $\text{Cu}^{2+}$ <sup>6c</sup> as well as anions *e.g.* fluorides,<sup>6d</sup> cyanides, *etc.* It is well established that the biological activity associated with hydrazone compounds is attributed to the presence of the active pharmacophore ( $-\text{CONH}-\text{N}=\text{C}-$ ),<sup>7</sup> and many hydrazone compounds containing this active moiety showed good anticancer, anti-tuberculosis and antibacterial activities.<sup>8</sup> Additionally, the azomethine ( $-\text{CONH}-\text{N}=\text{C}-$ ) group in hydrazone derivatives possesses strong binding ability due to the formation of stable chelates with metal ions. So, coumarin containing hydrazone derivatives are not only expected to have enhanced biological activities but also show high sensitivity towards metal ions. Indeed, chelation studies involving metal ions coordinated to coumarin derivatives at different positions have been reported in the literature.<sup>9</sup> 8-Substituted-4-methyl-7-hydroxycoumarin

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and 6-substituted-4-methyl-7-hydroxycoumarin have been investigated due to their complexation ability and exhibited anticoagulant and plant growth regulating properties.<sup>10</sup> Some of us previously reported one Cu(II) derivative mediated by *N*[(2-pyridyl)methylidene]-6-coumarin.<sup>11</sup>

Dinuclear transition metal complexes have gained significant interest over the years due to their diverse properties and reactivity, originating from the magnetic exchange occurring between neighbouring metal centres and potential applications in the field of magnetism<sup>12</sup> and biochemistry.<sup>13</sup> So far, several copper ions coordinated to oxygen and/or nitrogen atoms as structural and functional models in magneto-chemistry and metalloenzymes have been reported in the literature.<sup>14</sup> Copper is the third most abundant transition metal found in living systems,<sup>15</sup> and it plays an essential role in biological systems. The nature (antiferromagnetic or ferromagnetic) and magnitude of a magnetic exchange interaction between two metal ions are reasonably well understood in terms of the energetics and overlap of 'magnetic' orbitals.<sup>16</sup> In the last few years, the success in understanding magnetic exchange interactions in metal complexes has prompted efforts to see if these complexes can be used as molecular building blocks for materials exhibiting interesting properties. The magnetic spin exchange in dinuclear copper(II) complexes in relation to the participation of donor atoms of the corresponding ligand is of continuing interest.<sup>17</sup> The design of ligands capable of binding two metal ions in close proximity is, therefore, an important subject in this field. In order to gain an insight into the correlation between the structure and magnetism of dinuclear copper(II) complexes, detailed investigations based on crystal structure analyses are needed. Tuberculosis, which is caused primarily by *Mycobacterium tuberculosis* and rarely by *M. bovis* and *M. africanum*, is one of the oldest disease in the world affecting lungs and spreading to the other organs.<sup>18</sup> At present, a million children per year have died from this disease.<sup>19</sup> Mycobacteria resist to many chemicals, disinfectant antibiotics and chemotherapeutic agents.<sup>20</sup> Mycobacteria show a strong resistance to most of the antibiotics and chemicals due to the low permeability of the mycobacterial cell wall. Porins of the thickness and low permeability of this bilayer membrane allow inefficient permeability of the substances.<sup>20b</sup> The prospect of developing new and effective drugs is central to the global control of tuberculosis.

In the present contribution, the ligand [benzoic acid(7-hydroxy-4-methyl-2-oxo-2*H*-chromen-8-ylmethylene)-hydrazide] ( $H_2L$ ), a tridentate ONO donor,<sup>21</sup> was used in order to synthesize the Cu(II) dinuclear complex  $Cu_2[(L)_2(DMF)_2]$  (1). Besides studying the systematic spectral and magnetic properties, we have focused our attention on the anti-mycobacterial activity of human tuberculosis cells like *M. tuberculosis* H37Rv ATCC 27294 and *M. tuberculosis* H37Ra ATCC 25177 strains. We also have examined the cytotoxicity study on human adenocarcinoma cell lines (MCF7). The anti-microbacterial efficiency of  $H_2L$  is examined by molecular docking with the enoyl acyl carrier protein reductase of *M. tuberculosis* H37Rv (PDB ID: 4U0K).

## Experimental

### Materials

All chemicals and solvents were of reagent grade and used as received.  $Cu(NO_3)_2 \cdot 3H_2O$  was purchased from Merck, India. Benzohydrazide and 7-hydroxy-4-methyl-coumarin were purchased from Aldrich Chemicals, India.

### Physical measurements

Microanalytical data (C, H, and N) were collected on a Perkin-Elmer 2400 CHNS/O elemental analyzer. FTIR spectra were recorded on a Perkin-Elmer RX-1 spectrophotometer in the range 4000–400  $cm^{-1}$  as KBr pellets. Electronic spectra were measured on a Lambda 25 (UV-vis-NIR) spectrophotometer. Emission spectra were recorded using an LS 55 Perkin-Elmer spectrofluorometer at room temperature (298 K) in different solutions under degassed conditions. The fluorescence quantum yield of the complexes was determined using carbazole as a reference with a known  $\phi_R$  of 0.42 in benzene.<sup>22</sup> The complex and the reference dye were excited at the same wavelength, maintaining a nearly equal absorbance ( $\sim 0.1$ ), and the emission spectra were recorded. The area of the emission spectrum was integrated using the software available in the instrument and the quantum yield was calculated according to the following equation:

$$\frac{\phi_S}{\phi_R} = \left[ \frac{A_S}{A_R} \right] \times \left[ \frac{(Abs)_R}{(Abs)_S} \right] \times \left[ \frac{\eta_S^2}{\eta_R^2} \right]$$

here,  $\phi_S$  and  $\phi_R$  are the fluorescence quantum yield of the sample and the reference, respectively.  $A_S$  and  $A_R$  are the area under the fluorescence spectra of the sample and the reference, respectively,  $(Abs)_S$  and  $(Abs)_R$  are the respective optical densities of the sample and the reference solution at the wavelength of excitation, and  $\eta_S$  and  $\eta_R$  are the values of the refractive index of the respective solvent used for the sample and the reference. Fluorescence lifetimes were measured using a time-resolved spectrofluorometer from IBH, UK. The instrument uses a picosecond diode laser (NanoLed-03, 370 nm) as the excitation source and works on the principle of time-correlated single photon counting.<sup>23</sup> The goodness of fit was evaluated by the  $\chi^2$  criterion and the visual inspection of the residuals of the fitted function to the data. EPR spectra were recorded from 0 to 10 000 Gauss in the temperature range 77–298 K using an X-band (9.4 GHz) Bruker EMX spectrometer. EPR parameters reported in the text were obtained by simulating the spectra using the computer program Bruker WinEPR SimFonia.<sup>24</sup> Electronic absorption spectra were recorded using a Perkin-Elmer Lambda 35 spectrophotometer. Magnetic properties were investigated using a Quantum Design MPMS-XL superconducting quantum interference device magnetometer (SQUID) at an applied field 0.5 T in a temperature range 5–300 K. The diamagnetic correction was carried out by using Pascal constants.

### Synthesis of the ligand and the complex

**Synthesis of the ligand ( $H_2L$ ).** 8-Formyl-7-hydroxy-4-methyl-coumarin was prepared from 7-hydroxy-4-methyl-coumarin using the reported procedure.<sup>25</sup> Ligand  $H_2L$  was synthesized by



Table 1 Crystallographic data of **1**

Empirical formula	C <sub>42</sub> H <sub>38</sub> Cu <sub>2</sub> N <sub>6</sub> O <sub>10</sub>
Formula weight	913.86
Temperature	293(2)
Wavelength (Å)	0.71069
Crystal system	Monoclinic
Space group	P21/n
<i>a</i> (Å)	8.2580(10)
<i>b</i> (Å)	11.221(2)
<i>c</i> (Å)	21.508(3)
$\beta$ (deg)	97.190(2)
Volume (Å <sup>3</sup> )	1977.3(5)
<i>Z</i>	2
<i>D</i> <sub>calc</sub> (mg m <sup>-3</sup> )	1.535
$\mu$ (MoK $\alpha$ ) (mm <sup>-1</sup> )	1.144
<i>F</i> (000)	940
Crystal size, mm <sup>-3</sup>	0.15 × 0.12 × 0.09
$\theta$ range for data collection	1.91–30.63
Reflections collected/unique	31462/6071 [ <i>R</i> (int) = 0.0358]
Observed reflections [ <i>F</i> <sub>o</sub> > 4 $\sigma$ ( <i>F</i> <sub>o</sub> )]	4850
Data/restraints/parameters	6071/0/314
Final <i>R</i> indices [ <i>F</i> <sub>o</sub> > 4 $\sigma$ ( <i>F</i> <sub>o</sub> )] <sup>b</sup>	<i>R</i> <sub>1</sub> = 0.0358, <i>wR</i> <sub>2</sub> = 0.0963
<i>R</i> indices (all data)	<i>R</i> <sub>1</sub> = 0.0483, <i>wR</i> <sub>2</sub> = 0.1040
Goodness-of-fit on <i>F</i> <sup>2a</sup>	1.046
Largest diff. peak and hole, e Å <sup>-3</sup>	0.616 and -0.251

<sup>a</sup> Goodness-of-fit *S* =  $[\sum w(F_o^2 - F_c^2)^2/(n - p)]^{1/2}$ , where *n* is the number of reflections and *p* is the number of parameters. <sup>b</sup> *R*<sub>1</sub> =  $\sum \|F_o\| - |F_c\| / \sum |F_o\|$ ; *wR*<sub>2</sub> =  $[\sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2]]^{1/2}$ .

refluxing 8-formyl-7-hydroxy-4-methyl-coumarin (0.408 g, 2 mmol) and benzohydrazide (0.272 g, 2 mmol) for a period of 2 h in methanol to obtain the yellow colored compound. It was filtered, washed with excess of methanol and recrystallized from dioxane (see Scheme ESI†). Yield: 0.231 g (72%). Anal. calc. for C<sub>18</sub>H<sub>14</sub>N<sub>2</sub>O<sub>4</sub>: C, 66.72; H, 4.91; N, 8.45. Found: C, 67.07; H, 4.38; N, 8.69%. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>):  $\delta$  2.4 (s, 3H, C18-CH<sub>3</sub>), 6.25 (s, 1H, C3-H), 6.96–6.99 (d, 1H, C6-H, *J* = 9 Hz), 7.53–7.63 (m, 3H, C-14, C-15, C-16), 7.71–7.74 (d, 1H, C5-H, *J* = 9 Hz), 7.98–7.96 (d, 2H, C-13,C-17, *J* = 6.9 Hz), 8.70 (s, 1H, HC = N), 12.44 (s, 1H, NH), 12.87 (s, 1H, OH).

**Synthesis of Cu<sub>2</sub>[(L)<sub>2</sub>(DMF)<sub>2</sub>] (**1**).** To a methanol solution (10 mL) of Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O (0.482 g, 2 mmol), the ligand H<sub>2</sub>L (0.644 g, 2 mmol) in 15 mL of DMF was added upon constant stirring. The resulting green solution was kept undisturbed at room temperature. Dark-green square-shaped single crystals of **1** were generated after one week. These were separated by filtration and air-dried before X-ray diffraction analysis. Yield: 0.69 g. Anal. calc. for C<sub>42</sub>H<sub>38</sub>Cu<sub>2</sub>N<sub>6</sub>O<sub>10</sub>: C, 55.20; H, 4.20; N, 9.19. Found: C, 55.373; H, 3.87; N, 9.58%.

### X-ray crystallography

The crystal structure of **1** was determined by X-ray diffraction methods. Crystal data and experimental details of data collection and structure refinement are reported in Table 1. Intensity data and cell parameters were recorded at 293(2) K on a Bruker Breeze (MoK $\alpha$  radiation  $\lambda$  = 0.71069 Å) equipped with a CCD area detector and a graphite monochromator. The  $\omega$ : $2\theta$  scan technique was applied within a  $\theta$  range of 1.91–30.63°. No significant crystal decay was observed. A total of 31 462 reflections were collected of which 4850 were independent [*R*(int) = 0.0358] reflections.

The raw frame data were processed using SAINT and SADABS to yield the reflection data file.<sup>26</sup> The structure was solved by Direct Methods using the SIR97 program<sup>27</sup> and refined on *F*<sub>o</sub><sup>2</sup> by full-matrix least-squares procedures, using the SHELXL-97 program<sup>28</sup> in the WinGX suite v.1.80.05.<sup>29</sup> All non-hydrogen atoms were refined by anisotropic atomic displacements. The hydrogen atoms were included in the refinement at idealized geometry (C–H 0.95 Å) and refined “riding” on the corresponding parent atoms. The weighting scheme used in the last cycle of refinement was *w* = 1/[ $\sigma^2 F_o^2 + (0.0583P)^2 + 0.4073P$ ], where *P* = (*F*<sub>o</sub><sup>2</sup> + 2*F*<sub>c</sub><sup>2</sup>)/3. CCDC-938644 for **1**.

### Theoretical calculation

Full geometry optimization of **1** was carried out using density functional theory (DFT) at the B3LYP level.<sup>30</sup> All calculations were performed using the Gaussian 03 program package<sup>31</sup> with the aid of the Gauss View visualization program.<sup>32</sup> For C, H, N, O, and Cl the 6-31G(d) basis set was assigned, while for Cu and Ni the LanL2DZ basis set with effective core potential was employed.<sup>33</sup> The vibrational frequency calculations were performed to ensure that the optimized geometries represent the local minima and there are only positive eigen values. Vertical electronic excitations based on B3LYP optimized geometries were computed by the time-dependent density functional theory (TD-DFT) formalism<sup>34–36</sup> in acetonitrile using the conductor-like polarizable continuum model (CPCM).<sup>37</sup> The Gauss sum was used to calculate the fractional contributions of various groups to each molecular orbital.<sup>38</sup>

### Anti-mycobacterial activity

**Medium.** In the assays, Mycobacteria Growth Indicator Tubes (MGITs) and their supplements, BBL MGIT OADC enrichment and BBL MGIT PANTA were purchased from BD. The MGIT (Mycobacteria Growth Indicator Tube) contains 4 mL of modified Middlebrook 7H9 Broth base.

**Inoculum preparation.** For the cultivation of mycobacteria, the MGIT (Mycobacteria Growth Indicator Tube), a fluorescent compound, is embedded in silicone at the bottom, and then 4 mL of modified Middle brook 7H9 Broth base are added to the mixture. After that 0.5 mL of OADC (oleic acid, albumin, dextrose and catalase) enrichment, and the PANTA antibiotic mixture used to prevent the growth of any non-mycobacteria (0.1 mL) are added to this medium. Oleic acid plays an important role in the metabolism of mycobacteria; albumin acts as a protective agent; dextrose is an energy source; catalase destroys toxic peroxides. Tubes are incubated at 37 °C. For positive control, MGIT tubes are prepared by inoculating bacteria. An un-inoculated MGIT tube is used as a negative control. Blood agar is used for checking the growth of other bacteria. Daily tube reading starts on the second day of incubation using a Micro MGIT fluorescence reader which has long wave UV light.<sup>39</sup> To prepare inoculums from a positive BACTEC MGIT tube, the positive tubes (day 1 or day 2 positive) are used directly as inoculums. The positive tubes between day 3 and day 5 are diluted to 1:4 ratio using sterile saline. Inoculums, prepared from a Day 1 to Day 5 MGIT 7 mL positive tube, range between



$0.8 \times 10^5$  and  $3.2 \times 10^5$  CFU mL $^{-1}$ . Each assay is performed according to the MGIT manual fluorometric susceptibility test procedure recommended by the manufacturer.<sup>39,40</sup>

**Antimycobacterial susceptibility assay.** The activity of the ligand, H<sub>2</sub>L, and complex 1 against *M. tuberculosis* strains was tested using the Microplate Presto Blue Assay (MPBA) by the method described by Collins & Franzblau<sup>41</sup> and modified by Jimenez-Arellanes *et al.*<sup>42</sup> 128  $\mu$ L of compound 1 and 72  $\mu$ L of 7H9 broth were transferred into the first column, and then 100  $\mu$ L of 7H9 broth was transferred from column 2 to column 8. Column 9 and 10 were the positive and the negative control, respectively. 100  $\mu$ L of a mixture of broth and the compound were transferred from column 1 to column 2 and then mixed using pipettes three times; the procedure was repeated to provide serial 1:2 dilutions. 100  $\mu$ L of the excess medium was discarded from the wells in column 8. Afterwards, 20  $\mu$ L of *M. tuberculosis* inoculum was added to the wells of column 1 to 8 and to the positive control columns. Negative control columns were not inoculated with bacteria. Positive control columns included 7H9 broth and bacteria, while negative control columns contained 7H9 broth and compound 1. Final-test concentration ranges were 423.68–6.62  $\mu$ mol L $^{-1}$  in the mixture. Microplates were inoculated with the bacterial suspension (20  $\mu$ L per well) except for the negative control and incubated at 37 °C for 6 days. Presto blue (15  $\mu$ L, Life Technologies) was then added to the bacterial growth control wells (without compound 1) and the microplates were incubated at 37 °C for an additional 24 h. If the dye turned from blue to pink (indicating positive bacterial growth), the Presto blue solution was added to the other wells to determine the MIC values. All tests were performed in triplicate. The minimal inhibitory concentration (MIC) was defined as the lowest concentration of the sample that prevents a color change to pink. The minimal bactericidal concentration (MBC) corresponded to the minimum compound concentration which does not cause a color change in the cultures when re-incubated in fresh medium.<sup>43</sup> Streptomycin (STR) from Sigma and isoniazid (INH) from Fluka were used as standard drugs.

### Cytotoxicity assay

**Cell lines and culture conditions.** MCF7, a human breast adenocarcinoma cell line, was obtained from the American Type Culture Collection (ATCC, Bethesda, MD, USA). Cells were grown in DMEM (Invitrogen, USA) containing 10% heat inactivated fetal bovine serum (FBS) (GIBCO BRL, USA), at 37 °C in a humidified atmosphere containing 5% CO<sub>2</sub>.

**Cell viability test by MTT assay.** MTT [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide] assay was performed to determine the effect of compound 1 and the ligand (0.4, 0.8, 1.6, 3.12, 6.25, 12.50, 25, 50, and 100  $\mu$ M) on the proliferation of MCF7 cells by their utilization and conversion to formazan according to the method of Mosmann.<sup>44</sup> MCF7 cells were plated into 96-well microtiter plates at a density of  $2.5 \times 10^4$  cells per well. After 24 h, the culture medium was replaced by 200  $\mu$ L (0–100  $\mu$ M) of compound, and the cells were incubated for 24 and 48 h. DMSO treated cells were used as controls. After treatment, the media were replaced with serum free DMEM

media containing 1 mg mL $^{-1}$  of MTT and incubated at 37 °C in a 5% CO<sub>2</sub> incubator for additional 5 h. After 5 h, the unreacted dye was removed, and the insoluble formazan crystals were dissolved in 100  $\mu$ L per well DMSO and measured spectrophotometrically in a VersaMax tunable microplate reader (Molecular Devices, Sunnyvale, CA, USA) at 570 nm. The relative cell viability (%) related to control wells containing cell culture medium without samples was calculated by  $A_{570\text{nm}}[\text{sample}]/A_{570\text{nm}}[\text{control}] \times 100$ .

### Docking studies

Molecular docking is used to predict how a protein interacts with small molecules. The crystal structure of enoyl acyl carrier protein reductase of *M. tuberculosis* H37R<sub>v</sub> was downloaded from the RSCB protein data bank (<http://www.pdb.org>) and used for docking. The protein (PDB id: 4U0K) was co-crystallized with (3S)-N-(5-chloro-2-methylphenyl)-1-cyclohexyl-5-oxopyrrolidine-3-carboxamide and nicotinamide-adenine-dinucleotide. *In silico* docking studies were performed using the CDOCKER module of the Receptor–Ligand interaction protocol section of Discovery Studio client 3.5.<sup>45</sup> Initially, there was a pretreatment process for the protein and the ligand. The structure of the ligand was drawn using Chemdraw 5.0, saved in .mol file and finally the .mol file was imported to the Discovery studio 4.0 platform. Ligand preparation was done using the Prepare Ligand module in the Receptor–Ligand interaction tool of Discovery studio 4.0 and the prepared ligand was used for docking. Protein preparation was done under the Prepare Protein module of the Receptor–Ligand interaction tool of Discovery Studio 4.0 and that was used for docking. Then we have defined the protein as the total receptor and the active site was selected based on the ligand binding domain of (3S)-N-(5-chloro-2-methylphenyl)-1-cyclohexyl-5-oxopyrrolidine-3-carboxamide and nicotinamide-adenine-dinucleotide. Then the pre-existing ligand was removed and the prepared ligand was placed. The most favorable docked pose was selected according to the minimum free energy of the protein–ligand complex and analyzed to investigate the interaction.

### ADMET prediction

Absorption, distribution, metabolism, excretion and toxicity (ADMET) predictions were done in the ADMET descriptor module of the Small molecule protocol of Discovery studio client 4.0. The druglikeness of the compounds was also checked following Lipinski's rule of five.<sup>46,47</sup>

## Results and discussion

### Synthesis and formulation

The ligand, [benzoic acid(7-hydroxy-4-methyl-2-oxo-2H-chromen-8-yl-methylene)-hydrazide] (H<sub>2</sub>L), was prepared by the condensation of 8-formyl-7-hydroxy-4-methyl-coumarin with benzohydrazide (1:1 mole proportion) in methanol. H<sub>2</sub>L was then reacted with Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O in methanol to yield the complex Cu<sub>2</sub>[(L)<sub>2</sub>(DMF)<sub>2</sub>] (1), which was crystallized by slow evaporation of the solvent.



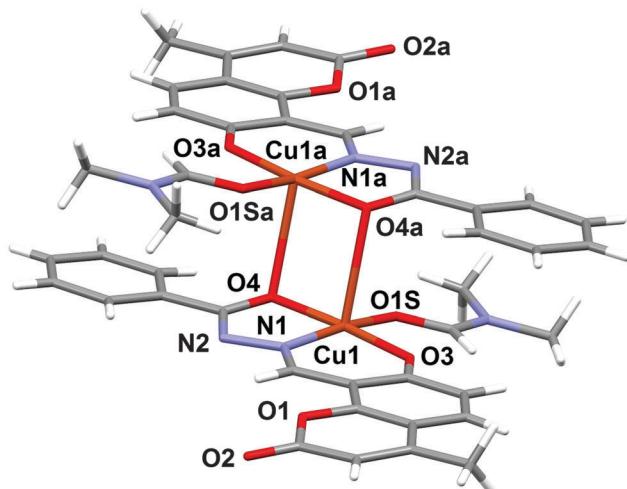


Fig. 1 Molecular structure of complex **1** with the partial labeling scheme.

The IR spectra of **1** show a  $\nu(\text{O}=\text{C}-\text{O})$  band at  $1720\text{ cm}^{-1}$  that represents the stretching frequency of the lactone (the corresponding peak appears at  $1707\text{ cm}^{-1}$  in  $\text{H}_2\text{L}$ ). The band at  $1618\text{ cm}^{-1}$  is assigned to coordinated  $\nu(\text{C}=\text{N})$ ,<sup>48</sup> whereas in the free ligand, the band appears at a lower wavenumber ( $1613\text{ cm}^{-1}$ ). A high intensity band in the region  $1303\text{--}1269\text{ cm}^{-1}$  is assigned to the phenolic  $\nu(\text{C}-\text{O})$  vibration, whereas the shifting of this band at  $1370\text{--}1400\text{ cm}^{-1}$  supports the coordination of the phenolic oxygen to the metal ion *via* deprotonation. The  $\nu_{(\text{M}-\text{N})}$  and  $\nu_{(\text{M}-\text{O})}$  stretching frequencies are observed at  $712$  and  $462\text{ cm}^{-1}$ .

### Crystal structure

The molecular structure of the dinuclear copper compound is shown in Fig. 1. Relevant bond lengths and angles are given in Table 2. The geometric centre of the molecule lies on a crystallographic inversion centre. The geometry of the  $\text{NO}_4$  environment around each copper ion is square pyramidal, as evidenced by the  $\tau$  value<sup>49</sup> of  $0.08$ . The dimer is assembled *via* the  $\mu_2$ -enolato oxygen atoms  $\text{O}4$  and its centrosymmetric analogue  $\text{O}4\text{a}$ , belonging to two distinct deprotonated ligand  $\text{L}^{2-}$ . Each of the oxygen atoms bridge two adjacent copper ions, thus occupying both the apical position of the pyramid around one metal center, and the equatorial position of its centrosymmetric counterpart. The coordination around copper is completed by the imine- $\text{N}(1)$  and phenolic- $\text{O}(3)$  atoms of the Schiff base and by the  $\text{O}1\text{S}$  atom of the DMF molecule. This arrangement gives rise to a  $\text{Cu}_2\text{O}_2$  distorted square and to a pair of five and six-membered chelate rings. The  $\text{Cu}1\text{-Cu}1\text{a}$  distance is  $3.429(6)\text{ \AA}$ , which is in good

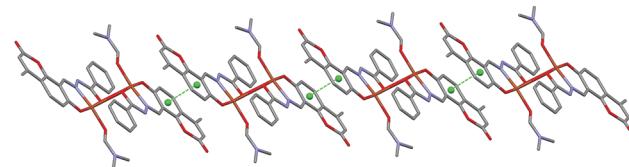


Fig. 2  $\pi\text{-}\pi$  stacking interactions present in the crystal structure of **1**, shown along the *b* axis of the unit cell. Centroids are represented as green spheres.

agreement with previously reported doubly phenolato-bridged copper complexes,<sup>50,51</sup> where the  $\text{Cu}\text{-Cu}$  separation is in the range  $2.901\text{--}3.345\text{ \AA}$ . The  $\text{Cu}1\text{-O}4$  distance is  $1.943(1)\text{ \AA}$ , which is comparable with similar bond distances [ $1.922(4)$  and  $1.941(4)\text{ \AA}$ ] reported in the literature.<sup>50a</sup> On the other hand, the  $\text{Cu}1\text{-O}4\text{-Cu}1\text{a}$  angle of  $95.33(4)^\circ$  is relatively small if compared with literature data,<sup>50a,d,52</sup> where the angular values span from  $95.7$  to  $107.5^\circ$ .<sup>53</sup> In the lattice, the crystal structure is stabilized by weak  $\pi\text{-}\pi$  stacking interactions involving the aromatic rings of the ligand. Fig. 2 shows the packing along the *b* axis of the unit cell, with centroid- $\cdot\cdot\cdot$ centroid distances of  $3.849(5)\text{ \AA}$ .

### Electronic spectra

The electronic spectra for  $\text{H}_2\text{L}$  and complex **1** were recorded in HPLC grade dimethyl formamide and methanol, respectively. For  $\text{H}_2\text{L}$ , the absorption bands at  $314$  ( $\varepsilon = 201\,600\text{ M}^{-1}\text{ cm}^{-1}$ ),  $347$  ( $\varepsilon = 115\,818\text{ M}^{-1}\text{ cm}^{-1}$ ),  $383$  ( $\varepsilon = 38218\text{ M}^{-1}\text{ cm}^{-1}$ ) and  $463\text{ nm}$  ( $\varepsilon = 28\,072\text{ M}^{-1}\text{ cm}^{-1}$ ) are attributed to  $\pi\rightarrow\pi^*$  and  $n\rightarrow\pi^*$  transitions.<sup>54</sup> Upon complexation, the  $\pi\rightarrow\pi^*$  band shifts from  $314$  to  $337\text{ nm}$  ( $\varepsilon = 38\,750\text{ M}^{-1}\text{ cm}^{-1}$ ), while the  $n\rightarrow\pi^*$  band moves to  $392$  ( $\varepsilon = 21\,583\text{ M}^{-1}\text{ cm}^{-1}$ ) and  $408\text{ nm}$  ( $\varepsilon = 20\,541\text{ M}^{-1}\text{ cm}^{-1}$ ). Furthermore for the complex, a pure d-d transition appears at  $650\text{ nm}$  ( $\varepsilon = 1008\text{ M}^{-1}\text{ cm}^{-1}$ ).<sup>55</sup>

The spectral characteristics are explained with the help of a DFT computation of the optimized structure of **1**. The orbital energies along with the contributions to some MOs from the ligands and the metal for a few MOs are given in Fig. 3, while the orbital energies of the ligands and the complex are given in Table S1 (ESI<sup>†</sup>). As complex **1** contains two unpaired electrons as  $s = 1/2$ , it shows two spin states designated as  $\alpha$  and  $\beta$ . Both filled and vacant  $\alpha$ -MOs from HOMO-5 to LUMO+5 constitute mainly of ligand orbitals ( $\text{L}^{2-}$ ). The metal and DMF contributions are insignificant in both filled and vacant MOs; only HOMO-5 and HOMO-4 carry  $15\%$  and  $14\%$  Cu functions whereas LUMO+4 and LUMO+5 carry  $9\%$  and  $12\%$  DMF functions in the  $\alpha$  state, respectively. Thus, the ligand  $\text{L}^{2-}$  controls the molecular functions and hence the electronic properties of the complexes. Thus, in the  $\alpha$  state all the transition bands are basically considered as ILCT in nature. The intensity of these transitions has been assessed by the oscillator strength ( $f$ ). HOMO  $\rightarrow$  LUMO is considered as  $\text{L}(\pi) \rightarrow \text{L}(\pi^*)$  with  $f = 0.5762$  at  $405.8\text{ nm}$ ; HOMO-1  $\rightarrow$  LUMO+3 is found at  $335.3\text{ nm}$  with  $f = 0.3549$  which is also represented as ILCT (see Table S2, ESI<sup>†</sup>). In a similar way, the orbital energy of the  $\beta$  state of **1** was also calculated (see Table S3, ESI<sup>†</sup>) and the calculated transitions are grouped. Here also the contribution of the ligand  $\text{L}^{2-}$  in

Table 2 Selected bond lengths [ $\text{\AA}$ ] and angles [ $^\circ$ ] for 1

Cu1-O3	1.892(1)	O4-Cu1-O3	174.34(5)
Cu1-O4	1.943(1)	O3-Cu1-N1	93.08(5)
Cu1-O4 <sup>a</sup>	2.651(1)	O3-Cu1-O1S	93.12(5)
Cu1-N1	1.919(1)	O4-Cu1-O1S	92.03(5)
Cu1-O1S	1.971(1)	N1-Cu1-O4	81.42(5)
		N1-Cu1-O1S	167.52(5)

<sup>a</sup>  $-x, -y + 1, -z$ .

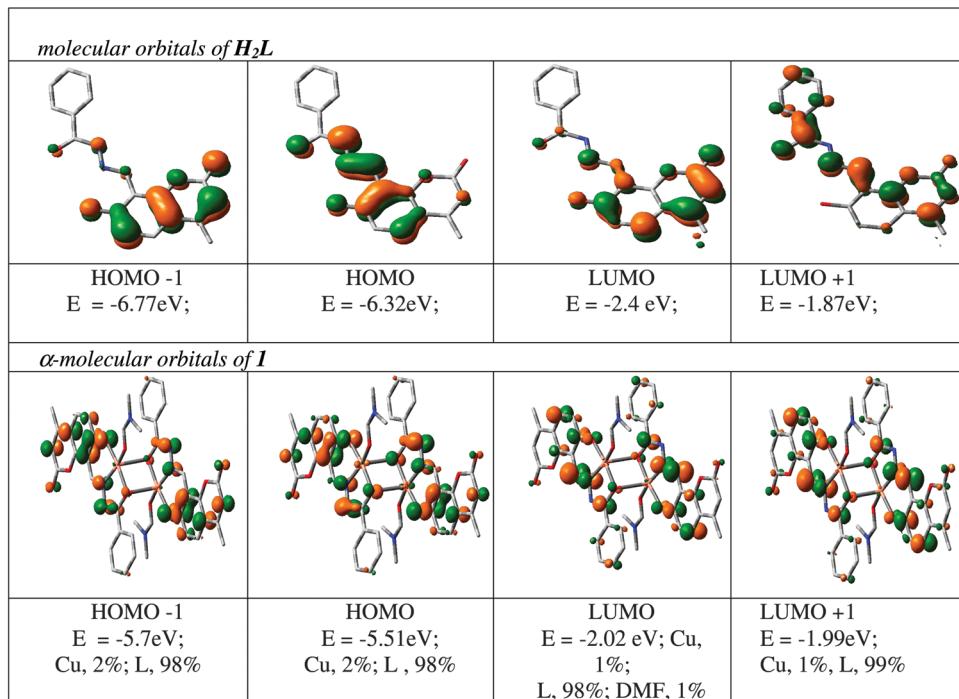


Fig. 3 Contour plots of some selected MOs of  $H_2L$  and  $Cu_2([L]_2(DMF)_2)$  (**1**).

both filled and vacant MOs is  $>80\%$  except for LUMO and LUMO+1 where the contribution is: 34% L; 62% Cu; 4% DMF; and 36% L; 60% Cu; 4% DMF, respectively. In this case the transition bands inferior to 400 nm are ILCT in nature except for the HOMO-9  $\rightarrow$  LUMO+1 transition at 360 nm ( $f = 0.0201$ ) which is a mixture of LMCT and ILCT. Additionally, absorption bands at 434 ( $f = 0.0025$ ) and 525 ( $f = 0.0046$ ) nm are also considered as the mixture of LMCT and ILCT which are due to HOMO-3  $\rightarrow$  LUMO+1 and HOMO  $\rightarrow$  LUMO+1 transitions.

#### Emission spectra

The emission spectra of  $H_2L$  and **1** were carried out in DMF and methanol, respectively, at room temperature. They show that  $H_2L$  fluoresces strongly when its  $\pi \rightarrow \pi^*$  transition band is excited,

whereas complex **1** is a weak emitter (Fig. 4). For  $H_2L$ , a strong emission band appears at 498 nm upon excitation at 314, 347 or 383 nm (which are  $\pi \rightarrow \pi^*$  bands). This means that  $H_2L$  shows a high Stokes shift value of 184 nm ( $\Delta\lambda' = \lambda_{\text{emission}} - \lambda_{\text{absorption}}$ ) (Fig. 5) which corresponds to the Stokes shift parameter of  $11\,767\text{ cm}^{-1}$ . The Stokes shift parameter indicates the difference in properties and structure between the ground state  $S_0$  and the first excited state  $S_1$ ,<sup>56</sup> and has been estimated according to the following equation:<sup>57</sup>

$$\Delta\nu' = \nu_{\text{abs}}' - \nu_{\text{em}}' = (1/\lambda_{\text{abs}} - 1/\lambda_{\text{em}}) \times 10^7\text{ cm}^{-1}$$

where:  $\lambda_{\text{abs}}$  and  $\lambda_{\text{em}}$  = absorbance and emission wavelength (nm),  $\nu_{\text{abs}}'$  and  $\nu_{\text{em}}'$  = wavenumbers for the absorption and emission maxima ( $\text{cm}^{-1}$ ).

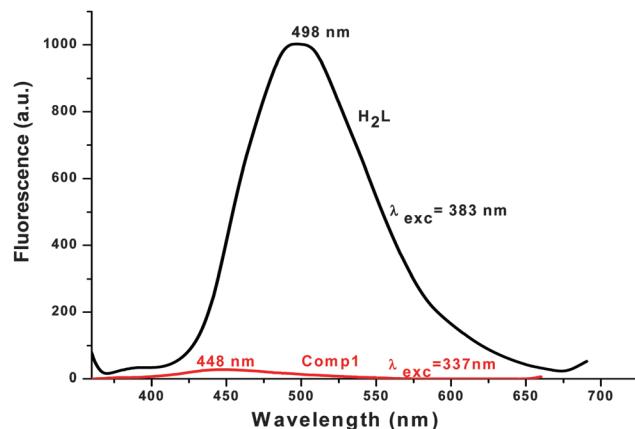


Fig. 4 Emission spectra of  $H_2L$  (black curve, in DMF) and of the Cu(II) complex (**1**) (red curve, in MeOH).

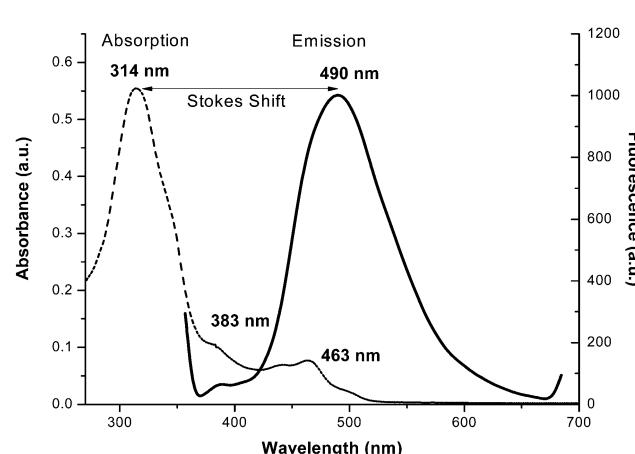


Fig. 5 Emission spectra with the Stokes shift value.



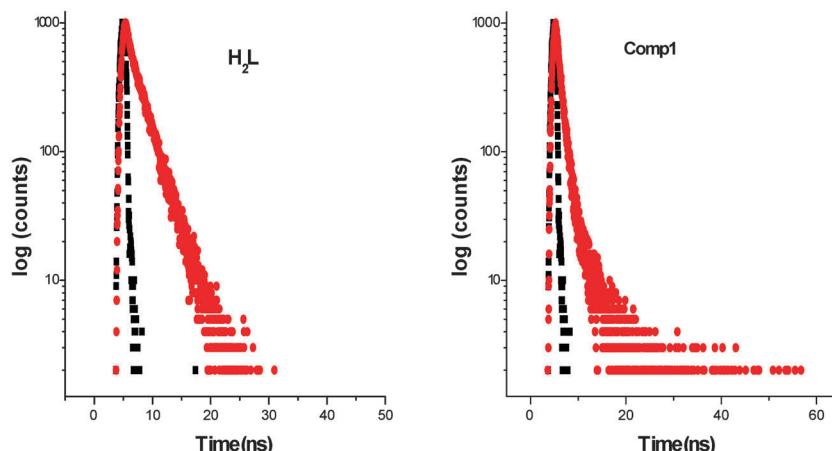


Fig. 6 Exponential decay profile (red) of  $\text{H}_2\text{L}$  and complex **1**.

Complex **1** fluoresces very weakly at 448 nm when it is excited at 337 nm. Longer wavelengths of excitation (MLCT/d-d bands) do not produce any emission. Though the Stokes shift parameter of the ligand is comparatively higher with respect to the well-known coumarin dye, coumarin-1 (C1; 7- $\text{NEt}_2$ -4- $\text{CH}_3$ -1,2-benzopyrone,  $\Delta\nu' = 4259\text{ cm}^{-1}$  in acetonitrile), the fluorescence quantum yield of the ligand ( $\phi_{\text{H}_2\text{L}}: 0.17$ ) is much less than C1( $\phi_{\text{C}1}: 0.62$ ). This is basically due to the presence of an OH oscillator, which acts as an efficient quencher enhancing the intersystem crossing process. In addition to that, the stretching vibrations of the CH, NH and C=O oscillators also have an effect on quenching of luminescence.<sup>58</sup> In the case of **1**, the fluorescence quantum yield ( $\phi_1: 0.008$ ) decreases by  $\sim 20$  times due to the high paramagnetic quenching of the Cu(II) metal (d<sup>9</sup>).<sup>59</sup> The decay studies of the free ligand and the complex have been performed in DMF and methanol respectively, upon excitation at 370 nm (Fig. 6, left). The decay profile of the ligand fits with a mono exponential decay whereas in the case of the complex, the nature of the decay is bi-exponential (Fig. 6, right). The mean lifetime ( $\tau_f = a_1\tau_1 + a_2\tau_2$ , where  $a_1$  and  $a_2$  are relative amplitudes of the decay process for bi-exponential) has been

calculated to compare the excited state stability of the complex (1 ns) and the free ligand (3 ns). This indicates that the excited states of **1** are unstable compared to those of  $\text{H}_2\text{L}$ . Radiative and non-radiative rate constant values have also been calculated showing, as usual,  $k_{\text{nr}}$  values [ $3.09 \times 10^8$  ( $\text{H}_2\text{L}$ ) and  $9.8 \times 10^8$  (**1**)] higher than the relative  $k_r$  values [ $6.32 \times 10^7$  ( $\text{H}_2\text{L}$ ) and  $7.97 \times 10^6$  (**1**)]. In the complex, the rate constant of non-radiative decay is much higher ( $\sim 100$  times) than the radiative decay, evidencing that **1** is a very weaker emitter than the ligand.

#### EPR spectra

The EPR spectra of the polycrystalline solid complex **1** were recorded at 298 and 77 K. The two spectra are shown in Fig. 7. No resonances below 2800 and above 3400 Gauss and no half-field transition  $\Delta M = \pm 2$ , attributable to a magnetic coupling between two 3d<sup>9</sup> Cu(II) ions, were detected. This indicates that the two ions, separated by 3.429 Å, are not interacting, in agreement with the EPR behaviour of other Cu dimeric complexes reported in the literature.<sup>60</sup> EPR spectra are 'tetragonal' at RT, with two *g* values, and 'rhombic' at 77 K, with three *g* values. This suggests a geometric distortion when lowering the temperature, which determines a slight *x*, *y* anisotropy. At RT, the order  $g_z \gg g_x = g_y > g_e$  (Table 3) is characteristic of a ground state  $\Psi$  based on the Cu-d<sub>x<sup>2</sup>-y<sup>2</sup></sub> orbital and it is compatible with the square pyramidal environment of each copper ion. At 77 K, the order  $g_z \gg g_y > g_y > g_e$  (Table 3) indicates a ground state  $\Psi$  which can be described as a linear combination of Cu-d<sub>x<sup>2</sup>-y<sup>2</sup></sub> and Cu-d<sub>z<sup>2</sup></sub> orbitals,  $\Psi = c_1|\text{d}_{x^2-y^2}\rangle + c_2|\text{d}_{z^2}\rangle$ ,<sup>61</sup> the parameter *R* ( $R = (g_y - g_x)/(g_z - g_y)$ , with  $g_z > g_y > g_x$ ) being indicative of the predominance of the Cu-d<sub>x<sup>2</sup>-y<sup>2</sup></sub> orbital ( $c_1 > c_2$ ) or Cu-d<sub>z<sup>2</sup></sub> ( $c_2 > c_1$ ).<sup>62</sup> If  $R < 1$ , the greater contribution to the ground state arises from the Cu-d<sub>z<sup>2</sup></sub> orbital, while if  $R > 1$  this arises from Cu-d<sub>x<sup>2</sup>-y<sup>2</sup></sub> orbital.<sup>61b</sup> For **1**, *R* is 2.7 and this means that  $c_1 > c_2$  and that a slight difference between the *x* and the *y* axis occurs at low temperatures, but the structure remains close to the square pyramidal geometry.<sup>61b</sup> Therefore, variations in temperature cause slight structural changes in bond lengths and angles in the *xy* plane, which results in a mixing

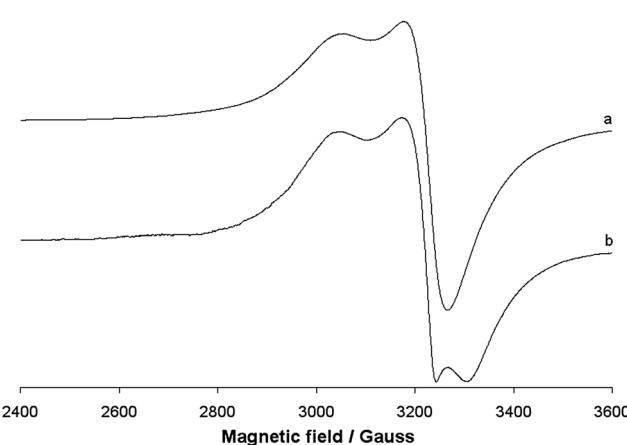


Fig. 7 X-band EPR spectra of the polycrystalline solid complex **1** at (a) 298 and (b) 77 K.



Table 3 EPR parameters of complex **1** in the solid state and in some organic solvents<sup>a</sup>

	$g_x$	$g_y$	$g_z$	$A_x$	$A_y$	$A_z$	$R$	Ground state
Solid at 298 K	2.080	2.080	2.205	—	—	—	$\infty$	$\text{Cu-d}_{x^2-y^2}$
Solid at 77 K	2.038	2.084	2.208	—	—	—	2.7	$\text{Cu-d}_{x^2-y^2}$
Solid dissolved in DMSO	2.040	2.069	2.253	13.0	15.0	186.9	6.3	$\text{Cu-d}_{x^2-y^2}$
Solid dissolved in MeOH	2.041	2.069	2.254	12.0	14.0	185.4	6.6	$\text{Cu-d}_{x^2-y^2}$
Solid dissolved in DMF	2.040	2.068	2.251	13.0	15.5	187.3	6.5	$\text{Cu-d}_{x^2-y^2}$

<sup>a</sup>  $A_x$ ,  $A_y$  and  $A_z$  values reported in  $10^{-4} \text{ cm}^{-1}$ .

between the  $\text{Cu-d}_{x^2-y^2}$  and  $\text{Cu-d}_{z^2}$  orbitals in the singly occupied molecular orbital that bears the unpaired electron. Similar effects induced by the temperature were observed from some other  $\text{Cu}(\text{II})$  compounds.<sup>63</sup> When **1** is dissolved in an organic solvent (DMSO, DMF or MeOH) at 298 K, the dinuclear  $\text{Cu}(\text{II})$  structure is destroyed; the solvent breaks the long  $\text{Cu-O4}$  bond, replaces the alkoxide oxygen in the axial position and mononuclear  $\text{Cu}(\text{II})$  species are formed (Fig. 8). This has been observed for other polynuclear  $\text{Cu}(\text{II})$  structures.<sup>64</sup> In all the three solvents, the spectra are slightly 'rhombic' with  $g_z \gg g_y > g_x > g_e$ , similarly to what were observed in the solid state (Table 3). Therefore, the ground state is based on the  $\text{Cu-d}_{x^2-y^2}$  orbital and the geometry around copper is a distorted square pyramid. The values of  $g_z$  and  $A_z$  are compatible with an equatorial  $\text{NO}_3^-$  coordination and a charge close to the neutrality around the metal ion.<sup>65</sup>

Interestingly, in the spectra recorded in MeOH and DMF, an EPR resonance centered at  $g \sim 2.24$  is detected. The intensity of this band increases from MeOH to DMF and cannot be ascribed to a residual presence of the dimeric structure because no other absorptions (nor at lower and higher magnetic fields) are observed. The broad and unresolved band can be assigned to an aggregation processes originated in MeOH and DMF, probably favored from the formation of  $\pi$  stacking interactions. Owing to such processes, the distance between the Cu centers decreases significantly and an isotropic band due to the magnetic interaction of the  $S = 1/2$  paramagnetic metal ions appears. These aggregations have been observed for other paramagnetic species

and depend on the solvent used; the effect can be detected using EPR spectroscopy but not using UV-vis spectrophotometry.<sup>66</sup>

The appearance of this band, characteristic of the solid samples, is not accompanied by the formation of a precipitate, ruled out by the electronic absorption spectra (Fig. 9). Therefore, this state can be interpreted as the intermediate between the solution and the solid state, in which the molecules undergo aggregation through  $\pi$  stacking favored by the presence of the two aromatic rings in the ligand structure.

### Magnetic properties

The variable-temperature magnetic properties (in the temperature range from 5 to 300 K under an external field of 10 000 Oe) of complex **1** in the form of  $\chi_M T$  vs.  $T$  ( $\chi_M$  vs.  $T$  inset) plots are illustrated in Fig. 10 ( $\chi_M$  is the molar susceptibility for two  $\text{Cu}(\text{II})$ ). At room temperature, the  $\chi_M T$  value is 6.74 emu K mol<sup>-1</sup>. When lowering the temperature, the  $\chi_M T$  value increases smoothly to a maximum value of 8.83 emu K mol<sup>-1</sup> at 30 K. It then decreases sharply and reaches a value of 7.94 emu K mol<sup>-1</sup> at 5 K. The behavior suggests that there is weak ferromagnetic interactions between the bridging dinuclear  $\text{Cu}(\text{II})$  ions. Furthermore, the effective magnetic moment at room temperature is 2.07 B.M., which is less than the spin only value of  $\text{Cu}(\text{II})$ , 2.45 B.M., calculated for two uncoupled spin 1/2 centers.

### Antimycobacterial activity

In the anti-mycobacterial assay,  $\text{H}_2\text{L}$  and complex **1** were tested against *M. tuberculosis* H37Rv ATCC 27294 and *M. tuberculosis* H37Ra ATCC 25177 strains as well as against two clinical strains

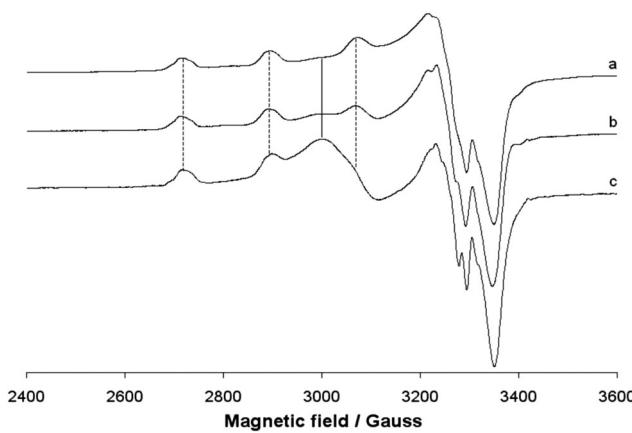


Fig. 8 Anisotropic X-band EPR spectra of the solid complex **1** dissolved in (a) DMSO, (b) MeOH and (c) DMF. The dotted lines indicate the first three parallel resonances due to the coupling between the unpaired electron and the<sup>63,65</sup>  $\text{Cu}$  nucleus, while the full line shows the position of the isotropic band due to the aggregate observed in MeOH and DMF.

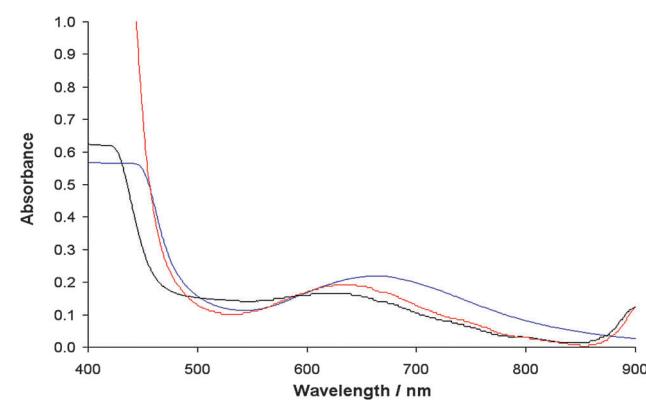


Fig. 9 Electronic absorption spectra of the solid complex **1** dissolved in DMSO (blue), DMF (red) and MeOH (black). The complex concentration was  $1 \times 10^{-3}$  M.



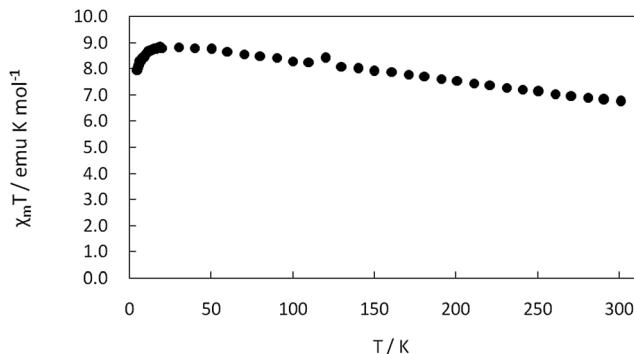


Fig. 10 The  $\chi_M T$  vs.  $T$  plots for complex **1** in the temperature range 5–300 K.

(strain 1 and strain 2). The results are shown in Table S4 (ESI†). *M. tuberculosis* H37Rv and *M. tuberculosis* H37R, well-known indicators, were used for the drug sensitivity tests.<sup>20</sup> The results showed that the ligand H<sub>2</sub>L and complex **1** both exhibited antimycobacterial activity against all the tested *M. tuberculosis* strains, with MIC values of 52.96  $\mu\text{mol L}^{-1}$  and MBC values in the range of

52.96–423.68  $\mu\text{mol L}^{-1}$ . Both H<sub>2</sub>L and complex **1** showed the bactericidal activity and killed the bacteria (Table S4 and Fig. S4, ESI†). Trias *et al.*<sup>67</sup> showed that *Mycobacterium chelonae* developed a pore-forming protein as a mycobacterial porin. In the study of permeability of *Mycobacterium smegmatis*, Trias and Benz<sup>68</sup> reported that the mycobacterial porin formed an important hydrophilic structure where small molecules pass through the pores and diffuse into the cell. In this study, we evidence that all compounds showed a considerable efficacy on the Mycobacterial strains. The mycobacterial cell wall includes a large amount of complex lipids, lipopolysaccharides and mycolic acids. This constitution makes the cell wall a hydrophobic strong barrier against antimicrobial agents.<sup>69</sup> These results are well comparable with previous literature.<sup>70</sup> The MIC (Minimal Inhibitory Concentration) and MBC (Minimal Bactericidal Concentration) values are higher in the present case. The concerned data respond well while comparing with known drugs like streptomycin and isoniazid.

#### Cytotoxicity study

Fig. 11 shows the effects of complex **1** and ligand H<sub>2</sub>L in human adenocarcinoma cells. In order to determine the cytotoxic effect

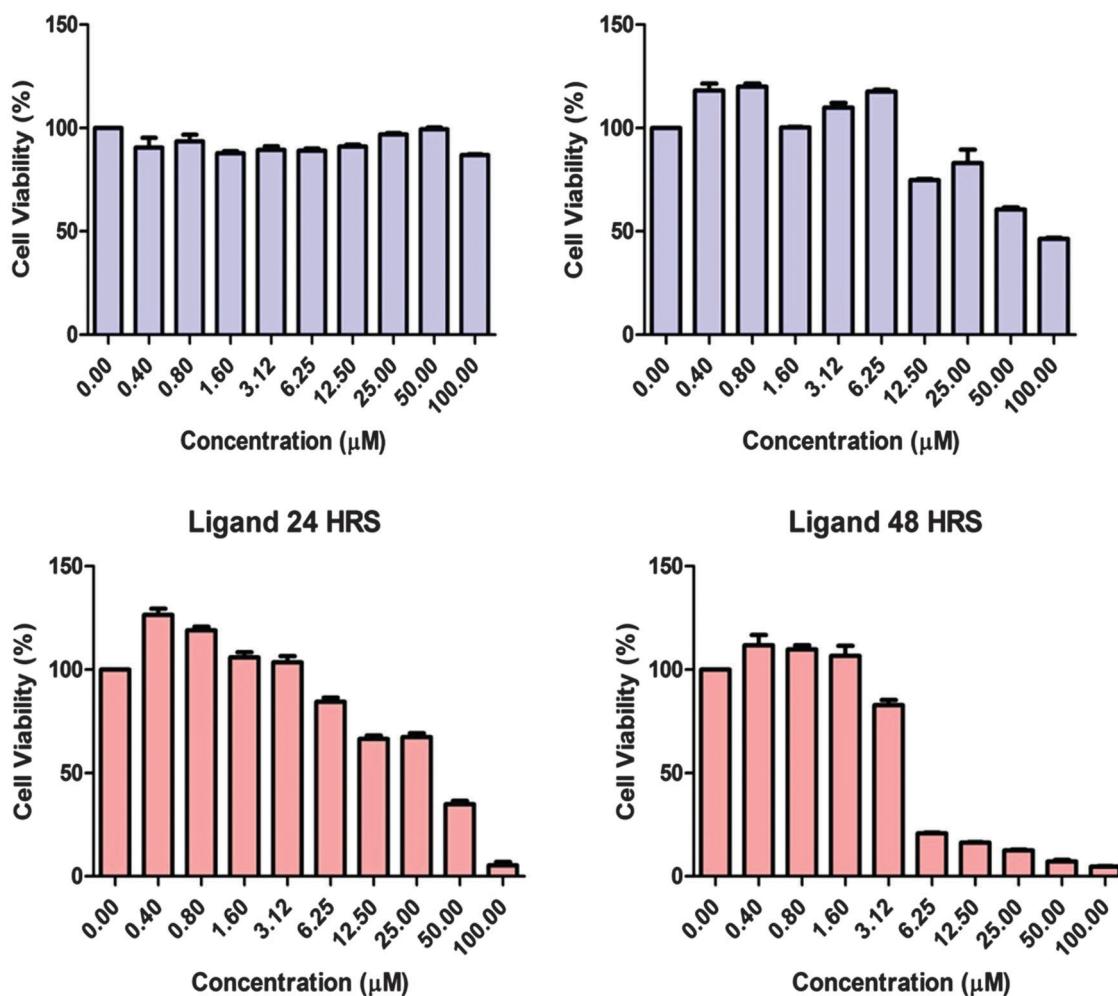


Fig. 11 Evaluation of the cytotoxic effect on the adenocarcinoma cell line. MCF7 cells were seeded ( $2.5 \times 10^4$  cells per mL) and treated with the compounds (0.4, 0.8, 1.6, 3.12, 6.25, 12.50, 25, 50 and 100  $\mu\text{M}$ ) and cytotoxicity was determined after 24 and 48 hours by MTT assay.

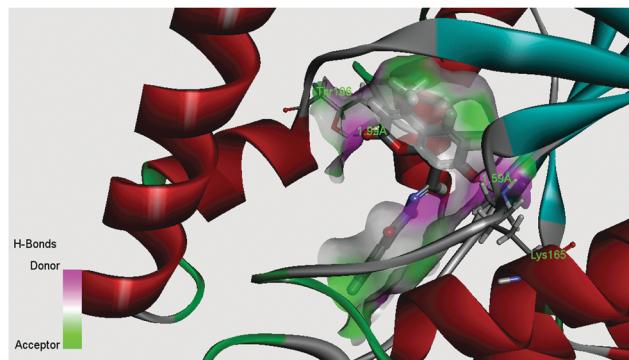


Fig. 12 Surface area (with respect to H-bond donors & acceptors) of the binding pocket in the best docked pose of the ligand and the protein.

of complex **1** and ligand  $\text{H}_2\text{L}$ , the adenocarcinoma cells were treated with or without the compounds and the cell viability was tested after 24 and 48 hours by MTT Assay. The results showed that the cell population was significantly decreased after 48 hours of treatment with  $\text{H}_2\text{L}$ . However complex **1** produced the cytotoxic effect only after 48 hours of incubation. Our results suggest that both complex **1** and the ligand are able to induce cytotoxicity in MCF7 cells with an  $\text{IC}_{50}$  value of 100  $\mu\text{M}$  and 5  $\mu\text{M}$ . Hence the ligand and complex **1** can have potential anticancer properties.

### Docking study

The enoyl acyl carrier protein reductase (ENR) from *M. tuberculosis* is one of the key enzymes involved in the mycobacterial fatty acid elongation cycle and has been validated as an effective antimicrobial target. In the current study we have docked our ligand inside the binding pocket of enoyl acyl carrier protein reductase of *M. tuberculosis* to get an idea about the mode of action. The crystallographic structure of enoyl acyl carrier protein reductase of *M. tuberculosis* H37R<sub>v</sub> was downloaded from the RCSB protein data bank (PDB ID: 4U0K), which was resolved at 1.09  $\text{\AA}$  by X-ray diffraction. Energy minimized structure of the ligand ( $\text{H}_2\text{L}$ ) was used for *in silico* protein ligand docking studies in the cavity of the enzyme. A total of 16 amino acids (Ile21, Gly96, Phe97, Met98, Met103, Met147, Asp148, Phe149, Lys165, Ala191, Gly192, Pro193, Ile194, Thr196, and Met199) were present in the cavity sites and involved in the binding process within 3.5  $\text{\AA}$  radius of the ligand. The ligand,  $\text{H}_2\text{L}$ , interacts with the protein by ionic interaction with Lys165 (1.59  $\text{\AA}$ , Fig. S5 and S6, ESI<sup>†</sup>) and hydrogen bond interaction with Thr196 (1.99  $\text{\AA}$ , Fig. S5 and S6, ESI<sup>†</sup>). The relevant data of the docking study are given in Tables S5 and S6 (ESI<sup>†</sup>).

Table 4 ADMET prediction of  $\text{H}_2\text{L}$

Compounds	Molecular weight	ADMET solubility (aqueous)	ADMET solubility level	ADMET absorption level <sup>a</sup>	ADMET_A log P98	No. of H-bond acceptor	No. of H-bond donor	Lipinski's filter	Druglikeness inference	Ames prediction
$\text{H}_2\text{L}$	322.3	-4.039	2	0 (good)	2.957	5	2	Yes	Yes, low	Non-mutagen

<sup>a</sup> ADMET absorption level: 0, good; 1, moderate; 2, low.

The surface area of the active site cavity (with respect to the H-bond donor/acceptor) is depicted in Fig. 12; the carbonyl group and the -OH group of the coumarin moiety is oriented towards the H-bond donor rich surface of the protein.

### Druglikeness and ADMET prediction

The druglikeness of  $\text{H}_2\text{L}$  was checked following Lipinski's rule of five. The ADMET modules of Discovery Studio were used to check the ADMET (absorption, distribution, metabolism, excretion and toxicity) properties of the compounds. Toxicity prediction, aqueous solubility, blood brain barrier penetration, human intestinal absorption, and Ames mutagenicity were predicted under the Calculate Molecular Property module of the Small molecule tool of Discovery Studio client 4.0. The predicted data are summarized in Table 4. According to these predictions, the ligand,  $\text{H}_2\text{L}$ , is well absorbed and non-mutagenic.

## Conclusions

The coumarin-assisted tridentate ligand, [benzoic acid(7-hydroxy-4-methyl-2-oxo-2*H*-chromen-8-ylmethylene)-hydrazide], has been used to synthesize a phenoxy-bridged Cu(II) dinuclear complex,  $\text{Cu}_2[(\text{L})_2(\text{DMF})_2]$  (**1**). The solid state structure of **1** shows that the Cu atoms are present in a square-pyramidal arrangement. The variable temperature EPR spectra suggest that a geometrical distortion takes place upon lowering the temperature and that the dinuclear structure is destroyed if the complex is dissolved in coordinating solvents. The temperature dependent magnetic moment supports the presence of weak ferromagnetic interactions between the bridging dinuclear Cu(II) ions. Both the ligand  $\text{H}_2\text{L}$  and complex **1** exhibit anti-mycobacterial activity and considerable efficacy on *M. tuberculosis* H37Rv ATCC 27294 and *M. tuberculosis* H37Ra ATCC 25177 strains. Our cytotoxicity study on human adenocarcinoma cells suggests that both complex **1** and the ligand  $\text{H}_2\text{L}$  are able to induce cytotoxicity in MCF7 cells. In summary, the studies here discussed have evidenced the activity in the biomedical field of a new copper complex; further investigations in this area with other transition metal ions and chemosensor activities of the ligand for selective detection of metal ions are currently being carried out in our laboratories.

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