

Cite this: *Dalton Trans.*, 2025, **54**, 4187

## The relative impact of ligand flexibility and redox potential on the activity of Cu superoxide dismutase mimics†

Sharon Signorella,<sup>‡a</sup> Micaela Bruno,<sup>‡a</sup> Gianfranco Frattini,<sup>a</sup> Claudia M. Palopoli,<sup>a</sup> Diego M. Moreno,<sup>‡a</sup> Nora Pellegri,<sup>b</sup> Verónica A. Daier<sup>‡a\*</sup> and Sandra R. Signorella<sup>‡a\*</sup>

Two copper(II) complexes, [Cu(salbn)] and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub>, formed with the Schiff-base ligands 1,4-bis(salicylidenamino)butane (H<sub>2</sub>salbn) and 1,4-bis(pyridin-2-ylmethyleneamino)butane (py<sub>2</sub>bn), have been prepared and characterized in solid state and in solution, and their ability to catalyse the dismutation of O<sub>2</sub><sup>•−</sup> has been evaluated in homogeneous medium and immobilized in a mesoporous matrix. The crystal structures show that [Cu(salbn)] possesses a distorted square-planar geometry, while [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> adopts a *cis*-distorted octahedral geometry. The two complexes experience structural changes in solution, and different spectroscopies were used to examine them. Moreover, their redox potentials are strongly affected by the solvent. In water, the complexes exist as [Cu(salbn)(H<sub>2</sub>O)] and [Cu(py<sub>2</sub>bn)(H<sub>2</sub>O)]<sup>2+</sup> with Cu(II)/Cu(I) reduction potential at −361 mV and −229 mV, respectively, well different from redox potentials measured in acetonitrile. Although with a more unfavourable redox potential, [Cu(salbn)(H<sub>2</sub>O)] reacts with O<sub>2</sub><sup>•−</sup> faster than [Cu(py<sub>2</sub>bn)(H<sub>2</sub>O)]<sup>2+</sup>, with catalytic rate constants of 3.3 × 10<sup>7</sup> and 2.9 × 10<sup>7</sup> M<sup>−1</sup> s<sup>−1</sup>, respectively, at pH = 7.8. Both complexes exhibit higher superoxide dismutase activity than the analogues with a shorter central alkyl chain. The observed catalytic rates essentially correlate with the ligand flexibility, rather than with the redox potential, which is also supported by the slower O<sub>2</sub><sup>•−</sup> dismutation rate when the complexes are immobilized by encapsulation into the channels of well-ordered mesoporous SBA-15 silica where the pore modifies the complex structures and restraints the ligand rearrangement.

Received 7th December 2024,  
Accepted 31st January 2025

DOI: 10.1039/d4dt03403a

rsc.li/dalton

## Introduction

The superoxide radical (O<sub>2</sub><sup>•−</sup>) is a byproduct of respiration and a key component of the immune defence, which together with H<sub>2</sub>O<sub>2</sub>, constitute the primary reactive oxygen species (ROS) generated from the leakage of the electron transport chain during the oxidative phosphorylation in humans.<sup>1</sup> Under

normal conditions, the concentration of O<sub>2</sub><sup>•−</sup> is maintained at an optimal level by the release of the superoxide dismutase (SOD) enzyme.<sup>2</sup> Even when superoxide is a crucial biological messenger and essential antibacterial agent, under certain pathological conditions, like Parkinson's disease, arthritis, cancer, *etc.*, the O<sub>2</sub><sup>•−</sup> production overcomes the endogenous defences and tissues become vulnerable to damage.<sup>3,4</sup> The use of native SODs for the therapeutic treatment presents several drawbacks including their large size, poor stability under harsh reaction conditions, high cost and immunologic responses.<sup>5</sup> To surmount these limitations, efforts have been directed towards the design of low molecular weight metal-based antioxidant catalysts.<sup>6–9</sup>

Among SOD enzymes, copper–zinc SOD (CuZnSOD) catalyses O<sub>2</sub><sup>•−</sup> dismutation at a CuZn bimetallic centre through a ping-pong mechanism that involves the redox switch of the copper ion between Cu(II)/Cu(I) oxidation states, with Zn(II) engaged in a structural role contributing to the enzyme stability.<sup>10,11</sup> Since only the Cu ion participates in the redox cycle of O<sub>2</sub><sup>•−</sup> dismutation, a number of mononuclear Cu(II)

<sup>a</sup>IQUIR (Instituto de Química Rosario), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Facultad de Ciencias Bioquímicas y Farmacéuticas, Universidad Nacional de Rosario, Suipacha 531, 2000 Rosario, Argentina. E-mail: daier@iquir-conicet.gov.ar, signorella@iquir-conicet.gov.ar

<sup>b</sup>FIR (Instituto de Física Rosario), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Facultad de Ciencias Exactas, Ingeniería y Agrimensura, Universidad Nacional de Rosario, 27 de Febrero 210 bis, 2000 Rosario, Argentina

† Electronic supplementary information (ESI) available: X-ray crystallography summary tables, additional spectroscopic, kinetics, DRX and microscopic data, results from DFT calculations. CCDC 2405330 and 2405337. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4dt03403a>

‡ Equal contribution.

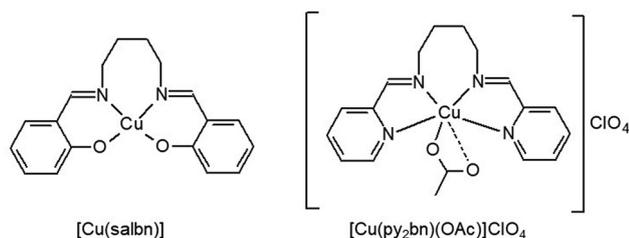


Fig. 1 Complexes studied in this work.

complexes with non-heme ligands bearing N/O donor sites, have been tested as SOD mimics.<sup>12–27</sup> It is currently accepted that copper centres with more flexible environment can better accommodate the structural rearrangements that take place when the metal switches between Cu(II) and Cu(I) oxidation states during  $O_2^{\cdot-}$  dismutation.<sup>28–31</sup> Also, it is known that the metal-centred redox potential is a crucial criterion for the activity of SOD mimics, and it indicates a thermodynamically favoured process when it falls between the one electron reduction and oxidation potentials of  $O_2^{\cdot-}$ .<sup>32</sup> However, little is known about the relative importance of these two factors on the activity of Cu SOD mimics. Moreover, the redox potential of the copper(II) complexes has rarely been measured in aqueous medium, even when it is well known that the solvent can substantially modify it.<sup>33,34</sup> In this work, we report the SOD activity, geometry and redox potential in aqueous solution of copper(II) complexes of two simple tetradentate Schiff-base ligands (Fig. 1), 1,4-bis(pyridin-2-ylmethyleneamino)butane ( $py_2bn$ ) and 1,4-bis(salicylideneamino)butane ( $H_2salbn$ ), bearing the same  $-(CH_2)_4-$  spacer between the two imino N-atoms and well different redox potentials in non-aqueous solvent, with the intention of ascertaining the relative influence of ligand flexibility and redox potential on the reactivity of the copper centre with  $O_2^{\cdot-}$ . Furthermore, in order to improve the complex stability and facilitate separation, recovery and handling, the complexes were immobilized by encapsulation into SBA-15 mesoporous silica. This strategy has been successfully employed to control the nuclearity, avoid undesired side reactions and replicate isolation and confinement of the metal centre as in the protein.<sup>7,35</sup> SBA-15 silica was selected because of its large surface area, highly ordered mesostructure, biocompatibility and open porosity that allows good accessibility of the substrate to interact with the catalyst.<sup>36,37</sup> The SOD activity of the heterogeneous catalysts was evaluated and differences in reactivity with respect to homogeneous complexes interpreted in terms of the geometrical changes that occur within the silica matrix.

## Experimental

### Materials

In this study, all the used reagents and solvents were commercial products of the highest available purity and, when necessary, were further purified by conventional methods.

### Synthetic procedures

Ligands  $N,N'$ -bis(2-pyridin-2-ylmethylene)butane-1,4-diamine ( $py_2bn$ ), and  $N,N'$ -bis(salicylidene)butane-1,4-diamine ( $H_2salbn$ ) were prepared as previously described by J. Lange *et al.*<sup>24</sup> and E. Kadwa *et al.*,<sup>38</sup> respectively.

**Synthesis of [Cu(salbn)].** A solution of  $Cu(OAc)_2 \cdot H_2O$  (216 mg, 1.08 mmol) in MeOH (20 mL) was added to a solution of  $H_2salbn$  (296 mg, 1.0 mmol) in MeOH (4 mL). The mixture was stirred at reflux for 30 min, and at room temperature for 2 h. Then, the volume was reduced and the solid was filtered, washed with cold MeOH and hexane, and dried under vacuum. Yield: 210 mg (0.58 mmol, 58%). Anal. calcd for  $C_{18}CuH_{18}N_2O_2$ : C 60.4, Cu 17.7, H 5.1, N 7.8%; found: C 60.3, Cu 17.3, H 5.5, N 7.8%. UV-vis,  $\lambda_{max}$  nm ( $\epsilon$   $M^{-1}$   $cm^{-1}$ ) in methanol: 240 (40 300), 273 (22 926), 296 (sh), 368 (9318), 634 (190); in buffer phosphate pH 7.8: 271 (27 140), 295 (sh), 355 (10 600), 631 (175). Significant IR bands (KBr,  $\nu$ ,  $cm^{-1}$ ): 3048, 3012, 2931, 2911, 2852, 1620, 1607, 1534, 1471, 1325, 1148, 760. Dark blue crystals suitable for X-ray diffraction were obtained from the reaction mother liquor after 3 days, at 4 °C.

**Synthesis of [Cu( $py_2bn$ )(OAc)]ClO<sub>4</sub>·1.5 H<sub>2</sub>O.** Powdered  $Cu(OAc)_2 \cdot H_2O$  (351 mg, 1.76 mmol) was slowly added to a solution of  $py_2bn$  (470 mg, 1.76 mmol) in 4.0 mL of methanol. The blue reaction mixture was left stirring for 24 h. Then,  $NaClO_4 \cdot H_2O$  (246 mg, 1.76 mmol) was added, and the mixture was stirred 1 day and left at 4 °C. A deep green solid formed and was filtered, washed with cold methanol and dried under vacuum. Yield: 654 mg (1.27 mmol, 72%). Anal. calcd for  $C_{18}H_{21}ClCuN_4O_6 \cdot 1.5 H_2O$ : C 41.9, Cu 12.3, H 4.7, N 10.9%; found: C 42.1, Cu 12, H 4.3, N 10.6%. UV-vis,  $\lambda_{max}$  nm ( $\epsilon$   $M^{-1}$   $cm^{-1}$ ) in methanol: 281 (20 224), 656 (151); in buffer phosphate of pH 7.8: 281 (15 360), 407 (sh), 720 (111), 865 (89). Significant IR bands (KBr,  $\nu$ ,  $cm^{-1}$ ): 3080, 2939, 1644, 1601, 1570, 1478, 1440, 1091, 625. Conductivity: 154  $\Omega^{-1}$   $cm^2$   $mol^{-1}$  in methanol. Green crystals suitable for X-ray diffraction were obtained from the reaction mother liquor after 3 days, at room temperature.

Caution! The perchlorate salts used in this study are potentially explosive and should be handled with care.

**Preparation of Cu-salbn@SBA-15.** SBA-15 mesoporous silica with high surface area was prepared by hydrothermal synthesis, using tetraethoxysilane (TEOS) as Si source and the triblock copolymer Pluronic 123 as template in acid medium, as already reported by M. Patriarca *et al.*<sup>35</sup> Insertion of complex [Cu(salbn)] into the mesoporous silica was carried out by slow addition of a solution of the complex (72 mg, 0.20 mmol) in methanol (20 mL, 35 °C) to SBA-15 silica (168 mg). The mixture was stirred for 24 h, filtered and the solid was suspended in 5 mL of methanol and left stirring for 24 h. The hybrid material was filtered, washed with methanol, and dried under vacuum. Yield: 140 mg. Anal. (wt%): Cu 0.32. Catalyst content: 5.2 mmol per 100 g. Significant IR bands (KBr,  $\nu$   $cm^{-1}$ ): 2966 (w, ligand), 2870 (w, ligand), 1640 ( $\delta$ , H–O–H), 1545 (w, ligand), 1460 (w, ligand), 1080 ( $\nu_{as}$ , Si–O), 795 ( $\nu_s$ , Si–O), 672 (w, ligand), 463 ( $\delta$ , Si–O–Si).

**Preparation of Cu-py<sub>2</sub>bn@SBA-15.** Following the same procedure, a methanol solution (20 mL) of complex [Cu(py<sub>2</sub>bn)]<sup>2+</sup> (98 mg, 0.19 mmol) was added to the silica (300 mg). The mixture was stirred for 24 h and filtered. The obtained material was suspended in 5 mL of methanol and left stirring for 24 h. The solid was filtered, washed with methanol, and dried under vacuum at room temperature. Yield: 354 mg. Anal. (wt%): Cu 1.78. Catalyst content: 28.1 mmol per 100 g. Significant IR bands (KBr,  $\nu$  cm<sup>-1</sup>): 2940 (w, ligand), 1640 ( $\delta$ , H–O–H), 1480 (w, ligand), 1450 (w, ligand), 1080 ( $\nu_{\text{as}}$ , Si–O), 795 ( $\nu_{\text{s}}$ , Si–O), 635 (w, ligand), 463 ( $\delta$ , Si–O–Si).

### Analytical and physical measurements

Infrared spectra were recorded in the 4000–400 cm<sup>-1</sup> range on a PerkinElmer Spectrum One FT-IR spectrophotometer provided with a DTGS detector, resolution = 4 cm<sup>-1</sup> and 10 accumulations. FT-IR spectra were registered from KBr sample pellets. Electronic spectra were recorded on a Jasco V-550 spectrophotometer. An Inductively coupled plasma mass spectrometer (ICP-MS) PerkinElmer NexION 350× was used to measure the metal content. For quantification of Cu in the hybrid materials, the samples (15 mg) were treated with 1 mL of H<sub>2</sub>SO<sub>4</sub> and 0.5 mL of HNO<sub>3</sub> and heated at 100 °C for 3 h. The solid was filtered and the filtrate diluted to a final volume of 25 mL. Cu in solution was analysed by ICP-MS. The copper content was verified by measuring the complex remaining in solution after the encapsulation. In this case, after filtering and washing the hybrid material, the filtrate was diluted to 250 mL. Then, 5 mL of this solution were diluted to 1 L with deionized water and Cu in solution was analysed by ICP-MS. Both methods afforded the same result. Each sample was prepared in triplicate. CHN analyses were performed on a PERKINELMER 2400 series II Analyzer. Electron Paramagnetic Resonance (EPR) spectra were obtained at 120 K on a Bruker EMX-Plus spectrometer with a microwave frequency of ~9.3 GHz. Simulations were performed in MATLAB using the EasySpin 6.0.6 program package for EPR.<sup>39</sup> <sup>1</sup>H NMR spectra were recorded on a Bruker AC 400 NMR spectrometer at ambient probe temperature (ca. 25 °C). Chemical shifts (in ppm) are referenced to tetramethylsilane and paramagnetic NMR spectra were acquired employing superWEFT sequence, with acquisition time of 80 ms. Conductivity measurements were performed on 1.0 mM solutions of the complexes using a Horiba F-54 BW conductivity meter. The electrochemical experiments were performed with a Princeton Applied Research potentiostat, VERSASTAT II model, with the 270/250 Research Electrochemistry Software. Studies were carried out with a standard three-electrode cell, under Ar, using 0.1 M Bu<sub>4</sub>NBF<sub>4</sub> as a supporting electrolyte and  $\approx 10^{-3}$  M of the complex. The working electrode was a glassy carbon disk, and the reference electrode was a saturated calomel electrode (SCE) isolated in a fritted bridge with a Pt wire as the auxiliary electrode. Under the employed conditions,  $E(\text{ferrocene/ferrocenium}) = 388$  mV in MeCN, 474 mV in DMF, 409 mV in MeOH (vs. SCE), at room temperature. Studies in aqueous buffered solutions were carried out on  $\sim 10^{-3}$  M of the complex in

50 mM buffer phosphate, pH = 7.8. Under these conditions, the reduction peak of the Zobell's solution = 185 mV vs. SCE. The size and morphology of the solid materials were analysed using an AMR 1000 Leitz scanning electron microscope (SEM) operated at variable accelerating voltages and with EDX detector NORAN System SIX NSS-200. Samples for SEM observation were prepared by dispersing a small amount of powder of dry silica and hybrid samples on double-sided conductive adhesive tabs on top of the SEM sample holders. Then, the samples were covered by a thin layer of gold deposited by sputtering to avoid charge accumulation on the surfaces. The selected accelerating voltage used in the shown images was 20 kV at high vacuum condition. Transmission electron microscopy (TEM) analysis was performed with a TEM/STEM JEM 2100 Plus with operational voltage of 200 kV (variable), with a LaB<sub>6</sub> filament. The samples were prepared by placing a suspension of silica and hybrid samples in ethanol onto a square mesh copper grid (400 mesh), coated with a layer of Formvar and carbon. The suspension of material in ethanol was left to dry, evaporating the ethanol, leaving the dispersed particles to adhere to the Formvar/Carbon surface. SEM and TEM images were processed using public domain ImageJ program. N<sub>2</sub> adsorption-desorption isotherms were obtained at 77 K on a Micrometric ASAP 2020 V4.02 (V4.02 G) apparatus. The samples were degassed at 10<sup>-3</sup> Torr and 200 °C for 6 h prior to the adsorption experiment. Surface area (SBET) was calculated using the Brunauer-Emmett-Teller (BET)<sup>40</sup> equation over the pressure range ( $p/p^0$ ) of 0.05–0.20. The volume of micropores and mesopores ( $V_{\text{MP}}$  and  $V_{\text{MP}}$ ) was determined by the alpha-plot method using the standard Licospher isotherm. The total pore volume ( $V_{\text{TP}}$ ) was determined with the Gurvich rule<sup>40</sup> at 0.98 $p/p^0$ . The pore size distributions were calculated using the Villarroel-Bezerra-Sapag (VBS) model,<sup>41</sup> on the desorption branch of the N<sub>2</sub> isotherms.

### Crystal data collection and refinement

Crystallographic data for compounds [Cu(salbn)] and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> were collected at 298(2) K on a Bruker D8 QUEST ECO Photon II CPAD Diffractometer, using graphite monochromated Mo-K $\alpha$  radiation ( $\lambda = 0.71073$  Å). Data collection was carried out using the Bruker APEX3<sup>42</sup> and the Bruker APEX4<sup>43</sup> packages, cell refinement and data reduction were done with the program SAINT V8.40B.<sup>44</sup> The structures were solved by direct methods with SHELXT V 2018/2.<sup>45</sup> and refined by full matrix least-squares on  $F^2$  data with SHELXL-2019/1.<sup>46</sup> Crystal data collection and the results of the refined parameters for compounds [Cu(salbn)] and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> are summarized in Table S1.† Molecular graphics were done with ORTEP-3,<sup>47</sup> with 50% probability displacement ellipsoids. Crystallographic data for [Cu(salbn)] and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> has been deposited at the CCDC under 2405330 and 2405337, respectively.†

### SOD activity measurement

The SOD activity of the free and immobilized complexes was determined by an indirect assay based on the inhibition of the

photoreduction of nitro blue tetrazolium (NBT).<sup>48</sup> In a typical experiment, the reaction mixture (3.2 mL) containing riboflavin (3.35  $\mu\text{M}$ ), methionine (9.52 mM), NBT (38.2  $\mu\text{M}$ ) and different concentrations of the complex either free (diluted aliquots (50  $\mu\text{L}$ ) of the complex in methanol added to the reaction mixture) or immobilized, was prepared in 50 mM phosphate buffer of pH 7.8. Riboflavin was added last and the mixture was illuminated for 15 min with a 20 W led lamp placed at a distance such that the rate of increase in absorbance was 0.017  $\text{min}^{-1}$ , at 25  $^{\circ}\text{C}$ . The reduction of NBT was measured at 560 nm. Control reactions were performed to verify that the complexes did not react with NBT or riboflavin directly. Inhibition percentage (IC) was calculated according to:

$$\text{IC} = \frac{[(\Delta\text{abs}/t)_{\text{without catalyst}} - (\Delta\text{abs}/t)_{\text{with catalyst}}]}{(\Delta\text{abs}/t)_{\text{without catalyst}}} \times 100$$

The  $\text{IC}_{50}$  values were determined from plots of % inhibition vs. catalyst concentration and used to calculate the McCord–Fridovich second-order rate constant ( $k_{\text{MCF}}$ ).<sup>49</sup> At 50% inhibition,  $\text{O}_2^{\cdot-}$  reacts with NBT and the mimic at the same rate, so  $k_{\text{MCF}} \text{IC}_{50} = k_{\text{NBT}} [\text{NBT}]$ , with  $k_{\text{NBT}}$  (pH = 7.8) =  $5.94 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ .

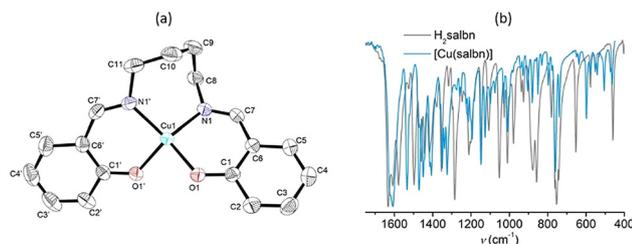
### Computational calculations

All calculations were performed at B3LYP/6-31G\*\* level<sup>50</sup> using Gaussian 16 package.<sup>51</sup> The solvent effects were computed by the polarizable continuum model (PCM)<sup>52</sup> with the DMF, DMSO or water dielectric constants. All the optimized structures are proved to be local minimum using frequency calculations.

## Results and discussion

### Geometrical and electronic properties of catalysts in solid state and solution

**Characterization of [Cu(salbn)].** The reaction of  $\text{H}_2\text{salbn}$  and  $\text{Cu}(\text{OAc})_2$  in refluxing methanol afforded [Cu(salbn)], which precipitated from the reaction mixture after concentrating and cooling. Dark blue crystals of [Cu(salbn)] suitable for X-ray diffraction studies formed from the mother liquors after several days. The molecular structure of [Cu(salbn)] is shown in Fig. 2(a) and relevant bond lengths and angles are given in

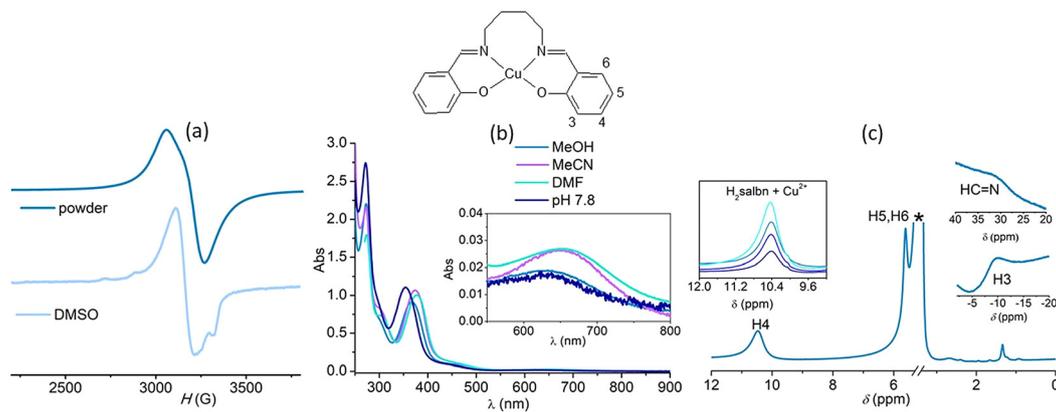


**Fig. 2** (a) Molecular structure of [Cu(salbn)] at the 50% probability level with atom numbering. (b) FT-IR spectra of  $\text{H}_2\text{salbn}$  and [Cu(salbn)].

Table S2.† The complex crystallizes in the *Pbca* space group as previously reported for crystals of this compound obtained by recrystallization from ethanol,<sup>53</sup> or by a high dilution and slow addition method, at room temperature,<sup>54</sup> but differs from the *P2<sub>1</sub>/c* space group of turquoise blue crystals obtained in aqueous medium.<sup>24</sup> In the complex, the Cu(II) ion is bound to the  $\text{N}_2\text{O}_2$  donor set of the deprotonated Schiff-base ligand, with Cu–N/O bond lengths in the range 1.898–1.988 Å typical of the Cu(II) oxidation state.<sup>55</sup> The twisted angle of 30° between the two phenolate planes as well as the value calculated for the four-coordinate geometry index  $\tau_4 = 0.44$ , (where  $\tau_4 = 0$  for square-planar and  $\tau_4 = 1$  for tetrahedral geometry)<sup>56</sup> indicate that the coordination geometry of the Cu centre may be described as distorted square-planar. The geometry around Cu(II) distinguishes from the essentially square-planar of [Cu(salpn)] (salpn = 1,3-bis(salicylideneamine)propane), for which  $\tau_4 = 0.13$  and O–Cu–N angles are closer to 180° (169 and 172°)<sup>57</sup> than in [Cu(salbn)]. Powdered samples and crystals of [Cu(salbn)] exhibit identical FT-IR spectra, both displaying the “fingerprint” pattern of the Cu(II) bound ligand. As shown in Fig. 2(b), the C=N stretching band at 1635  $\text{cm}^{-1}$  in  $\text{H}_2\text{salbn}$  shifts to 1627  $\text{cm}^{-1}$  in the complex as a consequence of the Cu(II) coordination to the imino N-atom, while the peak at 1379  $\text{cm}^{-1}$ , assigned to O–H in plane bending vibration of the phenol groups in the free ligand, is absent in the spectrum of the complex, as expected for deprotonated phenol groups bound to copper.

The low-temperature EPR spectrum of a polycrystalline sample of [Cu(salbn)] (Fig. 3(a)) shows an anisotropic signal for which the principal components of the *g*-tensor  $g_z = 2.20$ , and  $g_y = 2.06$ ,  $g_x = 2.05$  were obtained by simulation of the spectrum (Fig. S1(a)†). These values are consistent with the distorted square-planar geometry of the Cu(II) centre observed in the crystals. The lack of hyperfine structure in the powdered sample is the result of strong exchange interactions between neighbouring Cu(II) centres. In frozen DMSO glass, the EPR spectrum of [Cu(salbn)] shows axial anisotropy with the expected hyperfine splitting pattern for an  $I = 3/2$  nucleus of naturally abundant Cu along the direction parallel to the magnetic field (Fig. 3(a)). The simulation of the spectrum (Fig. S1(b)†) afforded spectral parameters  $g_{\parallel} = 2.20$ ,  $A_{\parallel} = 173 \times 10^{-4} \text{ cm}^{-1}$ ,  $g_{\perp} = 2.10$ , typical of Cu(II) complex possessing a  $d_{x^2-y^2}$  ground state. The calculated *f*-factor ( $g_{\parallel}/A_{\parallel}$ ) = 127 cm for this complex is in the range of 105 to 135 cm expected for tetrahedrally distorted square-planar geometry.<sup>58</sup> This tetrahedral distortion decreases as the central chain shortens. Therefore,  $f^{\text{Cu(salbn)}} > f^{\text{Cu(salpn)}} (119 \text{ cm in frozen DMSO})^{59} > f^{\text{Cu(salen)}} (108 \text{ in frozen DMF})^{60}$ , the last with the  $\text{CuN}_2\text{O}_2$  chromophore arranged almost in the plane.

The absorption spectra of [Cu(salbn)], shown in Fig. 3(b), exhibit an intense band at 273 nm and a shoulder around 295 nm corresponding to  $\pi \rightarrow \pi^*$  and  $n \rightarrow \pi^*$  intraligand transitions. The band at 355–377 nm experiences a bathochromic shift as the solvent donor capacity increases and can be assigned to the phenolate-to-copper(II) charge transfer (LMCT) process. The d–d transitions are observed as a broad absorp-

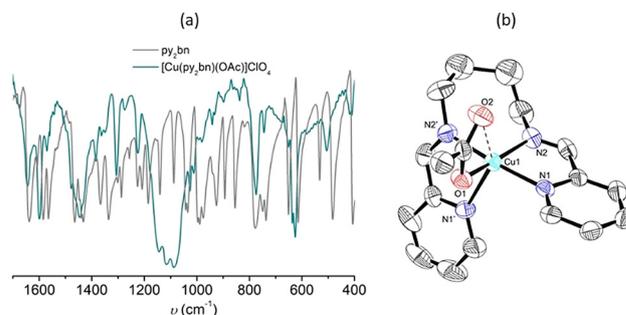


**Fig. 3** (a) X-band EPR spectra of powdered and frozen [Cu(salbn)] in DMSO;  $T = 120$  K;  $\nu = 9.31$  GHz. (b) Electronic spectra of 0.1 mM [Cu(salbn)] in different solvents. (c)  $^1\text{H}$  NMR spectrum of 10 mM [Cu(salbn)] in  $\text{D}_4$ -methanol. \*solvent. Inset: 9–12 ppm region of the  $^1\text{H}$  NMR spectrum of  $\text{H}_2\text{salbn}$  in  $\text{D}_4$ -methanol after addition of 0.2 (royal), 0.4 (blue), 0.8 (teal) and 1 (cyan) equivalent of  $\text{Cu}^{2+}$ .

tion band centred at 655 nm ( $\epsilon = 264\text{--}269$   $\text{M}^{-1} \text{cm}^{-1}$ ) in DMF and MeCN solution. This band appears at wavelengths longer than observed for [Cu(salpn)] (605 nm in toluene or  $\text{Cl}_3\text{CH}$ ) or [Cu(salen)] (565 nm in toluene),<sup>54</sup> consistent with a more distorted geometry around the copper ion in [Cu(salbn)], in agreement with the EPR results. The position of the d–d transitions varies with the solvent, and shifts to slightly shorter wavelengths (631–634 nm,  $\epsilon = 175\text{--}190$   $\text{M}^{-1} \text{cm}^{-1}$ ) in water or methanol, as shown in Fig. 3(b). Both, the lower energy and higher molar intensity of the d–d transitions support tetrahedral perturbation of the tetragonal Cu(II) environment in both DMF and MeCN greater than in methanol or water.<sup>33,60,61</sup>

Further evidence of the coordination of the ligand to Cu(II) is provided by the  $^1\text{H}$  NMR spectrum of the complex (Fig. 3(c)) that shows the broadening and shift of the resonances of the aromatic ring protons as well as those from the azomethine protons as a result of the paramagnetic relaxation induced by the Cu(II) ion. Aromatic protons undergo differential isotropic shift depending on the distance to the metal centre. Therefore, the broad signal at 31 ppm can be assigned to the azomethine protons, H4/H4' are observed at 10.5 ppm with peak half-width  $\omega = 189$  Hz, H5/H5' and H6/H6' protons appear overlapped at 5.6 ppm ( $\omega = 95$  Hz), and the resonance shifted up-field at -10 ppm can be attributed to H3/H3' on the basis of comparison with reported spectra for related Cu(II) complexes.<sup>62</sup> Given the large relaxation values of the phenolate proton signals, distortions from the square-planar geometry are not resolved and protons belonging to both aromatic rings cannot be distinguished. In addition, the coordination to the metal also affects the methylene resonances, displaying broad peaks that span in the 0.5–3 ppm range. The spectral assignment was verified recording  $^1\text{H}$  NMR spectra of  $\text{H}_2\text{salbn}$  after addition of increasing amounts of  $\text{Cu}(\text{ClO}_4)_2$ . The intensity of the broad resonances grows with increasing amounts of salt as a result of metal coordination (shown in the inset of Fig. 3(c) for the peak of H4), while those of the free ligand decrease and disappear. For the 1 : 1 metal-to-ligand mixture the spectrum is analogous to that of the solution of the complex.

**Characterization of [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub>·1.5H<sub>2</sub>O.** When the tetradentate ligand py<sub>2</sub>bn was reacted with one equivalent of  $\text{Cu}(\text{OAc})_2 \cdot \text{H}_2\text{O}$ , the complexation of the metal ion was evidenced by the deep green colour developed by the reaction mixture. Given the high solubility of the complex, addition of sodium perchlorate was necessary to induce the precipitation of the mononuclear copper complex [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub>·1.5H<sub>2</sub>O as a deep green solid, after 1 day at 4 °C. The FT-IR spectrum of this complex retains the pattern of the ligand and shows intense bands in the regions corresponding to the stretching frequencies of the acetate co-ligand and perchlorate counterion. The spectrum displays intense O–H (from solvation water) at 3434  $\text{cm}^{-1}$ , C=N and pyridine ring stretching vibrations at 1644 and 1601  $\text{cm}^{-1}$ , respectively, two strong absorption bands at 1091 and 625  $\text{cm}^{-1}$  corresponding to free perchlorate, and bands at 1652 (overlapped to the azomethine stretching band) and 1478  $\text{cm}^{-1}$  assignable to the asymmetrical and symmetrical stretching modes of bound acetate (Fig. 4(a)). This spectral pattern is retained in the FT-IR spectrum of crystals of [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> formed from the mother liquors (Fig. S2<sup>†</sup>), evidencing that the crystal structure is analogous to that of the powdered compound.



**Fig. 4** (a) FT-IR spectra of py<sub>2</sub>bn and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub>. (b) Molecular structure of the complex cation [Cu(py<sub>2</sub>bn)(OAc)]<sup>+</sup> at the 50% probability level.

The molecular structure of the complex cation determined by X-ray diffraction analysis is shown in Fig. 4(b), and selected bond lengths and angles are listed in Table S3.† The compound crystallizes in the  $P2_1/n$  space group with the asymmetric unit containing a discrete Cu(II) complex cation and one non-coordinated  $\text{ClO}_4^-$  anion. The  $\text{N}_4$ -tetradentate Schiff-base ligand is tightly bound to the Cu(II) ion *via* the two  $\text{N}_{\text{imine}}$  and two  $\text{N}_{\text{py}}$  atoms adopting a *cis-β*-configuration, in which the angle between the two aromatic ring planes is  $102.6^\circ$ . The two other coordination positions are occupied by acetate acting as a bidentate ligand bound to copper asymmetrically, with one short Cu1–O1 bond of  $1.934(5)$  Å and one long Cu1–O2 distance of  $2.754(5)$  Å. In line with this, the C11–O2 bond length is shorter than the C11–O1 one in the carboxylate group. The overall geometry may be described as *cis*-distorted octahedral<sup>63</sup> with large deviations of bond angles around Cu(II), up to  $24^\circ$  from  $90^\circ$ .

The low temperature X-band EPR spectrum recorded on the powdered complex (Fig. 5(a)) shows an anisotropic signal with  $g_1 = 2.05$ ,  $g_2 = 2.12$  and  $g_3 = 2.20$  (obtained from the simulated spectrum, Fig. S3(a)†) consistent with the rhombically distorted octahedral geometry around the Cu(II) ion observed in the crystal structure and also found in other  $\text{CuN}_4\text{O}_2$  chromophores with asymmetrically coordinated acetate groups.<sup>64,65</sup> The EPR spectra of the powder show only slight variation with increasing temperature from 120 K to 290 K, compatible with the retention of the *cis*-distorted octahedral configuration. In DMSO and 120 K, the spectrum is still broad and anisotropic, with  $g$  components at  $g_1 = 2.02$ ,  $g_2 = 2.05$ , and  $g_3 = 2.11$  (Fig. 5(a) and Fig. S3(b)†), suggesting that the rhombic distortion persists, probably with a less constrained geometry in solution. This was confirmed by optimizing the geometry of the complex using DFT calculations with DMF (as well as DMSO) as the implicit solvent, where the lowest-energy species corresponds to  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]^+$  retaining essentially the same geometry observed in the crystal (Table S3 and Fig. S4(a)†). Electronic spectra of  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]^+$  in different solvents are shown in Fig. 5(b). In the UV region the complex displays intense absorption bands around 280 nm originated from intraligand and charge transfer transitions, whereas in the visible region, the absorption features depend

on the solvent. In DMF, the spectrum displays a weak broad band centred at 608 nm ( $\epsilon = 206 \text{ M}^{-1} \text{ cm}^{-1}$ ), that can be assigned to d–d transitions in a distorted tetragonal environment. As observed in the inset of Fig. 5(b), the d–d transitions shift to the red in protic solvents. In methanol  $\lambda_{\text{max}} = 656$  nm with a shoulder at longer wavelengths, while in aqueous medium deconvolution of the d–d transitions afforded two bands at  $\lambda_{\text{max}} = 720$  and 865 nm. This trend suggests solvolysis of acetate in the protic solvents to form  $[\text{Cu}(\text{py}_2\text{bn})(\text{solv})]^{2+}$ . These two d–d bands are characteristic of the distortion of the Cu(II) centre towards trigonal bipyramidal geometry.<sup>66,67</sup> This geometry is in line with the preference for the trigonal bipyramidal coordination reported for the related complexes  $[\text{Cu}(6\text{-Me-py}_2\text{bn})(\text{H}_2\text{O})]^{2+}$ <sup>25</sup> and  $[\text{Cu}(\text{py}_2\text{bn})\text{Cl}]^+$ ,<sup>68</sup> with water/chloride occupying an equatorial position. Acetate dissociation in the protic solvent was also confirmed by the molar conductivity of  $154 \text{ } \Omega^{-1} \text{ cm}^2 \text{ mol}^{-1}$  of the complex measured in methanol, a value expected for a 1 : 2 electrolyte in this solvent.<sup>69</sup> The optimized structure of  $[\text{Cu}(\text{py}_2\text{bn})(\text{HOAc})]^{2+}$  in water as implicit solvent, shows that the Cu–O1 bond lengthens when acetate protonates and that the preferred structure for the protonated complex is a trigonal bipyramid with the HOAc coligand occupying an equatorial position (shown in Fig. S4(b) and Table S3†). DFT calculations show that the labilised HOAc ligand can be substituted by water when the entering water molecule approaches the edge opposite to the equatorial pyridine of the trigonal bipyramidal complex (the attack on another position did not lead to the substituted product), as can be seen in Fig. S5.†  $[\text{Cu}(\text{py}_2\text{bn})(\text{H}_2\text{O})]^{2+}$  retains the geometry with water placed in the equatorial plane and Cu–OH<sub>2</sub> bond distance of 2.085 Å. For comparative purposes, in  $[\text{Cu}(\text{py}_2\text{en})]^{2+}$  ( $\text{py}_2\text{en} = N,N'$ -bis(2-pyridin-2-ylmethylene)ethane-1,2-diamine), where the metal has a distinctively flat  $\text{CuN}_4$  plane as part of a tetragonally elongated octahedral geometry,<sup>70</sup> the d–d bands are centred at 612 nm in MeCN, while for  $[\text{Cu}(\text{py}_2\text{pn})(\text{DMF})]^{2+}$  ( $\text{py}_2\text{pn} = N,N'$ -bis(2-pyridin-2-ylmethylene)ethane-1,3-diamine) with a square-pyramidal structure in solution, the d–d transitions are centred at 626 nm in DMF.<sup>15</sup>

The  $^1\text{H}$  NMR spectrum of the paramagnetic  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]^+$  complex was recorded in  $\text{Cl}_3\text{CD}$  in the 50 to  $-50$  ppm spectral range. Due to the copper(II) paramagnetic effect, all

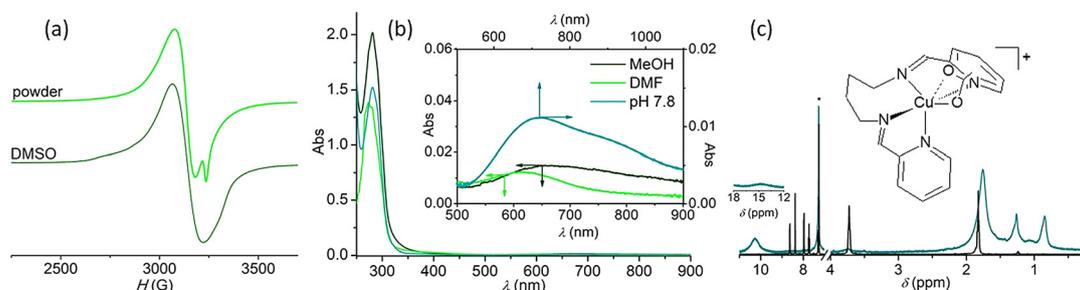


Fig. 5 (a) Low-temperature X-band EPR spectra of powdered and DMSO frozen solution  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]\text{ClO}_4$ ,  $\nu = 9.30$  GHz,  $T = 120$  K. (b) Electronic spectra of 0.1 mM solutions of  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]\text{ClO}_4$  in various solvents. (c)  $^1\text{H}$  NMR spectra of ligand (black line) and 15 mM complex (green line) in  $\text{Cl}_3\text{CD}$ ; \*solvent.

resonances are broadened to some extent, and broadening increases with the isotropic shift, so that a number of resonance lines become extremely wide precluding their detection. Fig. 5(c) shows the peaks that could be measured for this system using a superWEFT sequence with an acquisition time of 80 ms. The signals in the 0–2 ppm range can be assigned to the methylene protons of the alkyl chain of the ligand.<sup>71</sup> The resonance of the  $-(\text{CH}_2)_2-$  moiety at  $\delta = 1.75$  is slightly shifted from its position in the free ligand and broadens ( $\omega = 68$  Hz). The two up-field peaks at 1.26 ppm ( $\omega = 36$  Hz) and 0.84 ppm ( $\omega = 31$  Hz) can correspond to diastereotopic methylene  $-\text{CH}_2-\text{N}=\text{C}$ , while the broad resonance at 14.7 ppm ( $\omega = 650$  Hz) is compatible with the methyl acetate shifted down-field as a result of spin density delocalization through contact shift.<sup>72</sup> For the pyridine rings, it is expected that all protons exhibit a downfield shift from their respective positions in the free ligand arising from the positive spin density generated through the  $\sigma$ -metal-pyridine interactions, with downfield shifts for  $H_{\alpha\text{-py}}$  larger than for  $H_{\beta,\gamma\text{-py}}$ .<sup>73</sup> The broad signal observed in the spectrum at 10.3 ppm ( $\omega = 162$  Hz) could arise from the  $H_{\beta/\gamma}$  of one of the non-equivalent pyridine rings of the complex, the other resonances probably broaden beyond recognition under the experimental conditions used here or are outside the spectral sweep window.

### Electrochemical studies

The potential of the Cu(II)/Cu(I) redox couple depends on the metal first coordination sphere and the flexibility of the chelate ring, which in turn control the reactivity of the metal centre in electron transfer reactions. Therefore, the electrochemical behaviour of the complexes was studied by cyclic voltammetry in different solvents (Fig. 6). In  $[\text{Cu}(\text{salbn})]$ , the  $\text{N}_2\text{O}_2$  first coordination sphere of the metal stabilizes the Cu(II) oxidation state showing a quasi-reversible redox wave at  $-980$  mV vs. SCE in MeCN, characteristic of the Cu(II)/Cu(I) couple in an elongated tetragonal environment.<sup>33</sup> In DMF, the redox couple shifts to  $-901$  mV vs. SCE, most likely with DMF at the apex of a slightly distorted square-pyramid. The adoption of a flatter conformation with lowering in the flexibility of the ligand as the length of the alkyl spacer decreases is reflected in the more negative redox potential values reported for the Cu(II)/Cu

(I) redox couple of  $[\text{Cu}(\text{salpn})]$  ( $E_{1/2}^{\text{DMF}}$ :  $-1016$  mV vs. SCE) and  $[\text{Cu}(\text{salen})]$  ( $E_{1/2}^{\text{DMF}}$ :  $-1176$  mV vs. SCE).<sup>74</sup> In agreement with the spectroscopic results described above, the anodic and cathodic waves of the Cu(II)/Cu(I) couple of  $[\text{Cu}(\text{salbn})]$  shift to less negative potentials with the decreasing donor capacity of the solvent to bind copper at the apex of a square-pyramid (Fig. 6(a)).<sup>33</sup> Therefore,  $E_{1/2}^{\text{DMF}}$ :  $-901$  mV  $<$   $E_{1/2}^{\text{MeOH}}$ :  $-618$  mV  $<$   $E_{\text{pc}}^{\text{H}_2\text{O, pH 7.8}}$ :  $-361$  mV vs. SCE.

The absence of a redissolution peak in the anodic scan, indicates that, even in the aqueous buffer, no copper is released from the Cu(I) complex generated at the electrode. In the aqueous medium, the irreversibility of the process is probably related to the dissociation of the labile water molecule during reduction of  $[\text{Cu}(\text{salbn})(\text{OH}_2)]$  to form the negatively charged tetracoordinated Cu(I) complex  $[\text{Cu}(\text{salbn})]^-$  which is then re-oxidised at  $E_{\text{pa}}^{\text{H}_2\text{O}} = +33$  mV in the anodic scan.

In MeCN,  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]^+$  exhibits a quasi-reversible redox wave at 93 mV corresponding to the Cu(II)/Cu(I) couple (Fig. 6(b)). The large peak to peak separation  $\Delta E = 186$  mV, suggests that geometrical changes occur during the redox process. Probably the long Cu–N<sub>py</sub> bond breaks in the reduced form of the complex, retaining the bound acetate, so that Cu(I) can adopt a flattened tetrahedral geometry. Then, pyridine re-binds the metal in the oxidized complex to adopt the starting geometry. This electrochemical behaviour is different from that observed for  $[\text{Cu}(\text{py}_2\text{bn})]^{2+}$  in acetonitrile which shows a cathodic peak at 195 mV corresponding to the Cu(II)/Cu(I) reduction, and two oxidation peaks at 289 and 519 mV.<sup>68</sup> It has been reported that in MeCN, when the complex concentration is  $>5 \times 10^{-5}$  M,  $[\text{Cu}(\text{I})(\text{py}_2\text{bn})]^+$  is in equilibrium with the dimer  $[\text{Cu}(\text{I})_2(\text{py}_2\text{bn})_2]^{2+}$ , and the two observed anodic waves were assigned to  $[\text{Cu}(\text{I})(\text{py}_2\text{bn})]^+ \rightarrow [\text{Cu}(\text{II})(\text{py}_2\text{bn})]^{2+}$  and  $[\text{Cu}(\text{I})_2(\text{py}_2\text{bn})_2]^{2+} \rightarrow 2 [\text{Cu}(\text{II})(\text{py}_2\text{bn})]^{2+}$  oxidation processes, respectively.<sup>68</sup> The oxidation of the metal in the second process is accompanied by partial dissociation of the ligand so that oxidation of the Cu(I) centre in the dimer should require more energy than in the monomer. In the present study,  $[\text{Cu}(\text{py}_2\text{bn})(\text{OAc})]^+$  shows a similar electrochemical behaviour in methanol. As described before, in this protic solvent, acetate can protonate and dissociate affording  $[\text{Cu}(\text{py}_2\text{bn})(\text{MeOH})]^{2+}$  which is reduced at  $E_{\text{pc}} = 49$  mV, while  $[\text{Cu}(\text{I})(\text{py}_2\text{bn})]^+$  is oxidized at  $E_{\text{pa1}} = 183$  mV and  $[\text{Cu}(\text{I})_2(\text{py}_2\text{bn})_2]^{2+}$  at  $E_{\text{pa2}} = 319$  mV. A behaviour that also agrees with the molar conductivity of the complex in methanol. Additionally, the formation of the solvolyzed complex was verified by the identical voltammogram obtained for  $[\text{Cu}(\text{py}_2\text{bn})]^{2+}$  generated *in situ* by mixing  $\text{py}_2\text{bn}$  and  $\text{Cu}(\text{ClO}_4)_2$  in methanol. In water, acetate hydrolyses, and, as described previously, the aquo complex adopts a trigonal-bipyramidal geometry, with water bound to Cu(II) in the equatorial plane. Therefore, the reduction peak at  $E_{\text{pc}} = -229$  mV must involve dissociation of the water molecule from the meridional plane and geometrical rearrangement of the ligand, while the re-oxidation process of the tetracoordinated Cu(I) to the aquo-Cu(II) complex occurs at  $E_{\text{pa}} = 159$  mV. In water, no dimerization was observed. The redox potentials of the two complexes in different solvents are summarized in Table S4.†

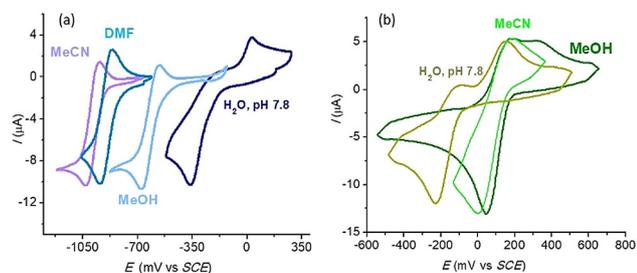


Fig. 6 Cyclic voltammograms of (a)  $[\text{Cu}(\text{salbn})]$  and (b)  $[\text{Cu}(\text{py}_2\text{bn})(\text{AcO})]^+$  in different media. Scan rate:  $100 \text{ mV s}^{-1}$ .

As expected, the  $N_2O_2$ -donor set of salbn stabilizes the Cu (ii) oxidation state more than the  $N_4$ -donor set of  $py_2bn$ . Therefore, in aqueous buffer,  $[Cu(salbn)]$  is reduced at a more negative potential than  $[Cu(py_2bn)]^{2+}$ . Anyway, it must be emphasized that in the aqueous buffer the redox potentials of both complexes lay between the redox couples  $E(O_2^{\cdot-}/H_2O_2) = 642$  mV and  $E(O_2/O_2^{\cdot-}) = -404$  mV vs. SCE, at pH 7,<sup>75</sup> and thus, they are suitable to act as SOD mimics.

### Characterization of the heterogeneous catalysts

Complexes  $[Cu(salbn)]$  and  $[Cu(py_2bn)]^{2+}$  were successfully encapsulated inside the channels of well-ordered mesoporous SBA-15 silica particles through wet impregnation, yielding Cu-salbn@SBA-15 and Cu- $py_2bn$ @SBA-15. The loading of the divalent complex cation was more than 5-times higher than the neutral complex as the result of the strong electrostatic interactions with the negative silanolate groups of the pores surface that step-up the retention of the cationic complex through ionic exchange with the silanol protons. In the FT-IR spectra (Fig. S6 and S7<sup>†</sup>) of both hybrid materials, the strong bands of the Si-O-Si framework and Si-O/OH<sub>2</sub> groups mask those from

the complexes, even in the case of the material where the catalyst is present in higher proportion. However, a few weak bands are not completely hidden by those of the silica matrix and can be observed in the 3000–2800, 1600–1350 and 700–600  $cm^{-1}$  spectral regions.

The textural properties of SBA-15 and the hybrid materials were analysed by nitrogen adsorption–desorption measurements at 77 K. All the samples exhibit type IV isotherms with an H1 hysteresis loop at relative pressure  $p/p^0 = 0.55$ –0.75 (Fig. 7(a and b)) typical of ordered mesoporous materials, and a steep step distinctive of the uniform mesopore size distribution. The plateau after the sharp step indicates the lack of significant secondary mesoporosity. Textural data of the mesoporous materials are summarized in Table 1. The prepared SBA-15 possesses a high specific surface area and pore width of 5.2 nm appropriate for hosting the complexes, which are  $\approx 0.93$ –1.25 nm wide calculated from the crystal structures described above.

The insertion of the catalyst causes the decrease of the BET surface area and total pore volume, indicating the compound is filling the pores, but leaves the overall shape of the BET

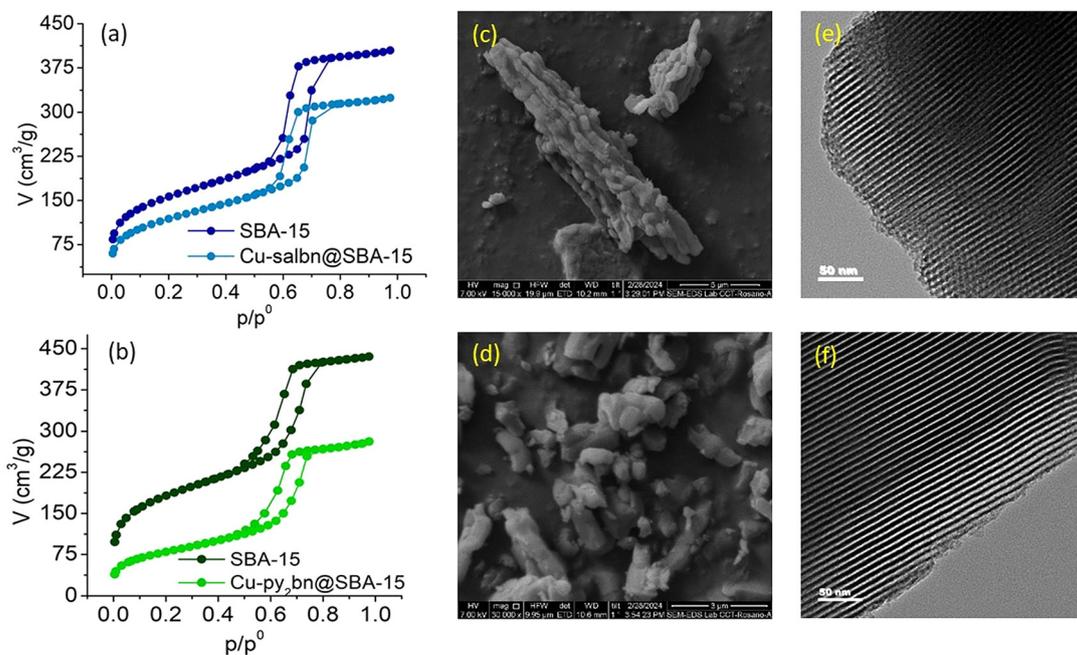


Fig. 7 Adsorption–desorption  $N_2$  isotherms of SBA-15 and (a) Cu-salbn@SBA-15 and (b) Cu- $py_2bn$ @SBA-15, at 77 K. SEM images of (c) Cu-salbn@SBA-15; 15 000x and (d) Cu- $py_2bn$ @SBA-15; 30 000x. TEM images of (e) Cu-salbn@SBA-15 and (f) Cu- $py_2bn$ @SBA-15.

Table 1 Textural characterization of mesoporous materials

	$S_{BET}$ ( $m^2 g^{-1}$ )	$V_{\mu P}$ ( $cm^3 g^{-1}$ )	$V_{MP}$ ( $cm^3 g^{-1}$ )	$V_{TP}$ ( $cm^3 g^{-1}$ )	$w_p$ (nm)	mmol complex per 100 g material
SBA-15	481	0.06	0.57	0.63	5.2	—
Cu-salbn@SBA-15	375	0.03	0.47	0.50	5.2	5.2
Cu- $py_2bn$ @SBA-15	261	0.01	0.37	0.43	5.4	28.1

$V_{TP} = V_{\mu P} + V_{primary MP} + V_{secondary MP}$ , MP = mesopore;  $\mu P$  = micropore;  $w_p$  = pore diameter.

curves almost unchanged, as well as the pore diameter, suggesting the ordered mesostructure of SBA-15 remains invariant, as confirmed through transmission electron microscopy (TEM) measurements described below. Cu-salbn@SBA-15 experiences a smaller decrease in the overall adsorption volume and surface area compared to Cu-py<sub>2</sub>bn@SBA-15, in line with the lower proportion of complex in the former, determined by ICP analysis.

The morphology and size of the particles of SBA-15 and the hybrid materials were analysed by scanning electron microscopy (SEM). SBA-15 particles possess elongated shapes of  $1.1 \pm 0.2$   $\mu\text{m}$  length with average aspect ratio of  $1.8 \pm 0.5$ , calculated from statistical analysis over 309 particles, that are lined up forming chains of 10–30  $\mu\text{m}$  long (Fig. S8(a) and S9(a)†). Particles of Cu-salbn@SBA-15 are somewhat smaller than SBA-15 ones and retain their shape and arrangement (Fig. 7(c)), whereas Cu-py<sub>2</sub>bn@SBA-15 develops entangled assemblies of more irregular spindly particles (Fig. 7(d)). The average aspect ratios calculated from *Image-J* analysis of SEM images of selected particles are  $2.0 \pm 0.4$  and  $1.8 \pm 0.5$  for Cu-salbn@SBA-15 and Cu-py<sub>2</sub>bn@SBA-15, respectively, and the corresponding histograms are shown in Fig. S9(b and c)†.

TEM images of the hybrid materials display a regular array of cylindrical channels along the long axis of the particles (Fig. 7(e and f)), confirming that incorporation of the complexes inside the pores preserve the highly ordered mesostructure of the SBA-15 (Fig. S8(b)†). Integration over the grey scales in carefully selected zones where the electron beam was perpendicular to the channels, afforded average pore diameter of  $4.5 \pm 0.6$  and  $4.3 \pm 0.9$  nm, and wall thickness of  $3.5 \pm 0.5$  and  $3.5 \pm 0.6$  nm, for Cu-salbn@SBA-15 and Cu-py<sub>2</sub>bn@SBA-15, respectively (Fig. S10†), which are in the order of pores size calculated from the adsorption isotherms. These channels reach the particle surface and adopt a hexagonal arrangement that can be observed in the images taken when the electron beam is parallel to the pores (Fig. S11†). This open porosity serves as an entry channel to the substrate to interact with the complex inside the pores, an essential aspect for catalysis.

### SOD activity studies

As described previously, in water the two complexes possess labile coordination sites and display redox potentials appropriate to dismutate O<sub>2</sub><sup>•−</sup>, so that they are expected to act as SOD mimics. Before carrying out the SOD activity tests, the stability of the catalysts in phosphate buffer was verified. For this, the free and immobilized complexes were incubated in a phosphate buffer of pH 7.8. UV-vis spectra of the free complexes recorded during an interval of 2 h showed constant spectral features and band intensities (Fig. S12†), evidencing the stability of the complexes in the reaction medium of the SOD assay. For the hybrid materials, the UV-vis spectra were registered on the supernatant after centrifugation of the suspensions. Spectra taken at different incubation times were compared to those of the free complexes in solution, verifying negligible complex leaching (Fig. S13†), even in the case of the neutral

complex. This behaviour highlights the efficiency of encapsulation that retains the complex inside the silica channels, and was confirmed by high angle X-ray diffractograms of Cu-salbn@SBA-15, which do not show peaks associated to crystallites of [Cu(salbn)] on the external surface of the silica particles (Fig. S14†).

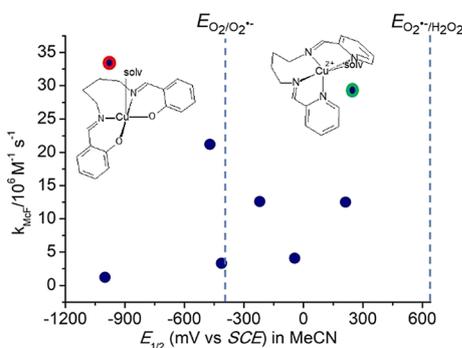
The SOD activity of free and encapsulated [Cu(salbn)] and [Cu(py<sub>2</sub>bn)]<sup>2+</sup> was examined by the Beauchamps and Fridovich indirect assay using nitroblue tetrazolium (NBT) in phosphate buffer of pH 7.8.<sup>48</sup> In this test, the catalyst competes with NBT for reacting with the photogenerated O<sub>2</sub><sup>•−</sup>, and the reduction of NBT by O<sub>2</sub><sup>•−</sup> leads to the formation of deep blue formazan determined at 560 nm after irradiating the reaction mixtures during a fixed time. In this way, the SOD activity of the catalysts is inversely related to the concentration of formazan, and the catalyst concentration required to decrease by 50% the reduction of NBT by O<sub>2</sub><sup>•−</sup> (IC<sub>50</sub>), can be determined from plots of % inhibition vs. [catalyst]. The IC<sub>50</sub> values calculated for [Cu(salbn)] and [Cu(py<sub>2</sub>bn)]<sup>2+</sup> are 0.068  $\mu\text{M}$  and 0.078  $\mu\text{M}$ , respectively. These values are dependent on the concentration and type of detector and cannot be contrasted with values reported for other synthetic catalysts. Therefore, the second-order rate constants  $k_{\text{MCF}} = k_{\text{NBT}}[\text{NBT}]/\text{IC}_{50}$ , which are independent of the detector, were calculated and used to compare the present results with others from the literature.<sup>76</sup> Both, [Cu(salbn)] and [Cu(py<sub>2</sub>bn)]<sup>2+</sup>, displayed excellent SOD activity (Fig. S15†), with  $k_{\text{MCF}}$  (Table 2, entries 2 and 3) higher than most of the reported Cu(II) complexes with open-chain N<sub>2</sub>O<sub>2</sub><sup>−</sup>, N<sub>3</sub>O<sup>−</sup>, and N<sub>4</sub>-tetradentate ligands, as shown in Table 2. Ligand structures of complexes listed in Table 2 are depicted in Chart S1. The selected Cu(II) catalysts are those for which  $k_{\text{MCF}}$  was reported or could be calculated from the IC<sub>50</sub> value.<sup>14,15,19–21,23,25–27,77,78</sup>

If the redox potential of the catalyst is one of the major features affecting the catalysed O<sub>2</sub><sup>•−</sup> dismutation, the compounds with  $E_{1/2}$  closer to 119 mV vs. SCE – the midpoint of the reduction/oxidation potentials of superoxide – should be among the most active ones. However, when the SOD activities of the complexes listed in Table 2 are plotted against their redox potentials in MeCN (the solvent used in the electrochemical studies of most of the listed complexes), no correlation is observed (Fig. 8). On the contrary, [Cu(salbn)], with  $E_{(\text{Cu(II)}/\text{Cu(I)})}$  well outside the potential range required for O<sub>2</sub><sup>•−</sup> dismutation, is the best SOD mimic of the set of  $\{k_{\text{MCF}}, E\}$  data pairs plotted in the graph. As shown before for the two complexes studied in this work, the solvent affects the geometry and properties of species in solution, in particular the redox potentials, and the magnitude of this effect depends on the electronic structure of the complex, and, if coordinated, the position occupied by the solvent in the coordination sphere.<sup>33,34</sup> In the present case, the redox potential of [Cu(salbn)] shifts by 600 mV to less negative potentials, while that of [Cu(py<sub>2</sub>bn)]<sup>2+</sup> shifts 474 mV to more negative values, when going from MeCN to water, resulting close reduction potentials for the two complexes in water, in spite of the large difference

**Table 2** SOD activity of [Cu(salbn)], [Cu(py<sub>2</sub>bn)]<sup>2+</sup> and other functional SOD models

Catalyst	Ligand donor sites	10 <sup>6</sup> <i>k</i> <sub>MCF</sub> (M <sup>-1</sup> s <sup>-1</sup> )	<i>E</i> (II/I) (mV vs. SCE), solvent	Ref.
1	[Cu(PuPhePy)] <sup>2+</sup>	N <sub>4</sub>	—	26
2	[Cu(salbn)(H <sub>2</sub> O)]	N <sub>2</sub> O <sub>2</sub>	−361 ( <i>E</i> <sub>pc</sub> ), H <sub>2</sub> O; −980, MeCN	This work
3	[Cu(py <sub>2</sub> bn)(H <sub>2</sub> O)] <sup>2+</sup>	N <sub>4</sub>	−229 ( <i>E</i> <sub>pc</sub> ), H <sub>2</sub> O; 245, MeCN <sup>66</sup>	This work
4	[Cu(MPBMPA)Cl <sub>2</sub> ]	N <sub>3</sub>	−471, MeCN	21
5	[Cu(pypapn)] <sup>2+</sup>	N <sub>4</sub>	−220 ( <i>E</i> <sub>pc</sub> ), MeCN	14
6	[Cu(PBMBA)Cl]	N <sub>3</sub> O	213, MeCN	21
7	[Cu(Pu-6-MePy)(H <sub>2</sub> O)] <sup>2+</sup>	N <sub>4</sub>	—	25
8	[Cu(aptn)] <sup>2+</sup>	N <sub>4</sub>	—	27
9	[Cu(py <sub>2</sub> pn)] <sup>2+</sup>	N <sub>4</sub>	−44, MeCN	15
10	[Cu(PCINOL)Cl] <sup>+</sup>	N <sub>3</sub> O	−413, MeCN	19 and 77
11	[CuL <sup>1</sup> ] <sup>2+</sup>	N <sub>4</sub>	—	20
12	[Cu(salpn)] <sup>c</sup>	N <sub>2</sub> O <sub>2</sub>	−1000, MeCN	This work
13	[Cu(4-OMe-salen)ZnCl <sub>2</sub> ]	N <sub>2</sub> O <sub>2</sub>	—	23
14	[salpnCuZnCl <sub>2</sub> ]	N <sub>2</sub> O <sub>2</sub>	−689, DMF	78
15	CuZnSOD	N <sub>4</sub>	156	2

aptn = [*N,N'*-butylenebis(2-acetylpyridineimino)]; HPBMPA = *N*-propanoate-*N,N*-bis-(2-pyridylmethyl)amine; HPCINOL = 1-[bis(pyridin-2-ylmethyl)amino]-3-chloropropan-2-ol; L<sup>1</sup> = *N*<sup>1</sup>,*N*<sup>2</sup>-bis(phenyl(pyridine-2-yl)-methylene)ethane-1,2-diamine; MPBMPA = *N*-methylpropanoate-*N,N*-bis-(2-pyridylmethyl)amine; Pu-6-MePy = *N,N'*-bis(2-(6-methyl-pyridyl)methylene)-1,4-butanediamine; PuPhePy = *N,N'*-bis(2-pyridyl-phenyl)methylene-1,4-butanediamine; pypapn = 1,3-bis[(pyridin-2-ylmethyl)(propargyl)amino]propane; py<sub>2</sub>pn = *N,N'*-bis(2-pyridinylmethyl)propane-1,3-diamine; 4-OMe-salen = *N,N'*-bis(4-methoxysalicylidene)cyclohexane-1,2-diamine; salpn = 1,3-bis(salicylideneamino)propane. <sup>a</sup> Previous values determined by the xanthine-xanthine oxidase method: *k*<sub>MCF</sub> = 26.4 M<sup>-1</sup> s<sup>-1</sup> at pH 7.4;<sup>79</sup> *k*<sub>MCF</sub> = 23.7 M<sup>-1</sup> s<sup>-1</sup> at pH 8.0.<sup>26</sup> <sup>b</sup> Using alkaline DMSP-NBT method, pH 8.6. <sup>c</sup> [Cu(salpn)] was prepared as described by Maurya *et al.*<sup>80</sup> and synthetic details are given in ESI.†



**Fig. 8** *k*<sub>MCF</sub> vs. redox potentials of complexes in Table 2. Lit. values (blue and blue/green dots). Dashed lines correspond to O<sub>2</sub><sup>•−</sup> reduction and oxidation potentials.

observed in MeCN. This means that comparing SOD activity in terms of redox potentials registered in non-aqueous solvents could lead to erroneous conclusions.

In addition to affecting the geometry or the distortion degree of the complex structure, the solvent may be involved in the solvolysis of a weakly bound ligand, as observed in the present work for [Cu(py<sub>2</sub>bn)(OAc)]<sup>2+</sup> in the protic solvents. Therefore, given that the redox potential depends on the geometry of the complex, solvent parameters and type of the ligands bound to the metal ion,<sup>33,34</sup> only the potentials measured in aqueous medium can be directly correlated to the observed SOD activity. In the present case, in aqueous medium [Cu(salbn)] reacts with O<sub>2</sub><sup>•−</sup> just 15% faster than [Cu(py<sub>2</sub>bn)]<sup>2+</sup> despite having a redox potential more distant from the optimal value to disproportionate O<sub>2</sub><sup>•−</sup> (119 mV vs. SCE). This implies that in

addition to the redox potential, there are other factors affecting the reaction of these complexes with O<sub>2</sub><sup>•−</sup>.

A feature that influences the SOD activity is the ability of the ligand to adopt different dispositions around the metal centre to best adapt to the preferred geometries of each oxidation state of copper during catalysis. For this reason, the more flexible ligands adapt better to the required conformational changes when going from Cu(II) to Cu(I). The flexibility of ligands used here depends largely on the length of the central chain. The two ligands with the -(CH<sub>2</sub>)<sub>4</sub>- linker between the imino N-atoms form flexible seven-membered chelate rings which can adopt different conformations with energy barriers lower than six-membered and five-membered ones. Therefore, [Cu(salbn)] reacts 27-times faster than [Cu(salpn)] (Table 2, entry 12) and 39-times higher than [CuZn(salpn (or salen)Cl<sub>2</sub>)] (Table 2, entries 13 and 14) formed with more rigid ligands with 3C- and 2C-central alkyl chains. Also, [Cu(py<sub>2</sub>bn)]<sup>2+</sup> reacts with O<sub>2</sub><sup>•−</sup> 7.2-times faster than [Cu(py<sub>2</sub>pn)]<sup>2+</sup> (Table 2, entry 9) and even more than twice as fast as [Cu(pypapn)]<sup>2+</sup> (Table 2, entry 5), in which copper is bound to 2N<sub>py</sub>2N<sub>amine</sub> donor set and possesses a 3C-central aliphatic chain. The methyl substituent on the pyridine ring in the trigonal-bipyramidal [Cu(Pu-6-MePy)(H<sub>2</sub>O)]<sup>2+</sup> or on the imino C-atom in distorted square-planar [Cu(aptn)]<sup>2+</sup> decreases the SOD activity about 5- and 6-times, respectively, compared to the unsubstituted [Cu(py<sub>2</sub>bn)]<sup>2+</sup> (Table 2, entries 7 and 8), probably because the methyl induces a decrease of the positive charge on the metal to interact with the substrate. The opposite effect is observed for [Cu(PuPhePy)]<sup>2+</sup> (Table 2, entry 1), a distorted square-pyramidal complex with solvent at the apex and the 4N-donor set at the base, where the phenyl substituent on the imino C-atom improves the activity compared to [Cu(py<sub>2</sub>bn)]<sup>2+</sup>, placing it as the most active Cu-based catalyst of

open chain ligands known so far. Complex  $[\text{CuL}^1]^{2+}$  is an analogue of  $[\text{Cu}(\text{PuPhePy})]^{2+}$  with a shorter  $-(\text{CH}_2)_2-$  central alkyl chain. The shortening of the aliphatic chain increases the complex rigidity that adopts a square-planar geometry and reduces its SOD activity drastically (Table 2, entry 11) compared to  $[\text{Cu}(\text{PuPhePy})]^{2+}$ .  $[\text{Cu}(\text{MPBMPA})\text{Cl}_2]$ ,  $[\text{Cu}(\text{PBMPA})\text{Cl}]$  and  $[\text{Cu}(\text{PCINOL})\text{Cl}]^+$  (Table 2, entries 4, 6 and 10) are three square pyramidal complexes in which the Cu atom sits in the plane formed by the 3N-donor set of the ligand and one chloride anion, with the apical position occupied by a second chloride anion, an O atom from the carboxylate of the ligand, or an alcohol group, respectively. In  $[\text{Cu}(\text{PBMPA})\text{Cl}]$ , the carboxylate acts as a competing ligand lowering SOD activity by almost half of  $[\text{Cu}(\text{MPBMPA})\text{Cl}_2]$ , for which the apical chloride is rapidly exchanged with the solvent leaving a labile position to interact with the substrate. This fact favours the reactivity of  $[\text{Cu}(\text{MPBMPA})\text{Cl}]^+$  with  $\text{O}_2^{\cdot-}$  and places this complex among the more active ones. The apical alcohol group in  $[\text{Cu}(\text{PCINOL})\text{Cl}]^+$  (Table 2, entry 10) further decreases the SOD activity, probably as a consequence of the lower labilising effect of the alcohol disfavoured ligand exchange.

Even when all the complexes listed in Table 2 are less reactive than the SOD enzyme (entry 15), it is evident that the ligand flexibility plays a critical role in the  $\text{O}_2^{\cdot-}$  dismutation catalysed by Cu-based SOD mimics, placing the two complexes of this work among the three most reactive catalysts with open-chain ligands.

In the enzyme, the protein environment isolates the active site and modulates its geometry so that the structural reorganization required to switch between Cu(II)/Cu(I) oxidation states is minimal, and the superoxide dismutation occurs efficiently at a rate close to the diffusion limit ( $\approx 10^9 \text{ M}^{-1} \text{ s}^{-1}$ ).<sup>2</sup> Keeping in mind the confining effect of the protein matrix and with the intention of improving the stability and reuse of the catalysts, the hybrid materials prepared by encapsulation of the catalysts inside the SBA-15 silica channels were tested as SOD mimics. Besides, isolation of the catalyst within the pores would avoid undesired dimerization that can occur in homogeneous medium during the catalytic cycles competing with the target reaction.<sup>81</sup>  $\text{Cu-salbn@SBA-15}$  retains SOD activity but reacts at a lower rate than the homogeneous catalyst, affording  $\text{IC}_{50} = 0.11 \mu\text{M}$ , and  $k_{\text{MCF}} = 20.6 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$  (Fig. S15(a)†). In the case of  $\text{Cu-py}_2\text{bn@SBA-15}$ , the decrease in rate is more evident (Fig. S15(b)†), giving  $\text{IC}_{50} = 0.28 \mu\text{M}$ , and  $k_{\text{MCF}} = 8.1 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ . The drop in activity might be related to a change in the geometry of the catalyst within the silica channels. This was confirmed by low-temperature X-band EPR spectra of the hybrids. The EPR spectrum of  $\text{Cu-salbn@SBA-15}$  exhibits an axial signal (Fig. 9(a), left) with spectral parameters  $g_{\parallel} = 2.17$  and  $g_{\perp} = 2.07$  obtained from the simulated spectrum (Fig. S1(c)†). These features are diagnostic of a tetragonal geometry around the Cu(II) centre within the SBA-15 pores, although less distorted than the complex in frozen DMSO solution. The EPR spectrum is broad probably as a result of the overlap of species with slightly different geometries or orientations within the SBA-15 matrix. The low-temperature EPR

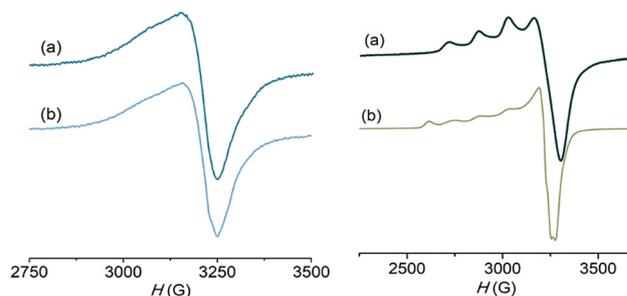


Fig. 9 Low-temperature EPR spectra of  $\text{Cu-salbn@SBA-15}$  (left) and  $\text{Cu-py}_2\text{bn@SBA-15}$  (right) before (a) and after (b) reaction with 10 equiv. of  $\text{KO}_2$  in DMSO.  $\nu = 9.31 \text{ GHz}$ ,  $T = 120 \text{ K}$ .

spectrum of  $\text{Cu-py}_2\text{bn@SBA-15}$  (Fig. 9(a), right) displays the typical signature of Cu(II) in a tetragonal geometry with  $g_{\parallel} = 2.25$ ,  $g_{\perp} = 2.07$ ,  $A_{\parallel} = 164 \times 10^{-4} \text{ cm}^{-1}$ , obtained from the simulated spectrum (Fig. S3(c)†), and an empirical distortion factor  $f(g_{\parallel}/A_{\parallel}) = 137 \text{ cm}$ , denoting a tetrahedrally distorted  $\text{N}_4$ -square-planar geometry. In this case, encapsulation constrains the ligand to dispose in the equatorial plane, instead to adopt the trigonal bipyramidal geometry observed in aqueous solution. The two hybrids react rapidly with  $\text{KO}_2$  at  $25 \text{ }^\circ\text{C}$  in DMSO, where  $\text{O}_2^{\cdot-}$  is a powerful reductant and nucleophile, and rather stable towards self-dismutation. When  $\text{Cu-salbn@SBA-15}$  was treated with excess of  $\text{KO}_2$  in DMSO, the EPR spectrum of the hybrid after reaction (Fig. 9(b), left) confirmed that the immobilized complex keeps the geometrical arrangement of the ligand around the Cu(II) ion. A different behaviour is observed for  $\text{Cu-py}_2\text{bn@SBA-15}$ , for which the low-temperature EPR spectrum taken after reaction with  $\text{O}_2^{\cdot-}$  still shows axial symmetry (Fig. 9(b), right), but well different spectral parameters:  $g_{\parallel} = 2.36$ ,  $g_{\perp} = 2.06$ ,  $A_{\parallel} = 154 \times 10^{-4} \text{ cm}^{-1}$ , and  $f = 153 \text{ cm}$ , obtained from the simulated spectrum (Fig. S3(d)†). These spectral parameters correspond to a species much more tetrahedrally distorted than that of the starting material. Although with a distortion similar to the enzyme (for the enzyme,  $f = 160 \text{ cm}$ ),<sup>82</sup> the species formed during the catalytic cycle is almost 3-times less reactive than the free complex probably because the silanolate-copper interaction restrains the ligand flexibility around the metal centre, resulting in a rate of reaction with  $\text{O}_2^{\cdot-}$  similar to that reported for  $\text{Cu-py}_2\text{pn@SBA-15}$  ( $k_{\text{MCF}} = 6.9 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ ).<sup>14</sup>

The absence of the superoxide signal ( $g_{\parallel} = 2.1021$ ,  $g_{\perp} = 2.003$ )<sup>83</sup> in the EPR spectra recorded a few seconds after reaction of the hybrids with 10-times excess of  $\text{KO}_2$  in DMSO, confirms that these materials act as catalysts for  $\text{O}_2^{\cdot-}$  dismutation. Also, the lack of EPR signal in the supernatant solution after treatment of the materials with excess of  $\text{KO}_2$  in DMSO indicates the absence of complex leached during the reaction with  $\text{O}_2^{\cdot-}$ . In this way, encapsulation confers stability and protection to the catalyst emulating the protein matrix in the enzyme, both for the neutral and cationic catalyst. Furthermore, to determine if the activity of the free and encapsulated catalysts was retained after several cycles, the conver-

sion of NBT was measured after successive illuminations of the reaction mixture with and without catalyst, adding after each new illumination the amount of NBT necessary to restore the initial concentration, so that the total NBT remained constant. When the homogeneous catalysts were used, the percentage of NBT converted to formazan decreased after each illumination period. In contrast, when the hybrid materials were used the NBT conversion remained almost invariant, indicating that immobilization prolongs the lifetime of the catalyst.

## Conclusions

This comparative study highlights that ligand flexibility has a more significant impact on SOD activity than the redox potential of the copper centre. For copper complexes of open-chain diimino ligands with terminal phenolate or pyridine, the flexibility increases with the length of the aliphatic linker between the imino N-atoms and the size of the chelate rings. Therefore, [Cu(salbn)(H<sub>2</sub>O)] and [Cu(py<sub>2</sub>bn)(H<sub>2</sub>O)]<sup>2+</sup>, with -(CH<sub>2</sub>)<sub>4</sub>- alkyl chain, react with O<sub>2</sub><sup>•-</sup> much faster than the analogues with shorter central alkyl chains. In addition, the effect of the solvent on geometry and redox potential of these complexes denotes it is essential to characterize them in aqueous medium to correlate their structure, redox potential and SOD activity. The aqueous buffered medium levels the redox potential of the two complexes studied here to values within the range required to catalyse O<sub>2</sub><sup>•-</sup> dismutation. However, [Cu(salbn)(H<sub>2</sub>O)], with a more unfavourable redox potential, reacts with O<sub>2</sub><sup>•-</sup> faster than [Cu(py<sub>2</sub>bn)(H<sub>2</sub>O)]<sup>2+</sup>, pointing out the strong influence of the flexibility of the adjacent six-seven-six-membered chelate rings of [Cu(salbn)(H<sub>2</sub>O)] to accommodate the Cu(II)/Cu(I) geometric preferences. This is supported by the effect of immobilization of the complexes in SBA-15 on the catalytic activity. Encapsulation of the complexes in the silica matrix not only modifies the complex structures but, probably, the pore restrains the ligand flexibility to encompass the switch between Cu(II)/Cu(I) oxidation states, decreasing the O<sub>2</sub><sup>•-</sup> dismutation rate. Nevertheless, although the two hybrids react with O<sub>2</sub><sup>•-</sup> slower than the free complexes, the confinement was successful in isolating, protecting and extending the useful life of the catalyst. Whereby, there must be a compromise between isolation and protection of the catalyst and restriction of the geometrical arrangement within the pore, and these two features should be considered to delineate heterogeneous SOD mimics.

## Data availability

The data supporting this article have been included as part of the ESI.†

Crystallographic data for [Cu(salbn)] and [Cu(py<sub>2</sub>bn)(OAc)]ClO<sub>4</sub> have been deposited at the CCDC under 2405330 and 2405337.†

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by the National University of Rosario (PIDs UNR 80020210200140UR, 8002019040023UR and 80020220700136UR), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET, PIP 0852 and PUE 0068), and Agencia Nacional de Promoción Científica y Tecnológica (ANPCyT, PICT-2019-03276). We thank Gustavo Terrestre for solving the crystal structures. The Bruker D8 QUEST ECO Photon II CPAD Diffractometer was purchased with funds from ANPCyT (PME 2015-0022) and CONICET.

## References

- 1 B. Yang, Y. Chen and J. Shi, *Chem. Rev.*, 2019, **119**, 4881–4985.
- 2 I. A. Abreu and D. E. Cabelli, *Biochim. Biophys. Acta*, 2010, **1804**, 263–274.
- 3 B. Halliwell and J. M. Gutteridge, *Free Radicals in Biology and Medicine*, Oxford University Press, 2015.
- 4 H. Sies, *Curr. Opin. Toxicol.*, 2018, **7**, 122–126.
- 5 R. Bonetta, *Chem. – Eur. J.*, 2018, **24**, 5032–5041.
- 6 G. Schanne, S. Demignot, C. Policar and N. Delsuc, *Coord. Chem. Rev.*, 2024, **514**, 215906.
- 7 C. Policar, J. Bouvet, H. C. Bertrand and N. Delsuc, *Curr. Opin. Chem. Biol.*, 2022, **67**, 102109.
- 8 S. Signorella, C. Palopoli and G. Ledesma, *Coord. Chem. Rev.*, 2018, **365**, 75–102.
- 9 I. Batinić-Haberle, J. S. Rebouças and I. Spasojević, *Antioxid. Redox Signal.*, 2010, **13**, 877–918.
- 10 Y. Sheng, I. A. Abreu, D. E. Cabelli, M. J. Maroney, A.-F. Miller, M. Teixeira and J. S. Valentine, *Chem. Rev.*, 2014, **114**, 3854–3918.
- 11 J. A. Tainer, E. D. Getzoff, K. M. Beem, J. S. Richardson and D. C. Richardson, *J. Mol. Biol.*, 1982, **160**, 181–217.
- 12 S. Pramanik and S. Chattopadhyay, *Inorg. Chim. Acta*, 2023, **552**, 121486.
- 13 M. A. Mekhail, K. J. Smith, D. M. Freire, K. Pota, N. Nguyen, M. E. Burnett and K. N. Green, *Inorg. Chem.*, 2023, **62**, 5415–5425.
- 14 M. Richezzi, J. Ferreyra, S. Signorella, C. Palopoli, G. Terrestre, N. Pellegri, C. Hureau and S. R. Signorella, *Inorganics*, 2023, **11**, 425.
- 15 M. Richezzi, J. Ferreyra, J. Puzzolo, L. Milesi, C. M. Palopoli, D. M. Moreno, C. Hureau and S. R. Signorella, *Eur. J. Inorg. Chem.*, 2022, e202101042.
- 16 Á. Martínez-Camarena, P. A. Sanchez-Murcia, S. Blasco, L. Gonzalez and E. Garcia-España, *Chem. Commun.*, 2020, **56**, 7511–7514.

- 17 J. D. Siqueira, S. F. de Pellegrin, S. S. dos Santos, B. Almeida Iglesias, P. C. Piquini, L. P. Arantes, F. A. Soares, O. A. Chaves, A. Neves and D. F. Back, *J. Inorg. Biochem.*, 2020, **204**, 110950.
- 18 K. N. Green, K. Pota, G. Tircso, R. A. Gogolak, O. Kinsinger, C. Davda, K. Blain, S. M. Brewer, P. Gonzalez, H. M. Johnston and G. Akkaraju, *Dalton Trans.*, 2019, **48**, 12430–12439.
- 19 T. P. Ribeiro, C. Fernandes, K. V. Melo, S. S. Ferreira, J. A. Lessa, R. W. A. Franco, G. Schenk, M. D. Pereira and A. Horn Jr, *Free Radicals Biol. Med.*, 2015, **80**, 67–76.
- 20 M. Choudhary, R. N. Patel and S. P. Rawat, *J. Mol. Struct.*, 2014, **1070**, 94–105.
- 21 J. S. Pap, B. Kripli, I. Bors, D. Bogáth, M. Giorgi, J. Kaizer and G. Speier, *J. Inorg. Biochem.*, 2012, **117**, 60–70.
- 22 C. Wang, S. Li, D.-J. Shang, X.-L. Wang, Z.-L. You and H.-B. Li, *Bioorg. Med. Chem. Lett.*, 2011, **21**, 4320–4324.
- 23 Z.-L. You, L.-L. Ni, P. Hou, J.-C. Zhang and C. Wang, *J. Coord. Chem.*, 2010, **63**, 515–523.
- 24 J. Lange, H. Elias, H. Paulus, J. Müller and U. Weser, *Inorg. Chem.*, 2000, **39**, 3342–3349.
- 25 J. Müller, D. Schübl, C. Maichle-Mössmer, J. Strähle and U. Weser, *J. Inorg. Biochem.*, 1999, **75**, 63–69.
- 26 J. Müller, K. Felix, C. Maichle, E. Lengfelder, J. Strähle and U. Weser, *Inorg. Chim. Acta*, 1995, **233**, 11–19.
- 27 Q. Luo, Q. Lu, A. Dai and L. Huang, *J. Inorg. Biochem.*, 1993, **51**, 655–662.
- 28 J. Stanek, A. Hoffmann and S. Herres-Pawlis, Renaissance of the entatic state principle, *Coord. Chem. Rev.*, 2018, **365**, 103–121.
- 29 B. A. Vaughn, A. M. Brown, S. H. Ahn, J. R. Robinson and E. Borosm, *Inorg. Chem.*, 2020, **59**, 16095–16108.
- 30 N. W. G. Smits, B. van Dijk, I. de Bruin, S. L. T. Groeneveld, M. A. Siegler and D. G. H. Hetterscheid, *Inorg. Chem.*, 2020, **59**, 16398–16409.
- 31 E. Falcone and C. Hureau, *Chem. Soc. Rev.*, 2023, **52**, 6595–6600.
- 32 C. Policar, in *Redox Active Therapeutics*, ed. J. S. Reboucas, I. Batinic-Haberle, I. Spasojevic, D. S. Warner and D. St. Clair, Springer, 2016, ch. 17, pp. 125–164.
- 33 G. Tabbi, A. Giuffrida and R. P. Bonomo, *J. Inorg. Biochem.*, 2013, **128**, 137–145.
- 34 P. A. Lay, N. S. McAlpine, J. T. Hupp, M. J. Weaver and A. M. Sargesons, *Inorg. Chem.*, 1990, **29**, 4322–4328.
- 35 M. Patriarca, V. Daier, G. Camí, N. Pellegri, E. Rivière, C. Hureau and S. Signorella, *Microporous Mesoporous Mater.*, 2019, **279**, 133–141.
- 36 A. Larki, S. J. Saghanzhad and M. Ghomi, *Microchem. J.*, 2021, **169**, 106601.
- 37 M. N. Iqbal, G. R. Nicoud, M. Ciurans-Oset, F. Akhtar, N. Hedin and T. Bengtsson, *ACS Appl. Mater. Interfaces*, 2023, **15**, 9542–9553.
- 38 E. Kadwa, M. D. Bala and H. B. Friedrich, *Appl. Clay Sci.*, 2014, **95**, 340–347.
- 39 S. Stoll and A. Schweiger, *J. Magn. Reson.*, 2006, **178**, 42–55.
- 40 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 41 J. Villarroel Rocha, D. Barrera and K. Sapag, *Top. Catal.*, 2011, **54**, 121–134.
- 42 Bruker, *APEX3 v2019.10-1*, Bruker AXS Inc., Madison, WI, USA, 2019.
- 43 Bruker *APEX4 v2022.10-1*, Bruker AXS Inc., Madison, Wisconsin, USA, 2022.
- 44 Bruker *SAINT V8.40B*, Bruker AXS Inc., Madison, Wisconsin, USA, 2019.
- 45 G. M. Sheldrick, SHELXT-Integrated space-group and crystal-structure determination, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 3–8.
- 46 G. M. Sheldrick, Crystal structure refinement with SHELXL, *Acta Crystallogr., Sect. B: Struct. Sci., Cryst. Eng. Mater.*, 2015, **71**, 3–8.
- 47 L. J. Farrugia, ORTEP3 for Windows, *J. Appl. Crystallogr.*, 1997, **30**, 565.
- 48 C. Beauchamps, Fridovich, *Anal. Biochem.*, 1971, **44**, 276–287.
- 49 Z.-R. Liao, X.-F. Zheng, B.-S. Luo, L.-R. Shen, D.-F. Li, H.-L. Liu and W. Zhao, *Polyhedron*, 2001, **20**, 2813–2821.
- 50 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision C.01*, Gaussian, Inc., Wallingford CT, 2016.
- 51 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 52 J. Tomasi, B. Mennucci and E. Cancès, *J. Mol. Struct.: THEOCHEM*, 1999, **464**, 211–226.
- 53 H.-H. Yao, J.-M. Lo, B.-H. Chen and T.-H. Lu, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 1997, **53**, 1012–1013.
- 54 L. C. Nathan, J. E. Koehne, J. M. Gilmore, K. A. Hannibal, W. E. Dewhirst and T. D. Mai, *Polyhedron*, 2003, **22**, 887–894.
- 55 H. Keypour, M. Shayesteh, S. Salehzadeh, S. Dhers, F. Maleki, H. Ünverç and N. Dilek, *New J. Chem.*, 2015, **39**, 7429–7441.

- 56 L. Yang, D. R. Powell and R. P. Houser, *Dalton Trans.*, 2007, 955–964.
- 57 H. Kanso, R. M. Clarke, A. Kochem, H. Arora, C. Philouze, O. Jarjayes, T. Storr and F. Thomas, *Inorg. Chem.*, 2020, **59**, 5133–5148.
- 58 S. Muthuramalingam, K. Anandababu, M. Velusamy and R. Mayilmurugan, *Inorg. Chem.*, 2020, **59**, 5918–5928.
- 59 K. Das, S. Goswami, B. B. Beyene, A. W. Yibeltal, E. Garribba, A. Frontera and A. Datta, *Polyhedron*, 2019, **159**, 323–329.
- 60 M. M. Bhadbhade and D. Srinivas, *Inorg. Chem.*, 1993, **32**, 5458–5466.
- 61 E. Movahedi and H. Golchoubian, *J. Mol. Struct.*, 2006, **787**, 167–171.
- 62 M. Bühl, S. E. Ashbrook, D. M. Dawson, R. A. Doyle, P. Hrobárik, M. Kaupp and I. A. Smellie, *Chem. – Eur. J.*, 2016, 15328–15339.
- 63 B. Murphy and B. Hathaway, *Coord. Chem. Rev.*, 2003, **243**, 237–262.
- 64 A. Gubendran, J. Rajesh, K. Anitha and P. Athappan, *J. Mol. Struct.*, 2014, **1075**, 419–429.
- 65 F. Clifford, E. Counihan, W. Fitzgerald, K. Seff, C. J. Simmons, S. Tyagi and B. J. Hathaway, *J. Chem. Soc., Chem. Commun.*, 1982, 196–198.
- 66 Y. Yao, M. W. Perkovic, D. P. Rillema and C. Woods, *Inorg. Chem.*, 1992, **31**, 3956–3962.
- 67 A. Kunishita, J. D. Scanlon, H. Ishimaru, K. Honda, T. Ogura, M. Suzuki, C. J. Cramer and S. Itoh, *Inorg. Chem.*, 2008, **47**, 8222–8232.
- 68 L. Llanos, C. Vera, A. Vega, D. Aravena and L. Lemus, *Inorg. Chem.*, 2020, **59**, 15061–15073.
- 69 W. J. Geary, *Coord. Chem. Rev.*, 1971, **7**, 81–122.
- 70 J. N. Melville and P. V. Bernhardt, *Inorg. Chem.*, 2021, **60**, 9709–9719.
- 71 S. Odisitse and G. E. Jackson, *Inorg. Chim. Acta*, 2009, **362**, 125–135.
- 72 I. Y. Ahmed and A. L. Abu-Hijleh, *Inorg. Chim. Acta*, 1982, **61**, 241–246.
- 73 J. M. Brink, R. A. Rose and R. C. Holz, *Inorg. Chem.*, 1996, **35**, 2878–2885.
- 74 M. K. Taylor, J. Reglinski, L. E. A. Berlouis and A. R. Kennedy, *Inorg. Chim. Acta*, 2006, **359**, 2455–2464.
- 75 P. M. Wood, *Biochem. J.*, 1988, **253**, 287–289.
- 76 S. Durot, C. Policar, F. Cisnetti, F. Lambert, J. P. Renault, G. Pelosi, G. Blain, H. Korri-Youssoufi and J. P. Mahy, *Eur. J. Inorg. Chem.*, 2005, 3513–3523.
- 77 A. C. Fernandes, G. L. Parrilha, J. A. Lessa, L. J. M. Santiago, M. M. Kanashiro, F. S. Boniolo, A. J. Bortoluzzi, N. V. Vugman, M. H. Herbst and A. Horn Jr., *Inorg. Chim. Acta*, 2006, **359**, 3167–3176.
- 78 V. A. Daier, E. Rivière, S. Mallet-Ladeira, D. Moreno, C. Hureau and S. R. Signorella, *J. Inorg. Biochem.*, 2016, **163**, 162–175.
- 79 M. Linss and U. Weser, *Inorg. Chim. Acta*, 1986, **125**, 117–121.
- 80 M. R. Maurya, S. J. J. Titinchi and S. Chand, *Appl. Catal., A*, 2002, **228**, 177–187.
- 81 C. Palopoli, G. Gómez, A. Foi, F. Doctorovich, S. Mallet-Ladeira, C. Hureau and S. Signorella, *J. Inorg. Biochem.*, 2017, **167**, 49–59.
- 82 K. M. Beem, W. E. Rich and K. V. Rajagopa, *J. Biol. Chem.*, 1974, **249**, 7298–7305.
- 83 R. N. Bagchi, A. M. Bond, F. Scholz and R. Stoesser, *J. Am. Chem. Soc.*, 1989, **111**, 8270–8271.