Synthesis, Characterization, EPR Spectroelectrochemistry Studies and Theoretical Calculations of Manganese(II) Complexes with the Ligands H₃bpeten and H₃bnbpeten

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Este trabalho relata a síntese e a caracterização dos complexos de manganês(II) [Mn^{II}(Hbpeten)] e [Mn^{II}(Hbnbpeten)], respectivamente [*N*,*N*'-bis-(2-hidroxibenzil)-*N*-(2-piridilmetil)-*N*'-(2-hidroxietil)etano-1,2-diamine]manganês(II) e [*N*,*N*'-bis-(5-nitro-2-hidroxibenzil)-*N*'-(2-hidroxietil) etano-1,2-diamin]manganês(II), por análise elementar, voltametria cíclica, espectroscopias eletrônica (UV-Vis) e vibracional (FTIR), ressonância magnética nuclear de hidrogênio (¹H NMR) e espectroeletroquímica de ressonância paramagnética eletrônica (EPR). Os dados de espectroeletroquímica de EPR foram consistentes com a redução de um grupo nitro, tanto no H₃bnbpeten livre quanto no complexo de manganês. Além disso, os cálculos teóricos foram consistentes com os resultados experimentais, mostrando que somente um grupo nitro contribui para a formação dos orbitais LUMO e sugerindo que os resultados teóricos são adequados para explicar as propriedades eletrônicas dos complexos.

The synthesis and characterization of the manganese(II) complexes [Mn^{II}(Hbpeten)] and [Mn^{II}(Hbnbpeten)], where H₃bpeten and H₃bnbpeten are respectively [*N*,*N*'-bis-(2-hydroxybenzyl)-*N*-(2-pyridylmethyl)-*N*'-(2-hydroxyethyl)ethane-1,2-diamine] and [*N*,*N*'-bis-(5-nitro-2-hydroxybenzyl)-*N*'-(2-hydroxyethyl)ethane-1,2-diamine], are reported. The characterization was carried out by elemental analyses, cyclic voltammetry, spectroscopic methods (UV-Vis, FTIR, ¹H NMR), electronic paramagnetic resonance spectroelectrochemistry (EPR) and theoretical DFT calculations. The electrochemistry and EPR espectroelectrochemistry data were consistent with the reduction of one of the nitro groups in free H₃bnbpeten and in the respective manganese(II) complex. These results were supported by DFT calculations, which showed that only one nitro group contributes to the LUMO. The theoretical data appear to be suitable to describe the electronic properties of the compounds.

Keywords: manganese complex, EPR, N,O-donor ligands, DFT, computational chemistry

Introduction

Reactions catalyzed by transition metal cations have attracted great interest in view of their applications in industrial processes.¹ In fact, several M-salen complexes, where M is manganese(III),² chromium(III)³ or nickel(II),⁴ have been utilized as homogeneous catalysts for olefin epoxidation.

The design of many manganese complexes and the investigation of their properties and applications (for example as catalysts) are bio-inspired. This is the case of several manganese complexes whose preparation aims to model the structure, reactivity and spectroscopic properties of biomolecules that contain this transition metal in its various oxidation states. In this way, complexes with different ligand types and nuclearities have contributed substantially to the understanding of the role of manganese in biological systems such as SOD (superoxide dismutase), the oxygen-evolving complex (OEC) of photosystem II (PS II)^{5,6} and catalases,⁷ among others. Furthermore, manganese complexes based on Schiff bases and other *N*,*O*-donor ligands have been explored as model systems of non-heme biological manganese sites. These models have proven to be suitable for studying the influence of

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the coordination geometry and electronic properties on the rich spectroscopic features of the biological Mn sites. In fact, such features and the chemical structures of the complexes were intensively studied in the past decades by different techniques, such as UV-Vis, FT-IR, EPR, XANES and EXAFS spectroscopies, X-ray crystallography, etc.^{8,9}

In view of the development of molecular mechanics and quantum chemical calculations, geometry optimization, molecular orbital energy and vibrational frequency calculations of metallocomplexes by DFT methods have turned into valuable tools for the description of molecular properties. In recent years, this method has been extensively used in biomolecular chemistry involving coordination compounds of the first and second transition metal series.¹⁰⁻¹³ Recently, a detailed computational study of O-O bond formation, catalyzed by monomeric and dimeric Mncorrole complexes, was carried out by the density functional theory (DFT) approach, in the light of biomimetic models of OEC.¹⁴

Our interest in the chemistry of the H_3 bpeten and H_3 bnbpeten ligands led to the preparation and characterization (electrochemistry, vibrational and electronic spectroscopies) of the respective copper(II) complexes. The use of $-NO_2$ electron-withdrawing groups in Hbnbpeten stabilizes the copper(II) oxidation state by several hundred millivolts, as confirmed by the anodic shift of the E_{pa} value, when compared to the analogous H_3 bpeten complex which has no substituent in the phenolic moieties.¹⁵

In order to investigate the effect of the ligand structure in the coordination geometry and oxidation state of the manganese ion, complexes bearing the H₂bpeten and H_bbnbpeten ligands were synthesized and shown to present an interesting catalytic activity in oxidation reactions.^{16,17} They were designed aiming the study of the effect of the $-NO_{2}$ group on the physicochemical properties of the manganese complexes. The H₂bnbpeten ligand is characterized by the presence of the electron-withdrawing nitro group, which should stabilize the manganese(II) species and make the synthesis of the complex much easier, as the greater acidity of H₂bnbpeten¹⁵ facilitates the generation and coordination of the deprotonated phenol group to the metal ion. Thus, this work reports the synthesis and characterization of the manganese-containing model compounds [Mn^{II}(Hbpeten)] ([N,N'-bis-(2-hydroxybenzyl)-N-(2-pyridylmethyl)-N'-(2-hydroxyethyl)ethane-1,2-diamine]manganese(II)) and [Mn^{II}(Hbnbpeten)] ([N,N'-bis-(5-nitro-2-hydroxybenzyl)-N'-(2-hydroxyethyl)ethane-1,2-diamine]manganese(II)) by spectroscopic and electrochemical methods (UV-Vis, FTIR, cyclic voltammetry, EPR and EPR spectroelectrochemistry),18 as well as theoretical calculations envisaging the investigation

of the influence of the frontier orbitals on the reactivity of the products.

Experimental

The following abbreviations were used throughout the text: H_3 bpten: N,N'-bis-(2-hydroxybenzyl)-N-2pyridylmethyl)-N'-(2-hydroxyethyl)ethane-1,2-diamine; H_3 bnbpeten: N,N'-bis-(5-nitro-2-hydroxybenzyl)-N'-(2-hydroxyethyl)ethane-1,2-diamine; Hpeten: N-(2pyridylmethyl)-N'-(2-hydroxyethyl)ethane-1,2-diamine; Et₃N: triethylamine; dmf: N,N-dimethylformamide; DMF and [TBA][PF₆]: tetra-n-butylammonium hexafluorophosphate and Fc⁺/Fc: ferrocenium/ferrocene redox pair.

Materials

2-pyridylcarboxyaldehyde, 2-bromomethylphenol, 2-hydroxy-5-nitrobenzaldehyde (5-nitrosalicylaldehyde), N-(2-hydroxyethyl)ethylenediamine, 2-hydroxy-5nitrobenzyl bromide (α -bromo-4-nitro-o-cresol), manganese(II) acetate and [TBA][PF₆] were obtained from Aldrich Chemical Co. For the electrochemical, spectroelectrochemical and spectroscopic studies, highpurity solvents were purchased from Merck and used as received. High-purity nitrogen gas was used to deoxygenate solutions. All other chemicals and solvents were reagent grade.

Syntheses

 $H_{2}bpeten$ (L1):^{16,17} This unsymmetrical compound was obtained in two steps, according to Scheme 1. The first step was a condensation reaction of N-(2hydroxyethyl)ethylenediamine (2.60 g; 25 mmol) with 2-pyridinecarboxyaldehyde (2.68 g; 25 mmol), followed by reduction with NaBH₄ (1.90 g; 50 mmol) in methanol, producing N-(2-pyridylmethyl)-N'-(2-hydroxyethyl) ethane-1,2-diamine (Hpeten). The desired unsymmetrical hexadentate proligand was synthesized by nucleophilic substitution of the 2-bromomethylphenol (3.74 g; 20 mmol) with Hpeten (1.95 g; 10 mmol), in 50 mL of THF, under argon atmosphere. After the addition of Et₂N (22 mmol) and stirring for 24 h, Et₂NHBr precipitated out and was removed by filtration. A yellowish oil was obtained after concentration of the filtrate in the flash evaporator, and the ligand was extracted with eight 50 mL portions of CHCl₂. The extracted volumes were combined, washed with brine, dried over anhydrous MgSO₄ and concentrated under reduced pressure. Then, the product was precipitated

out with acetone, filtered out, washed with propan-2-ol and dried under vacuum. H₃bpeten was obtained as a pale yellow solid. Yield: 3.20 g (78.5%), mp 157 °C. Anal. Calc. for $C_{24}H_{29}N_3O_3\cdot 2C_3H_8O$: C, 68.24; H, 7.01; N, 7.97%. Found: C, 68.2; H, 7.1; N, 7.5%. IR v_{max} /cm⁻¹: v(O-H), 3188; v(C=N, C=C), 1592, 1488, 1456 and 1438; δ (O-H), 1368; v(C-O), 1252. ¹H NMR (DMSO- d_6) δ 2.31 (4H, CH₂ alcohol), 3.34 (4H, NCH₂CH₂N), 3.73 (2H, NCH₂Py), 3.59 (4H, NCH₂Ph), 7.50-6.72 (11H, PhH and PyH), 8.45 (1H, PyH). No signal was found for the -OH groups.

*H*₃*bnbpeten* (*L2*):¹⁵ Also obtained according to Scheme 1, but using 2-hydroxy-5-nitrobenzyl bromide instead of 2-bromomethylphenol. H₃bnbpeten was obtained as a yellow solid. Yield: 3.18 g (91.4%), mp 208 °C. Anal. Calc. for C₂₄H₂₇N₅O₇: C, 57.95; H, 5.43; N, 14.08%. Found: C, 58.2; H, 5.3; N, 14.6%. IR v_{max}/cm⁻¹: v(O-H), 3478; v(C=N, C=C), 1.663, 1.597, 1.538 and 1.418; δ (O-H), 1343; v(C-O), 1280; δ_{assim}(NO₂), 1597; δ_{sim}(NO₂), 1280 and δ_{C-N} (ArNO₂), 829. ¹H NMR (DMSO-d₆) δ 2.21 (4H, CH₂ alcohol), 2.82 (4H, NCH₂CH₂N), 3.77 (2H, NCH₂Py), 3.72 (4H, NCH₂Ph), 7.08-7.22-7.69-8.56 (4H, PyH), 6.79-7.83-8.05 (6H, PhH). No signal was found for the -OH groups.

[$Mn^{II}(Hbpeten)$] (1): This coordination compound was prepared by refluxing Mn(CH₃COO)₂ (0.173 g, 1 mmol) and H₃bpeten (0.528 g, 1mmol) in methanol for 1 h, under magnetic stirring. Et₃N (1 mL, 7.18 mmol) was added to the reaction mixture to yield a brown precipitate that was filtered out and recrystallized in acetonitrile at room temperature. The complex was obtained as a brown solid after work out (Yield: 0.24 g; 52.17 %). Anal. Calc. for C₂₄H₂₇N₃O₃Mn: C, 62.55; H, 5.86; N, 9.12%. Found: C, 62.63; H, 5.95; N, 9.16%.

[$Mn^{II}(Hbnbpeten)$] (2): The complex was prepared by refluxing Mn(CH₃COO)₂ (0.173 g, 1 mmol) and $\rm H_3$ bnbpeten (0.497 g, 1 mmol) in methanol for 1 h, under magnetic stirring. Et₃N (1 mL, 7.18 mmol) was added to the reaction mixture to yield a brown precipitate, that was filtered out and recrystallized in acetonitrile at room temperature. The desired complex was obtained as a brown solid after work out (Yield: 0.32 g; 58.18%). Anal. Calc. for C₂₄H₂₅N₅O₇Mn: C, 52.36; H, 4.55; N, 12.73%. Found: C, 52.53; H, 4.58; N, 12.79 %.

Physical measurements

IR spectra were recorded on a FTIR BOMEN-MICHELSON spectrophotometer, model MB, from KBr disks. Elemental analyses were performed on a Perkin Elmer model 2400 equipment. The ¹H NMR spectra were obtained with a Bruker AC-200F spectrometer in DMSO- d_6 . EPR spectra were measured at 298 K and 77 K on a Bruker ESP 300E spectrometer operating at a frequency of about 9.5 GHz (X-band), with a modulation frequency of 100 kHz, modulation amplitude of 10 G to 0.1 G, depending on the sample (see text), and microwave power of ca. 20 mW. Handling of EPR spectra was carried out using Win-EPR[®] computer programs. The g-values were determined using a Bruker weak pitch sample to calibrate the spectrometer. Molar conductivity was measured in dmf solution (10^{-3} mol L⁻¹) at 298 ± 0.1 K with a Digimed DM-31 conductivimeter. Visible and NIR spectra were recorded in dmf with a SHIMADZU model 2401 spectrophotometer. Cyclic voltammetry experiments were performed with a PAR 273 (Princeton Applied Research) equipment in dmf at room temperature and under nitrogen. A 0.1 mol L⁻¹ solution of [TBA][PF₄] was employed as supporting electrolyte. The voltammograms were obtained by using a standard three-component



Scheme 1. Synthesis of Hbpeten and Hbnbpeten.

system consisting of a carbon disk working electrode, a platinum wire auxiliary electrode, and an Ag/AgCl reference electrode. The redox potential of the Fc+/Fc couple¹⁹ (found at +0.406 V vs. Ag/AgCl) was used to calibrate the reference electrode. Spectropotentiostatic experiments were simultaneously performed using an optically transparent thin-layer cell constructed according to a procedure described in the literature.^{18,20} Potentials were applied to the cell using a Microquímica (MQPG-01) potentiostat/galvanostat, and the spectra were collected on a Bruker ESP 300E spectrometer. The performance of the reference electrode was monitored by measuring the redox potential of the Fc⁺/Fc couple, which was found at +0.518 V vs. Ag/AgCl, both before and after the experiments. Spectra were registered for a selected series of applied potentials after the system attained equilibrium.

Theoretical calculations

The geometry optimization for H_3 bnbpeten was performed with the Gaussian 03W package, using the B3LYP hybrid functional and the 6-31G (d,p) and HF 6-31G (d,p) basis sets, on a personal computer equipped with a 3.0 GHz Intel Pentium D processor, 2 Gb of RAM memory and 160 Gb of hard disk space. Theoretical vibrational frequencies were calculated to check if the optimized structures presented true energy minima.

Results and Discussion

Syntheses

The syntheses of the unsymmetrical H_3 bpeten and H_3 bnbpeten molecules were conveniently carried out by reacting 2-bromomethylphenyl acetate and 2-hydroxy-5-nitrobenzyl bromide, respectively, with the diamine Hbpeten (Scheme 1). The characterization of H_3 bpeten and H_3 bnbpeten was unambiguously carried out by elemental analysis, IR and ¹H NMR spectroscopies. They were then allowed to react with Mn(CH₃COO)₂, in methanolic solution, to form the [Mn^{II}(Hbpeten)] (1) and [Mn^{II}(Hbnbpeten)] (2) complexes, which are stable in air and were isolated in good yields.

The IR spectra of **1** and **2** were similar to those of the free H₃bpeten and H₃bnbpeten. They differed only in the following: (*i*) the appearance of well defined band at 3424 cm⁻¹ (**1**) and 3447 cm⁻¹ (**2**), respectively, attributed to the v(O-H) stretching mode of the noncoordinated 2-hydroxyethyl group and to the water molecule present in complex (**3**); (*ii*) the absence of in-plane deformation bands of phenol groups, δ (O-H), that appear at 1368 cm⁻¹ (**L1**) and 1343 cm⁻¹ (**L2**) in the free bases, indicating coordination to the manganese(II) center as phenolate groups.

The molar conductivities of **1** and **2** in DMF solutions at 298 K were insignificant, being consistent with neutral coordination compounds.²¹

Electronic absorption

The electronic spectra of H₃bpeten (L1), H₃bnbpeten (L2) and the respective manganese(II) complexes were collected in DMF, and the transitions (λ_{max} /nm (ϵ /L mol⁻¹ cm⁻¹)) are listed in Table 1. L1 exhibits absorption bands at 234 (5692), 260 (5731) and 282 nm (4360), while L2 showed bands at 275 (21508) and 428 nm (34016). In contrast, the complex 1 exhibited a band at 267 nm (12218) while 2 showed bands at 264 (13024) and 389 nm (21803).

The bands observed in L1 and L2 were attributed to $\pi \rightarrow \pi^*$ transitions in the pyridyl and nitrophenyl groups. The presence of the electron withdrawing -NO₂ substituent in the phenol ring of H₂bnbpeten enhanced the molar absorptivity of the bands because it is in resonance with the aromatic ring. The absorption band observed for the [Mn^{II}(Hbpeten)] complex (1) was attributed to $\pi \to \pi^*$ transitions of the pyridyl and phenyl groups. In the case of the $[Mn^{II}(Hbnbpeten)]$ complex (2), the band at 264 nm can be attributed to $\pi \to \pi^*$ transitions of the pyridyl group, while the 389 nm band was tentatively assigned to a charge transfer transition (LMCT), which overlapps with the $\pi \to \pi^*$ transitions of the nitrophenyl groups of the ligand, thus explaining its high molar absorptivity. No *d*-*d* transition could be found in the UV-Vis spectra of compounds 1 and 2, as expected for high spin d⁵ complexes.

Electron paramagnetic resonance

The EPR spectra of manganese complexes can provide valuable information regarding oxidation state, type of ligand and site symmetry. As expected, no signal was found in the X-band EPR spectra of free L1 and L2 in DMF solution at room temperature, while six intense lines (Figure 1) were observed for compounds 1 and 2. These spectra are characteristic of monomeric manganese(II) complexes with octahedral geometry and axial symmetry. The Hamiltonian parameters (Table 1) obtained from the analyses of the spectra were $A_{iso} = 91.0 \text{ G}$ and $g_{iso} = 2.0059$, and $A_{iso} = 94.1 \text{ G}$ and $g_{iso} = 2.0081$ for complexes 1 and 2 respectively.

Electrochemistry

The redox properties of L1, L2 and the corresponding



Figure 1. X-band EPR spectra of 1 (a) and 2 (b) in DMF at 298 K. The Hamiltonian parameters are: $A_{iso} = 91.0$ G and $g_{iso} = 2.0059$ for 1; and $A_{iso} = 94.1$ G and $g_{iso} = 2.0081$ for 2.

manganese(II) complexes in dmf solution were investigated by cyclic voltammetry. The results are summarized in Table 1.

The electrochemical study of H₃bpeten and H₃bnbpeten was focused on the first one-electron oxidation of the phenolic units and the effect of the nitro group on the redox potential. This may allow to predict the easiness of one-electron oxidation of the corresponding manganese(II) complexes because the oxidation potential of the protonated free proligand is similar to that of the corresponding manganese(II) complex (protonation parallels metallation).²² L1 exhibits an irreversible oxidation wave at E_{pa} = 0.980 V vs. Ag/AgCl (0.462 V vs. Fc⁺/Fc and 0.862 V vs. NHE) and L2 a quasi-reversible wave at $E_{1/2}$ = 0.910 V vs. Ag/AgCl (0.392 V vs. Fc⁺/Fc and 0.792 vs. NHE). These processes correspond to the oxidation of the phenolic moieties leading to unstable radical cations, undoubtedly of the phenoxyl type. Interestingly, **L1** showed an anodic peak shifted by 0.070 V to more positive potentials as compared with **L2**, in spite of the electron-withdrawing nitro group bound directly to the phenol group, probably due to the irreversible character of that wave. The cyclic voltammogram of **L2** also presents a quasi-reversible redox couple at -1.781 V *vs*. Ag/AgCl (-2.336 V *vs*. Fc⁺/Fc and -1.936 V *vs*. NHE), that was tentatively attributed to the reduction of the nitro group present in the phenolic residues of H₃ bnbpeten.

The cyclic voltammograms of **1** and **2** in dmf at 100 mV s⁻¹ showed a quasi-reversible redox couple, respectively at 0.268 V and 0.165 V vs. Ag/AgCl, which can be ascribed to the $[Mn^{II}(HL)]/[Mn^{III}(HL)]^+$ redox couple. Compared with **1**, the presence of an electron-withdrawing group such as NO₂ in the coordination environment of **2**

Table 1. Spectroscopic and electrochemical data for the compounds H_3 beten (L1), H_3 bnbpeten (L2), $[Mn^{II}(Hbpeten)]$ (1) and $[Mn^{II}(Hbnbpeten)]$ (2)

Compounds	UV-Vis data ^a λ_{max} /nm (ϵ / L mol ⁻¹ cm ⁻¹)	EPR parameters ^b		Cyclic voltammetry data ^c	
		$\overline{A_{iso}}(G)$	g _{iso}	E _{pa} V vs. Ag/AgCl (V vs. NHE)	E _{1/2} V vs. Ag/AgCl (V vs. NHE)
H ₃ bpeten (L1)	234 (5 692) 260 (5 731) 282 (4 360)	silent	silent	0.980 (0.862)	-
H_3 bnbpeten (L2)	275 (21 508) 428 (34 016)	silent	silent	-	0.910 (0.792) -1.781 (-1.936)
[Mn ^{II} (Hbpeten)] (1)	267 (12 218)	91.0	2.0059	_	0.268 (0.150)
[Mn ^{II} (Hbnbpeten)] (2)	264 (13 024) 389 (21 803)	94.1	2.0081	_	0.165 (0.040) -1.835 (-1.966)

^a In DMF, 1×10^{-3} mol L⁻¹ for L1 and 1 and 1×10^{-4} mol L⁻¹ for L2 and 2; ^b In DMF solution, 1×10^{-3} mol L⁻¹, at 298 K; c. In DMF solution, 1×10^{-3} mol L⁻¹, under nitrogen, at 298 K, with 0.1 mol L⁻¹ [TBA][PF₆] as the supporting electrolyte. Standard three-component system: carbon disk working electrode, a platinum wire auxiliary electrode and an Ag/AgCl reference electrode. Fc⁺/Fc internal standard.

decreases $E_{1/2}$ by about 0.100 V and is a consequence of the higher basicity of the H₃bnbpeten ligand, which leads to a lower resistance to oxidation of complex **2**. The cyclic voltammogram of compound **2**, similarly to the observed for **L2**, presents a quasi-reversible redox couple at –1.835 V *vs*. Ag/AgCl (–2.366 V *vs*. Fc⁺/Fc and –1.966 V *vs*. NHE), that can be attributed to the reduction of the nitro group present in the phenolic residues of the H₃bnbpeten ligand.

EPR spectroelectrochemistry

EPR spectroelectrochemistry was carried out in DMF to give support to the assignment of the redox process observed for free and coordinated L2 at the -1.800 V region. The spectral changes are shown in Figures 2 and 3.

The EPR spectroelectrochemistry studies (simultaneous EPR and electrochemistry measurements using an adapted EPR cell, see Experimental) of the [Mn^{II}(Hbnbpeten)] complex (Figure 2a) showed no spectral change in the 0.0 to -0.5 V range, with the six characteristic manganese(II) lines remaining unmodified. However, when a potential of -2.0 V (Figure 2b) was applied, there was a fast decrease of the manganese(II) signals while a signal constituted by three sharp lines of similar intensity appeared, until finally only the new signal remained (Figure 2c).

The complete disappearance of the characteristic manganese(II) lines suggests that the metal center may be involved in the electrochemical process. In fact, the applied potential is sufficient to promote reduction of the hydrated Mn^{2+} ions to Mn^0 ($Mn^{2+} + 2e \rightarrow Mn^0$, $E^0 = -1.185$ V), leading to the deposition of the metal on the electrode and the disappearance of the EPR signal. The hypothesis of formation of manganese(III) complexes, that are EPR silent, can be ruled out because of the negative potential applied to the system.

The set of three signals can be assigned to a species with I = 1 and should not involve the metal center, as discussed above. In this case, some of the nitrogen atoms of the coordinated H₃bnbpeten should be near one unpaired electron. It is improbable that the two saturated nitrogen atoms coordinated to the metal ion are involved, since a signal constituted by five lines, in agreement with the 2n + 1 equation, is expected in this case. Accordingly, the unpaired electron should be localized on the nitro groups or on the pyridine nitrogen atoms.

The Hamiltonian parameters determined for the threeline signal are A = 13.7 G and g = 2.0048. However, a closer analysis showed that the signals are rather non-symmetric (Figure 3) and are probably constituted by a set of non-





Figure 2. EPR spectroelectrochemistry of the compound [Mn^{II}(Hbnbpeten)] (2), in DMF, 10^{-3} mol L⁻¹ {0.1 mol L⁻¹ [TBA][PF₆]}. (a) enhancement of the six line signal when the potential is stepped from 0.0 to -0.5 V; (b) decrease of the six line signal and appearance of a three line signal at -2.0 V vs. Ag/AgCl, and (c) disappearance of the six line signal and rise of the sharp three line signal at A_{150} =14.643 G after 30 min.

resolved hyperfine lines. Thus, the acquisition parameters of the equipment were adjusted (sweep width from 20 to 5 ms, time constant from 40 to 5 ms, conversion time from 20 to 5 ms and modulation amplitude from 10 to 1 G) in order to try to improve the resolution of the spectra. The result is shown in Figure 3. Note that each line was resolved in three other lines separated by 3.0 G, denoting the hyperfine interaction of one unpaired electron with two atoms with I = $\frac{1}{2}$. Finally, a fine tuning of the acquisition parameters (sweep width to 0.64 ms, time constant to 0.64 ms, conversion time to 0.64 ms and modulation amplitude to 0.1 G) allowed a further increase in resolution, now splitting each one of the three lines in two lines separated by 0.7 G, indicating the superhyperfine interaction with one atom of I = $\frac{1}{2}$, that could be one hydrogen atom.

The EPR spectroelectrochemistry study described above was extended to H_3 bpeten (L1) and the [Mn^{II}(Hbpeten)] complex (1). As expected from the cyclic voltammograms, no spectral change could be observed in the –1.800 V region, suggesting that the above described redox process is associated with the nitro group in the phenolic moiety. In the case of L2, as the molecule is unsymmetric, the nitro group at the same side of the pyridine group can probably be reduced before the one located at the side of the 2-hydroxyethyl residue.

the frontier orbitals at -0.337 (HOMO-1), -0.314 (HOMO), -0.207 (LUMO) and -0.205 hartree (LUMO+1), as well as the mapping of the electronic density distribution of these orbitals in the molecule. The HOMO is mainly localized on the ethylenediamine and saturated substituents, while the LUMO is centered on the nitrophenol group in the pyridyl group side and the LUMO+1 is mainly localized on the nitrophenol group in the hydroxyethyl side (Figure 4). Thus, a strong participation of the nitro group and the phenolate ring is suggested, in agreement with the EPR spectroelectrochemistry data. This result reinforces the hypothesis that the unpaired electron in the reduced free H_bbnbpeten and the respective manganese(II) complex is localized on the nitro group of the pyridyl side. Similar results were obtained using the HF method and the 6-31G (d,p) basis set, reinforcing the validity of our theoretical calculations.

The validity of the method was also confirmed by carrying out the theoretical calculation for the symmetric H_2 bbpen-NO₂ ([*N*,*N*[']-bis-(2-hydroxy-5-nitro-benzyl)-*N*,*N*[']-bis-(pyridine-2-methyl) ethylenediamine]) molecule using the DFT method and the same basis set. As expected, the LUMO orbital had a similar contribution from both nitrophenol groups (Figure 5b), in contrast with the H_3 bnbpeten ligand, where the contribution was asymmetric, in agreement with EPR spectroelectrochemistry results. On the other hand, the HOMO orbital (Figure 5a) in both the symmetric and asymmetric molecules showed a similar electronic density distribution over the molecule, with no significant contribution from the nitrophenol groups.

Theoretical study

The analyses of the frontier orbitals of H_3 bnbpeten using DFT calculations allowed the determination of the energy of



Figure 3. EPR spectroelectrochemistry of H₃bnbpeten in DMF, 1×10^{-3} mol L⁻¹ {0.1 mol L⁻¹ [TBA][PF₆]}. Applied potential of -1.80 V vs. Ag/AgCl, during 10 min.

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Figure 4. Graphic representation of the HOMO (a), LUMO (b) and LUMO+1 (c) calculated for H₃bnbpeten using DFT.



Figure 5. Graphical representation of the HOMO (a) and LUMO (b) calculated for H,bbpen-NO, using DFT.

Conclusions

Two Mn^{2+} complexes containing two polyfunctional ligands, one symmetric bearing two phenol moieties (L1) and one asymmetric, with electron-withdrawing $-NO_2$ groups bound to the phenol rings (L2), were prepared and characterized electrochemically and spectroscopically. The infrared, UV-Vis and EPR spectroscopy results, as well as the conductivimetry, elemental analysis and electrochemistry results were in agreement with the formation of manganese(II) complexes, whose proposed structures are presented in Figure 6.

The spectroelectrochemical studies of the free ligands and their respective manganese(II) complexes evidenced that the nitro group shifted the redox potentials by up to about 0.100 V, and was responsible for the appearance of a new quasi-reversible wave at -1.800 V, attributed to a



Figure 6. Proposed structures of complexes 1 and 2.

monoelectronic process involving the nitrophenol moiety. The localization of the electron on the nitro and nitrophenol groups in complex **2** and in **L2**, respectively, was confirmed by EPR spectroelectrochemistry.

Thus, a potential of -2.0 V was applied to both compounds at different intervals of time. After 30 s, a three-line signal that can be assigned to a free radical localized on the nitro group, appeared in the middle of the typical manganese(II) signal. In the same reducing conditions, the free L2 presented an eighteen-line spectrum, due to the electronic spin interaction with the nuclear spins of the nitrogen atom of the nitro group, of two orto-hydrogen atoms and of one hydrogen atom bound to the meta-carbon atom relative to the nitro group. Accordingly, the electrochemistry and EPR espectroelectrochemistry data are consistent with the reduction of the nitro group bound to one of the phenol moieties of the free H₂bnbpeten and the respective manganese(II) complex. This result was confirmed by DFT calculations that showed a major contribution of the nitrophenol group to the LUMO, indicating that the method can be used to predict some of the electronic properties of these compounds.

Supplementary Information

Supplementary information is available free of charge at http://jbcs.sbq.org.br, as PDF file.



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References

- 1. Niu, S.; Hall, M. B.; Chem. Rev. 2000, 100, 353.
- Srinivasan, K.; Michaud, P.; Kochi, J. K.; J. Am. Chem. Soc. 1986, 108, 2309.
- Samsel, E. G.; Srinivasan, K.; Kochi, J. K.; J. Am. Chem. Soc. 1985, 107, 7606.
- 4. Yoon, H.; Burrows, C. J.; J. Am. Chem. Soc. 1988, 110, 4087.
- Manchanda, R.; Brudvig, G. W.; Crabtree, R. H.; *Coord. Chem. Rev.* 1995, 144, 1.
- Su, J.-H.; Havelius, K. G. V.; Ho, F. M.; Han, G.; Mamedov, F.; Styring, S.; *Biochemistry* 2007, *46*, 10703.
- 7. Wieghardt, K.; Angew. Chem., Int. Ed. 1989, 28, 1153.
- 8. Dismukes, G. C.; Chem. Rev. 1996, 96, 2909.
- Wu, A. J.; Penner-Hahn, J. E.; Pecoraro, V. L.; *Chem. Rev.* 2004, 104, 903.
- Lanznaster, M.; Neves, A.; Bortoluzzi, A. J.; Assumpção, A. M. C.; Vencato, I.; Machado, S. P.; Drechsel, S. M.; *Inorg. Chem.* 2006, 45, 1005.
- 11. Scarpellini, M.; Casellato, A.; Bortoluzzi, A. J.; Vencato, I.;

Mangrich, A. S.; Neves, A.; Machado, S. P.; *J. Braz. Chem. Soc.* **2006**, *17*, 1617.

- Paes, L. W.; Faria, R. B.; Machuca-Herrera, J. O.; Neves, A.; Machado, S. P.; *Can. J. Chem.* **2004**, *82*, 1619.
- Zhan, C. G.; Nichols, J. A.; Dixon, D. A.; *J. Phys. Chem. A.* 2003, 107, 4184.
- Privalov, T.; Sun, L.; Åkermark, B.; Liu, J.; Gao, Y.; Wang, M.; *Inorg. Chem.* 2007, 46, 7075.
- Romanowski, S. M. M.; Tormena, F.; Santos, V. A.; Hermann, M. F.; Mangrich, A. S.; *J. Braz. Chem. Soc.* 2004, *15*, 897.
- Neves, A.; Romanowski, S. M. de M.; Bortoluzzi, A. J.; Mangrich, A. S.; *Inorg. Chim. Acta* 2001, *313*, 137.
- Romanowski, S. M. de M.; Mangrich, A.S.; Neves, A.; *Quim. Nova* 2001, 24, 592.
- Friedermann, G. R.; Halma, M.; Castro, K. A. D. F.; Benedito, F. L.; Doro, F. G.; Drechsel, S. M.; Mangrich, A. S.; Assis, M. D.; Nakagaki, S.; *Appl. Catal.*, A **2006**, *308*, 172.
- Gagné, R. R.; Koval, C. A.; Lisensky, G.C.; *Inorg. Chem.* 1980, 19, 2854.
- 20. Poll, N.; J. Chem. Educ. 1974, 67, 970.
- 21. Geary, W. J.; Coord. Chem. Rev. 1971, 7, 81.
- Ménage, S.; Gellon, G.; Pierre, J-L.; Zurita, D.; Saint-Aman,
 E.; *Bull. Soc. Chim. Fr.* **1997**, *134*, 785.

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