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ARTICLE

Making a next step toward inert Mn²⁺ complexes of open-chain ligands: the case of the rigid PhDTA[‡] ligand[†]

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In line with our research to find inert Mn(II) complexes as contrast agents for Magnetic Resonance Imaging, we have studied the aromatic–ring rigidified EDTA-analogue *o*-phenylenediamine-*N,N,N',N'*-tetraacetic acid (PhDTA). The protonation constants (K_i^H) of PhDTA and stability constants of complexes formed between this open-chain ligand and several different biogenic metal ions (Ca²⁺, Mg²⁺, Zn²⁺, Cu²⁺, Mn²⁺) have been determined in 0.15 M NaCl at 25 °C and compared with the values reported in the literature previously. Protonation constants are lower than those of the corresponding *cis*- and *trans*-CDTA complexes, which might be attributed to the electron withdrawing effect of the phenylene group. The lower total basicity of the ligand leads to lower stability constants for all the examined metal complexes. On the contrary, we have found that the conditional stability constants of [Mn(PhDTA)]²⁻ and [Mn(*trans*-CDTA)]²⁻ are approximately the same, as both complexes are 100% formed by pH 5 and their pM values are also comparable. The relaxivity of [Mn(PhDTA)]²⁻ is nearly identical ($r_1 = 3.72 \text{ mM}^{-1}\text{s}^{-1}$) to that determined previously for the [Mn(*trans*-CDTA)]²⁻ complex ($r_1 = 3.62 \text{ mM}^{-1}\text{s}^{-1}$), and its pH-dependence confirms the equilibrium model used for the fitting of the titration data. The results of kinetic studies of the metal exchange reactions reveal that the [Mn(PhDTA)]²⁻ complex possesses a slightly better dissociation kinetics profile than [Mn(*trans*-CDTA)]²⁻, which has been tested *in vivo* recently (including human injections). The half-life of the dissociation of the complex near to physiological pH at 25 °C is 19 hours. By using the rate constant calculated for the dissociation (pH=7.4, $c_{\text{Cu}^{2+}}=10 \text{ }\mu\text{M}$) and the half-life of excretion (1.6 hour), the ratio of the dissociated complex is estimated to represent 8% of the injected dose. DFT studies reveal that the metal coordination environment of [Mn(PhDTA)]²⁻ is very similar to that of [Mn(EDTA)]²⁻, both containing an inner-sphere water molecule. Cyclic voltammetry studies indicate that [Mn(PhDTA)]²⁻ is slightly more resistant towards oxidation to the Mn³⁺ complex than the EDTA analogue.

Introduction

The effectiveness of Gd³⁺-based contrast agents (CAs) used in Magnetic Resonance Imaging (MRI) examinations, such as Magnevist[®] or Dotarem[®], is undisputable.¹ However the toxicity of the free Gd³⁺ ion is high (LD₅₀ = 0.2–0.5 mmol/kg), and has been related to the development of Nephrogenic Systemic Fibrosis (NSF) by patients with reduced renal acute

functions (e.g. chronic kidney disease) after administration of CAs.² Because of these toxicity problems, there is a growing interest in the development of new, less toxic complexes containing biogenic metal ions which are likely better tolerated by living organisms. Some transition metal ions, which possess lower electron spin than Gd³⁺, can be used as the central core of a potential imaging agent. Such ions are for example high spin Mn²⁺ (d⁵),³ Fe³⁺ (d⁵),⁴ Co²⁺ (d⁷)⁵ or Ni²⁺ (d⁸).⁶

In the case of the Mn²⁺ ion, oxidation to Mn³⁺ caused by oxygen may lead to a drastic decrease of the paramagnetic effect. Thus, the redox potential of Mn²⁺/Mn³⁺ system has to be raised above +0.8 V by means of a suitable chelating ligand to obtain a contrast agent that is not sensitive to the oxidative conditions. On the other hand, the complexes undergoing oxidation in biological conditions may also be valuable for MRI imaging as redox-activated CAs, as it was demonstrated by Aime⁷ and more recently by Caravan and co-workers.⁸ The thermodynamic stability and kinetic inertness of the Mn²⁺ complexes are generally lower compared to the Gd³⁺ complexes. In this respect, to develop Mn²⁺ complexes for

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[†] Electronic Supplementary Information (ESI) available: Structures of the ligands mentioned in the text, cyclic voltammogram of [Mn(EDTA)]²⁻, bond distances of the metal coordination environment and optimised Cartesian coordinates obtained with DFT calculations.

[‡] PhDTA = *o*-phenylenediamine-*N,N,N',N'*-tetraacetic acid

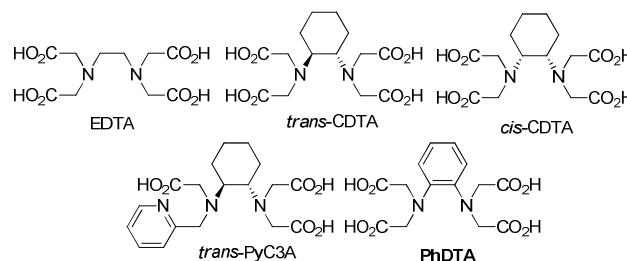
This paper is dedicated to professors Carlos F. G. C. Geraldes and Lothar Helm on the occasion of their retirement.

application as CAs is a great challenge, because it requires a good balance between the thermodynamic stability, relaxation and kinetic requirements, which are all crucial for *in vivo* MRI applications. There is only one commercially available (although already withdrawn from the European markets) non-Gd³⁺ based contrast agent, [Mn(DPDP)]⁴⁻ (Mangafodipir, Teslascan[®]), which is applied in MRI examinations of liver.⁹ The human body possesses mechanisms to eliminate the excess of Mn²⁺, but this works only in a limited concentration range (beyond that range it also possesses toxic effects with symptoms resembling Parkinson's disease).¹⁰

The properties of Mn²⁺ complexes formed with macrocyclic ligands (mostly derivatives of 9-aneN3, 12-aneN4, and pyridine containing macrocycles 12-PyaneN4, 15-PyaneN5 and 15-PyaneN3O2) synthesized and studied in the past 7-8 years predominantly by Éva Tóth and co-workers, see Figure S1) are also far from being optimal.¹¹⁻¹⁵ For instance Mn²⁺ complexes of macrocyclic ligands, such as NOTA and DOTA, possess high thermodynamic stability and suitable kinetic inertness for *in vivo* applications. However, the relaxivity (relaxation enhancement spurred by the presence of 1 mM paramagnetic substance) of these complexes is low because of the lack of the inner-sphere water molecule(s). On the other hand, coordinated water molecule(s) are observed in the complexes formed with ligands of lower denticity (e.g. ONO2A, NO2A, PC1A or PC1P). However, the decrease in the number of metal binding sites offered by the ligand affects negatively the kinetic inertness and the stability of the Mn²⁺ chelates. An acceptable balance of physicochemical properties (stability, redox potential, inertness and relaxivity of Mn²⁺ complexes) was found for *cis*-DO2A¹⁶ and its derivatives,¹⁷ opening a passable avenue to Mn²⁺-based MRI CAs. The Mn²⁺ complexes formed with linear ligands have almost the same thermodynamic stability as the corresponding macrocyclic ones, but their kinetic inertness are several orders of magnitude lower.^{12,13} For instance, while the [Gd(DTPA)]²⁻ complex presents a half-life of 33 hours at pH 5, the dissociation of [Mn(DTPA)]³⁻ is so fast at this pH that prevents the determination of the dissociation rate even with the stopped-flow technique (with a dead time of 8 ms).¹⁴ At the same time, linear ligands represent some advantages over the macrocyclic ones, for instance because their synthesis is less challenging. Moreover, careful ligand design can supply candidates with improved stability,¹⁸ and relaxivity,¹⁹ owing to the planned interaction of the complexes with biomolecules such as HSA,¹⁹⁻²¹ or even Mn²⁺-based responsive probes relying on the use of bifunctional ligands.²² Among the Mn²⁺ complexes studied so far [Mn(*trans*-CDTA)]²⁻ and related *trans*-CDTA derivatives (e.g. PyC3A) show by far the best kinetic parameters.²²⁻²⁵ More recently, these complexes were successfully tested *in vivo* in mice, in dogs and surprisingly even in human studies confirming their applicability.²⁶

The objective of this study is to determine the stability constants of [M(PhDTA)]²⁻ complexes and compare them with the stabilities of the complexes formed with the structurally-related EDTA, *cis*- and *trans*-CDTA ligands (Scheme 1). The PhDTA ligand has an even more rigid structure than CDTA due

to the presence of an *ortho*-phenylenediamine unit in the molecule. In the Mn²⁺ complexes of the *cis*- and *trans*-CDTA ligands the two N atoms of the iminodiacetate groups define N-C-C-N angles of ~50°,²⁷ while the unsaturated nature of the phenyl unit of PhDTA forces a planar N-C-C-N entity. As a result, the structure of the Mn²⁺ complex of PhDTA is expected to be somewhat different to that of [Mn(*trans*-CDTA)]²⁻, which in turn may lead to differences between the thermodynamic and kinetic properties of the two complexes. We do not expect a drastic change in relaxivity because of the presence of a coordinated water molecule and the identical overall charge of both complexes.²⁸ The thermodynamic properties of the PhDTA ligand and several of its metal complexes were studied in the 1980's and 1990's, but because of the differences in terms of the applied procedures (e.g. calorimetry, extraction method, etc.) and ionic strengths (1.0 M NaClO₄), as well as due to the large differences in the reported stability data (for [Cu(PhDTA)]²⁻ a difference of 10 log units can be found in the literature),²⁹⁻³³ we decided to determine these values using an ionic strength close to the physiological conditions (0.15 M NaCl). Therefore, we synthesized the PhDTA ligand and redetermined the stability constants of the Mn²⁺ complex and those with other biogenic metal ions. Furthermore, we investigated the dissociation kinetics, relaxometric and electrochemical properties of the [Mn(PhDTA)]²⁻ complex. Finally, the structure of the [Mn(PhDTA)]²⁻ complex in solution was studied by using DFT calculations.



Scheme 1. Ligands discussed in the present work.

Results and discussion

Equilibrium and Relaxometric Studies

The protonation constants of the PhDTA ligand must be determined to subsequently measure the stability constant of the [Mn(PhDTA)]²⁻ complex. For this purpose direct pH-potentiometric titrations have been carried out. During these studies it is essential to choose an appropriate ionic strength because the ions used to adjust the ionic strength interact with both the ligand and metal ion. We have chosen a 0.15 M NaCl ionic strength to model the NaCl background present in the body fluids. The protonation constants obtained by potentiometry are shown in Table 1 along with the ones determined previously using different ionic strengths^{30,31,33} and the corresponding values of the EDTA and *cis*- or *trans*-CDTA chelators.²⁷ The comparison of the constants excels that the log K_i^H values of PhDTA are lower than those of CDTA

ligands, with the total basicity of PhDTA being four orders of magnitude lower. In these ligands (EDTA, *cis*- and *trans*-CDTA and PhDTA) the first two protonation processes are attributed to the two amine nitrogen atoms, while the protonations occurring at acidic pH correspond to the acetate groups. The considerable difference between the first two protonation constants can be ascribed to the lower basicity of anilines compared to aliphatic amines, which is the result of delocalization of the nitrogen atom's lone pair onto the aromatic unit. The 3rd and the 4th protonation constant of the PhDTA chelator can be assigned to the protonation of carboxylate pendants while 5th and 6th protonation cannot be measured by pH-potentiometry. This suggests that these protons are very acidic and dissociate easily in aqueous media, similarly to the protons of strong acids. By comparing the protonation constants with the previously reported ones, we did not find significant deviations. The small differences may arise from the applied ionic strength and less likely from different measurement techniques.

In view of the protonation constants determined here, the stability and protonation constants of PhDTA complexes formed with biogenic metal ions are not expected to deviate notably from the literature data. The acquired stability constant values are also shown in Table 1, where the data reported previously for PhDTA, EDTA and CDAs are provided for comparison. As expected from the total basicity values, the stability constants of the PhDTA complexes are significantly lower compared to the EDTA and CDTA analogues. In the cases of Mg²⁺, Ca²⁺ and Mn²⁺ the stability constants drop by 3-4 orders of magnitude, but the decrease of the stability constants of the PhDTA complexes with Zn²⁺ and Cu²⁺ exceeds 6 orders of magnitude. However, the species distribution curves of [Mn(PhDTA)]²⁻ and [Mn(*trans*-CDTA)]²⁻ complexes (Figure 1) indicate that complex formation begins at pH 2.0 and ends at around pH 5.0 in both systems, which means that their apparent stability is nearly the same. This is supported by calculating and comparing the pMn values at pH 7.4 with 10 μM ligand and 10 μM Mn²⁺ concentrations (8.36 for [Mn(PhDTA)]²⁻ and 8.68 [Mn(*trans*-CDTA)]²⁻).¹⁵ In the case of Mg²⁺, Ca²⁺, Mn²⁺, Zn²⁺ and Cu²⁺ ions the formation of ML, mono- (MHL) and diprotonated (MH₂L) complex species was established. Potentiometric studies have also been carried out with 2:1 metal-ligand ratios with Mn²⁺, Zn²⁺ and Cu²⁺ to investigate the formation of dinuclear complexes. Both Zn²⁺ and Cu²⁺ form this kind of complexes with PhDTA, including protonated dinuclear species, though they possess low stability. The stability constant of the [Cu(PhDTA)]²⁻ complex could not be determined by using solely the pH-potentiometric data, as the Cu²⁺ ion is mostly in complexed form (~85%) even near pH = 1 (Figure 2). Therefore, the stability of this complex was assessed by using a combination of the UV-vis spectrophotometric method (applicable to very acidic samples) and pH-potentiometry titration data obtained at higher pH. The absorption spectra of the [Cu(PhDTA)]²⁻ complex as a function of H⁺ ion concentration are shown in Figure 3. In line with the equilibrium data obtained for the other metal ions studied, the stability constant of the [Cu(PhDTA)]²⁻ complex is

smaller than the value of the Cu²⁺ complexes of EDTA derivatives determined under identical conditions. On the other hand the stability constant determined in our study is in excellent agreement with the value determined previously by M. Tanaka and co-workers,³² indicating that the stability constants published recently for M²⁺ complexes by T. Zhang and co-workers³³ are not really reliable.

The pH dependence of the relaxivity of [Mn(PhDTA)]²⁻ was studied to acquire information about the relaxivity of the complex (r_{1p}) on one hand, and to support the equilibrium model used in the analysis of the pH-potentiometric titrations on the other hand. As shown in Figure 4, the formation of a monoprotated complex does not cause a considerable change in the relaxivity (only 0.3 – 0.5 mM⁻¹s⁻¹ units). The dissociation of the complex starts at pH < 3.5 leading to significant increase in the relaxivity because of the formation of [Mn(H₂O)₆]²⁺ (characterized by a r_{1p} value of 7.92 mM⁻¹s⁻¹ at 20 MHz and 25 °C). The measured relaxivity values remain constant in the pH range of 5-10, and then slightly decrease above pH 10. This can be interpreted by the oxidation of manganese upon the dissociation of a mixed hydroxo complex. The mixed hydroxo complex has not been studied because atmospheric oxygen oxidizes Mn²⁺ under the applied conditions. The relaxivity of the [Mn(PhDTA)]²⁻ complex is r_{1p} = 3.72 mM⁻¹s⁻¹ (between pH 5-10), which is very similar to that

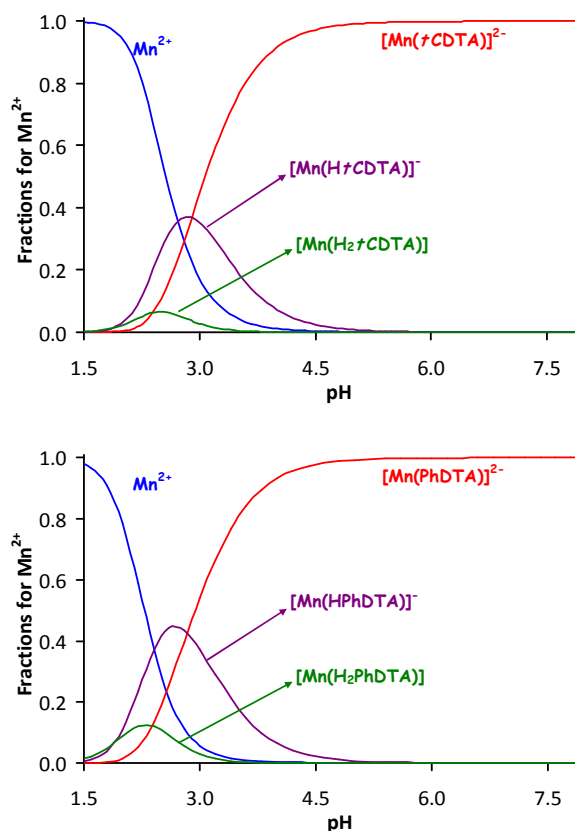


Figure 1. Concentration distribution curves calculated for [Mn(PhDTA)]²⁻ and [Mn(*trans*-CDTA)]²⁻ complexes ($c_{Mn^{2+}} = c_L = 0.001$ M).

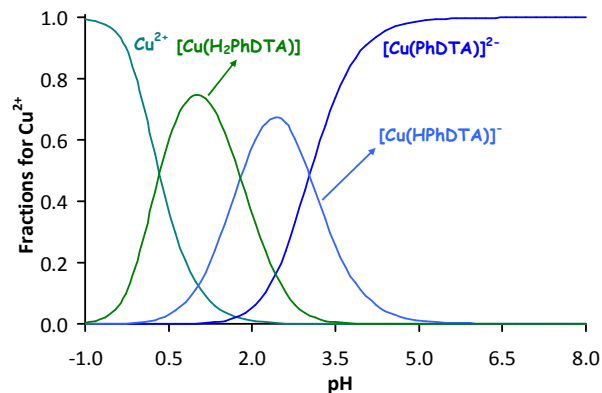


Figure 2. The species distribution curve calculated for the $[\text{Cu}(\text{PhDTA})]^{2-}$ complex by using the stability constants determined via combined UV-vis and pH-potentiometric titration data ($c_{\text{Cu}^{2+}} = c_{\text{L}} = 0.001 \text{ M}$).

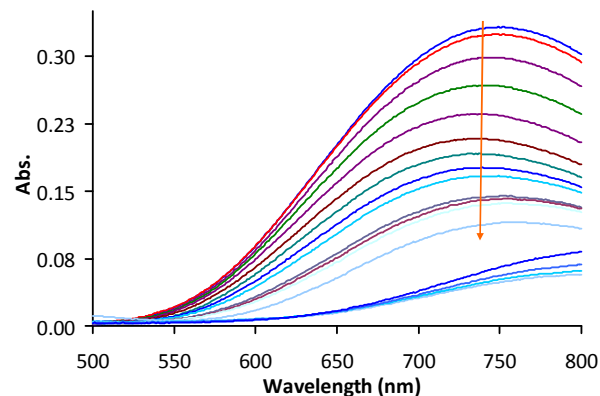


Figure 3. Absorption spectra of the $[\text{Cu}(\text{PhDTA})]^{2-}$ complex ($[\text{Cu}^{2+}]_{\text{tot}} = [\text{PhDTA}]_{\text{tot}} = 4 \times 10^{-3} \text{ M}$) recorded at different pH: 4.96, 4.06, 3.43, 2.99, 2.59, 2.30, 2.00, 1.71, 1.51, 1.30, 1.15, 1.0 and 0.7. The lower spectra correspond to the Cu^{2+} ion in solution at the same concentration and pH=1.0, 0.7, 0.3, and 0 ($\text{pH} = -\log[\text{H}^+]$).

Table 1. Protonation and stability constants of the EDTA, *cis*- and *trans*-CDTA and PhDTA ligands and their complexes formed with Ca^{2+} , Mg^{2+} , Mn^{2+} , Cu^{2+} and Zn^{2+} ions ($I = 0.15 \text{ M NaCl}$, $T = 25 \text{ }^\circ\text{C}$).

		EDTA ^a	<i>trans</i> -CDTA ^a	<i>cis</i> -CDTA ^c	PhDTAD ^d	PhDTA
H^+	$\log K_1^{\text{H}}$	9.17 ^b	9.36 ^b	11.00	6.41; 5.95 ^e	6.66(5)
	$\log K_2^{\text{H}}$	5.99 ^b	5.95 ^b	5.20	4.61; 3.70 ^e	4.85(6)
	$\log K_3^{\text{H}}$	2.73 ^b	3.62 ^b	3.41	3.53; 2.79 ^e	3.53(7)
	$\log K_4^{\text{H}}$	2.01 ^b	2.57 ^b	2.30	3.00; – ^e	3.32(6)
	$\log K_5^{\text{H}}$	1.38 ^b	1.49 ^b	1.41	–	–
	$\sum \log K_i^{\text{H}}$	21.28 ^b	22.99 ^b	23.32	17.55; 12.44 ^e	18.36
Mg^{2+}	$\log K_{\text{ML}}$	7.61	9.14	9.01	6.48 ^f ; 10.41 ^e	6.94(3)
	$\log K_{\text{MHL}}$	–	3.53	4.72	2.7 ^f	3.78(6)
Ca^{2+}	$\log K_{\text{ML}}$	9.53	10.23	9.65	8.27 ^f ; 11.25 ^e	8.79(2)
	$\log K_{\text{MHL}}$	2.92	3.53	4.55	3.0 ^f	3.38(3)
Mn^{2+}	$\log K_{\text{MnL}}$	12.46 ^b	14.32 ^b	14.19	11.37	11.79(2)
	$\log K_{\text{MnHL}}$	2.95 ^b	2.90 ^b	2.85	2.29	2.84(2)
	$\log K_{\text{MnH2L}}$	– ^b	1.89 ^b	–	1.70	1.91(5)
	pMn ^g	7.83	8.68	7.82	8.16	8.38
Cu^{2+}	$\log K_{\text{ML}}$	19.02 ^h	19.78 ^h	18.3(1) ^h	15.21 ^h ; 24.84 ^f	15.21(6) ^h
	$\log K_{\text{MHL}}$	3.15	2.91	3.65	3.04	3.35(1)
	$\log K_{\text{MH2L}}$	2.04	1.10	–	1.80	2.18(1)
	$\log K_{\text{M2L}}$	–	–	–	–	2.75(2)
Zn^{2+}	$\log K_{\text{ML}}$	15.92	16.75	17.06	12.89; 17.99 ^f	13.03(3)
	$\log K_{\text{MHL}}$	3.23	2.57	2.76	2.96	3.56(2)
	$\log K_{\text{MH2L}}$	1.50	–	–	1.30	1.90(3)
	$\log K_{\text{M2L}}$	–	–	–	–	2.69(7)

^a 0.15 M NaCl, 25°C from the diploma work of Veronika Józsa, 2015, University of Debrecen, Debrecen, Hungary; ^b Ref. 23.; ^c Ref. 27; ^d Ref. 32; ^e Ref. 33; ^f Ref. 31; ^g Calculated by using $c_{\text{Mn}} = c_{\text{Lig}} = 0.01 \text{ mM}$ at pH=7.4 according to Ref. 15.; ^h Determined by simultaneous fitting of the UV-vis and pH-potentiometric data.

Table 2. Rate and stability constants characterizing the dissociation of the $[\text{Mn}(\text{EDTA})]^{2-}$, $[\text{Mn}(\text{cis-CDTA})]^{2-}$, $[\text{Mn}(\text{trans-CDTA})]^{2-}$ and $[\text{Mn}(\text{PhDTA})]^{2-}$ complexes ($I = 0.15 \text{ M NaCl}$, $T = 25 \text{ }^\circ\text{C}$).

	EDTA ^{a, b}	<i>cis</i> -CDTA ^c	<i>trans</i> -CDTA ^{a, b}	PhDTA
$k_1 (\text{M}^{-1}\text{s}^{-1})$	5.2×10^4	1.02×10^5	4.0×10^2	$(2.01 \pm 0.04) \times 10^2$
$k_2 (\text{M}^{-2}\text{s}^{-1})$	2.3×10^8	–	–	$(4.9 \pm 0.3) \times 10^3$
$k_3 (\text{M}^{-1}\text{s}^{-1})$	45	–	–	0.21 ± 0.01
K_{MnLCu}	–	79	79	27 ± 4
$\log K_{\text{H}}$	2.65 ^d	2.85 ^d	2.90 ^d	2.84 ^d
$t_{1/2} (\text{h})^e$	0.076	0.47	12.3	19.1

^a Ref. 23. ^b For for $[\text{Mn}(\text{edta})]^{2-}$ $k_3 = 3.0 \times 10^{-1} \text{ M}^{-1}\text{s}^{-1}$, $k_4 = \sim 4.8 \times 10^1 \text{ M}^{-2}\text{s}^{-1}$ and $\log K_{\text{MnHL}} = 3.10$ while $[\text{Mn}(\text{trans-cdta})]^{2-}$ $k_1 = 3.2 \times 10^2 \text{ M}^{-1}\text{s}^{-1}$ and $t_{1/2} = 15 \text{ h}$ were found in Ref. 35. ^c Ref. 27. ^d Determined by pH-potentiometry and fixed in the data fitting. ^e Calculated at pH = 7.4 in the presence of $c_{\text{Cu}^{2+}} = 1 \times 10^{-5} \text{ M}$.

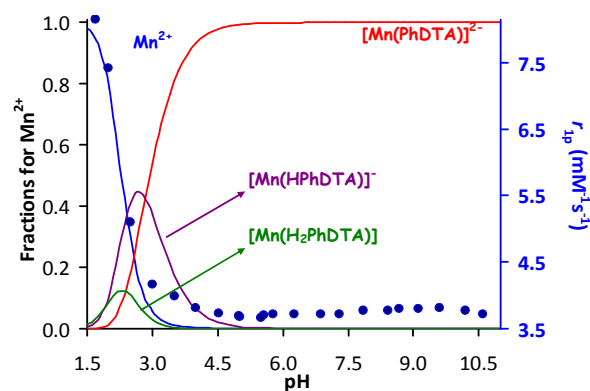
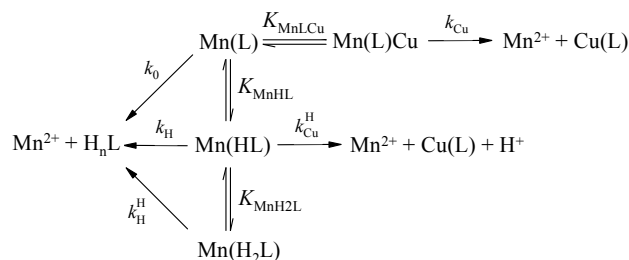


Figure 4. The speciation distribution of $[\text{Mn}(\text{PhDTA})]^{2-}$ complex with the pH dependence of its relaxivity ($I = 0.15 \text{ M NaCl}$, $T = 25 \text{ }^\circ\text{C}$, 20 MHz).

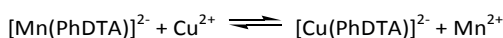


Scheme 2. Reaction mechanisms for the dissociation of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex.

determined previously for the $[\text{Mn}(\text{trans-CDTA})]^{2-}$ complex.^{23,27} Thus, we conclude that the replacement of the cyclohexanediamine unit by ortho-phenylenediamine bridge does not affect significantly the relaxivity of the Mn^{2+} complexes.

Dissociation kinetics of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex

Since Mn^{2+} is a biogenic metal ion, its toxicity is considerably lower than that of Gd^{3+} . Nevertheless Mn^{2+} is also harmful in large doses and causes symptoms similar to the ones associated with Alzheimer's disease.³⁴ Thus, our goal is to prepare Mn^{2+} -based contrast agents that do not dissociate *in vivo* to avoid potential toxicity issues. With an aim of gaining information about its inertness the rates of the exchange reactions of $[\text{Mn}(\text{PhDTA})]^{2-}$ have been studied in the presence of Cu^{2+} . The kinetic experiments have been conducted in the pH range of 3.6 – 5.2 with the use of Cu^{2+} as exchanging metal ion. These transmetalation reactions are slow enough in the mentioned pH range to allow for measurements with conventional spectrophotometric methods. The studied exchange reactions can be delineated as follows:



For the metal exchange reactions, the concentration of Cu^{2+} was 10 to 40 times higher than the concentration of the

complex to assure pseudo-first-order conditions. Fitting the acquired time-absorbance curves to the equation for a first order reaction gave the pseudo-first-order rate constant (k_{obs}) for each reaction. The H^+ concentration dependence of the dissociation rates are shown in Figure 5. It is obvious that the reaction rates show a second order dependence on proton concentration. Furthermore, the experiments performed using different concentrations of Cu^{2+} indicate that the role of the metal ion is not negligible, i.e. the dissociation can be realized through a heterodinuclear complex. The acquired rate constants increase with the rise of H^+ concentration and change by varying Cu^{2+} concentration. Accordingly, the reaction can occur both with the attack of protons and the exchanging metal ion. However the effect of the exchanging metal ion on the reaction rates depend strongly on pH. While the increase of the Cu^{2+} concentration leads to lower reaction rates in the more acidic samples, at higher pH it leads to higher reaction rates (Figure 6). The explanation for this behavior might be that the two reaction pathways (the proton assisted and the metal ion assisted paths) are in competition with each other. For the dissociation of the complex in the presence of Cu^{2+} the general reaction scheme shown in Scheme 2 can be established. Taking into account the different reaction pathways (characterized by the rate coefficients k_0 , k_{H} , k_{H}^{H} , k_{Cu} , k_{Cu}^{H}), the protonation constants and stability constants of the complexes (K_{MnHL} , $K_{\text{MnH}_2\text{L}}$ and K_{MnLCu}), the pseudo-first-order rate constant (k_{obs}) can be expressed by eqn. 1, where

$$\begin{aligned}
 K_{\text{MnHL}} &= [\text{Mn}(\text{HL})]/([\text{Mn}(\text{L})][\text{H}^+]), \\
 K_{\text{MnH}_2\text{L}} &= [\text{Mn}(\text{H}_2\text{L})]/([\text{Mn}(\text{HL})][\text{H}^+]), \\
 K_{\text{MnLCu}} &= [\text{Mn}(\text{L})\text{Cu}]/([\text{Mn}(\text{L})][\text{Cu}^{2+}]), \\
 k_2 &= k_{\text{H}}^{\text{H}} \cdot K_{\text{MnHL}} \cdot K_{\text{MnH}_2\text{L}}, \quad k_3 = k_{\text{Cu}} \cdot K_{\text{MnLCu}} \quad \text{and} \quad k_4 = k_{\text{Cu}}^{\text{H}} \cdot K_{\text{MnHL}}.
 \end{aligned}$$

$$k_{\text{obs}} = \frac{k_0 + k_1[\text{H}^+] + k_2[\text{H}^+]^2 + k_3[\text{Cu}^{2+}] + k_4[\text{Cu}^{2+}][\text{H}^+]}{1 + K_{\text{MnHL}}[\text{H}^+] + K_{\text{MnHL}}K_{\text{MnH}_2\text{L}}[\text{H}^+]^2 + K_{\text{MnLCu}}[\text{Cu}^{2+}]} \quad (1)$$

Initial attempts to fit the experimental data to eqn. 1 evidenced that beside two proton-assisted pathways (characterized by k_1 and k_2) the attack of the metal ion with the formation of a dinuclear complex also affects the dissociation reaction. The k_4 rate constant gave a negative value during the fitting thus the path was neglected. Taking into consideration the pathways mentioned above, the pseudo-first order rate constants of the exchange reaction were analyzed using the following simplified expression (eqn. 2):

$$k_{\text{obs}} = \frac{k_1[\text{H}^+] + k_2[\text{H}^+]^2 + k_3[\text{Cu}^{2+}]}{1 + K_{\text{MnHL}}[\text{H}^+] + K_{\text{MnLCu}}[\text{Cu}^{2+}]} \quad (2)$$

The rate and stability constants obtained by fitting the k_{obs} values are compared with the data acquired for the $[\text{Mn}(\text{EDTA})]^{2-}$, $[\text{Mn}(\text{cis-CDTA})]^{2-}$ and $[\text{Mn}(\text{trans-CDTA})]^{2-}$ complexes in Table 2. The results show that, in the case of the

trans-CDTA complex, the partial decoordination of the ligand from Mn^{2+} and coordination to Cu^{2+} is a slow process, which results in the formation of a quite stable dinuclear intermediate (a so-called “dead-end” complex). The formation of the dinuclear species slows down the reaction by decreasing the contribution of the proton assisted pathway. On the contrary, the dinuclear complex formed with PhDTA possesses a “Janus-face” nature in affecting the rate of the reaction depending on H^+ concentration. This can be explained by a positive contribution of the metal-assisted pathway to the overall exchange reaction above pH 4.6 (Fig 6), while at pH below pH 4.3 the slower metal-assisted dissociation competes with the proton assisted pathway, thus lowering the pseudo-first rate constants. It is obvious from the data shown in Table 2 that the rate of the proton assisted dissociation in the case of $[\text{Mn}(\text{PhDTA})]^{2-}$ is half of that of $[\text{Mn}(\text{trans-CDTA})]^{2-}$, which is reassuring with respect to a potential application of the complex. The half-life of dissociation of the Mn^{2+} complexes have been calculated using the first order rate constants at pH 7.4 and 25 °C ($c_{\text{Cu}^{2+}}=0.01$ mM). The results indicate that the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex presents the highest inertness with respect to complex dissociation among the four complexes ($t_{1/2, [\text{Mn}(\text{PhDTA})]^{2-}} = 19.1$ h considering seven times the depletion and not considering the reformation of the complex, and $t_{1/2, [\text{Mn}(\text{trans-CDTA})]^{2-}} = 12$ h) while the dissociation of the $[\text{Mn}(\text{EDTA})]^{2-}$ and $[\text{Mn}(\text{cis-CDTA})]^{2-}$ complexes is complete within few minutes under these conditions). Considering the typical excretion time of low molecular weight contrast agents ($t_{1/2} = 1.6$ h), the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex appears to be a good candidate for *in vivo* experiments.

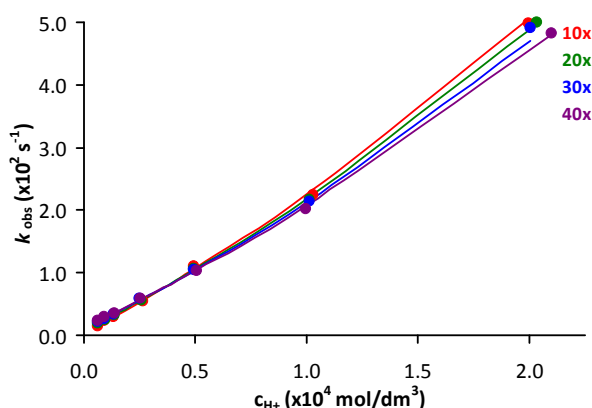


Figure 5. Dissociation rates (k_{obs}) of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex plotted as a function of H^+ ion concentration ($I = 0.15$ M NaCl, $T = 25$ °C). The colors represent different Cu^{2+} ion concentration as follows: 2 mM (red), 4 mM (green), 6 mM (blue) and 8 mM (purple).

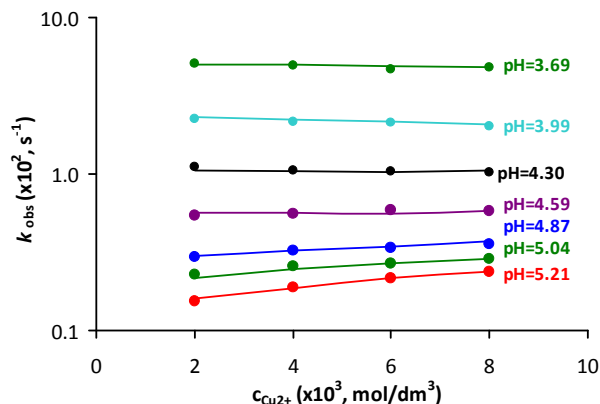


Figure 6. Dissociation rates (k_{obs}) of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex plotted as a function of Cu^{2+} ion concentration ($I = 0.15$ M NaCl, $T = 25$ °C).

DFT calculations

DFT calculations were carried out to gain insight into the structure of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex in solution. Geometry optimizations were performed on the $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ system, as the explicit inclusion of two second-sphere water molecules involved in hydrogen-bonding interactions with the coordinated water molecule are required for an accurate calculations of Mn-O_{water} distances and scalar hyperfine coupling constants at the ^{17}O nucleus of the coordinated water molecule.³⁶ The optimized geometry (Fig 7) calculated at the M062X/TZVP level shows a very good agreement with the X-ray crystallographic data reported by Tanaka et al.³⁷ The calculated bond distances deviate from the experimental ones by < 0.036 Å (Table S1, ESI[†]). A comparison of the structure obtained for $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ and that calculated for $[\text{Mn}(\text{EDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ using the same methodology reveals that the average Mn-N distance is somewhat shorter in the EDTA complex (2.411 Å) than in the PhDTA one (2.440 Å). An opposite situation is observed for the Mn-O_{carboxylate} distances, which show average values of 2.243 and 2.215 Å for $[\text{Mn}(\text{EDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ and $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$, respectively. The shorter Mn-O_{carboxylate} distances observed for $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2-}$ are likely responsible for the higher kinetic inertness of this complex with respect to the EDTA derivative, as the proton-assisted dissociation pathway likely involves decoordination of a protonated acetate group, while the metal-assisted mechanism also requires decoordination of acetate groups to form a dinuclear intermediate.

The Mn-O_{water} distance calculated for $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ (2.273 Å) is similar to that observed in the solid state (2.242 Å)³⁷ and only slightly longer than the value calculated for $[\text{Mn}(\text{EDTA})(\text{H}_2\text{O})]^{2-} \cdot 2\text{H}_2\text{O}$ (2.251 Å). The ^{17}O hyperfine coupling constant obtained using DFT ($A_0/\hbar = -42.6 \times 10^6$ rad s^{-1}) is in excellent agreement with the experimental value reported by Hunt ($A_0/\hbar = -38.0 \times 10^6$ rad s^{-1}),²⁸ which indicates that our calculations provide an adequate description of the spin density distribution on the Mn-O_{water} bond.

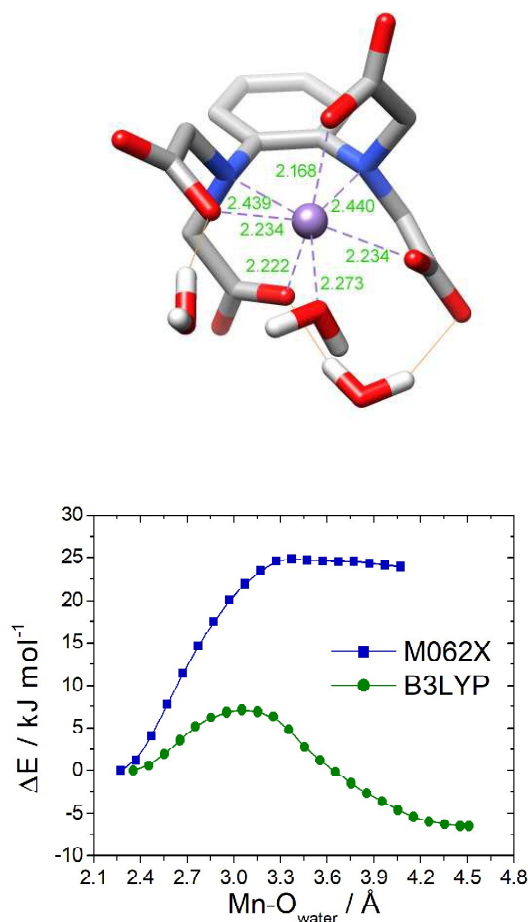


Figure 7. Top: Structure of the $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2+}\cdot 2\text{H}_2\text{O}$ system obtained using DFT calculations at the M062X/TZVP level. Bottom: Relaxed potential energy surface scans calculated for $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2+}\cdot 2\text{H}_2\text{O}$ using different functionals.

The experimental determination of the number of water molecules coordinated to Mn^{2+} in potential MRI contrast agents remains a difficult task. Caravan et al.³⁸ proposed a method relying on the analysis of the temperature dependence of the transverse ^{17}O NMR relaxation rates. This method can only be applied to systems undergoing a changeover from a slow exchange regime at low temperature to a fast exchange regime at high temperature. As a consequence, the hydration number of complexes whose transverse relaxation rates are in the fast exchange regime in the whole accessible temperature range cannot be estimated using this methodology. Thus, we wanted to explore whether our DFT calculations could provide information on the number of coordinated water molecules. For this purpose, we explored the potential energy surface (PES) of the $[\text{Mn}(\text{EDTA})(\text{H}_2\text{O})]^{2-}\cdot 2\text{H}_2\text{O}$ and $[\text{Mn}(\text{PhDTA})(\text{H}_2\text{O})]^{2+}\cdot 2\text{H}_2\text{O}$ systems by increasing the $\text{Mn}-\text{O}_{\text{water}}$ distance (Fig 7, see also Fig S2, ESI[†]). These relaxed PES scans provided the six-coordinate $[\text{Mn}(\text{EDTA})]^{2-}\cdot 3\text{H}_2\text{O}$ and $[\text{Mn}(\text{PhDTA})]^{2+}\cdot 3\text{H}_2\text{O}$ structures, which turned out to be less stable than the seven-coordinate forms at the M062X/TZVP level, in line with the experimental evidence. However, the PES

was found to differ considerably depending on the particular functional used. For instance the B3LYP functional stabilizes considerably the six-coordinate structures, which in the case of the PhDTA complex is the most stable form at the B3LYP/TZVP level. This is in line with our recent results that showed that M062X/TZVP calculations provided energy profiles for the water exchange reactions of $[\text{Li}(\text{H}_2\text{O})_4]^+$ and $[\text{Be}(\text{H}_2\text{O})_4]^{2+}$ in good agreement with the experiment, while B3LYP/TZVP calculations overstabilized a lower coordination numbers.³⁹ While an exhaustive evaluation of different functionals is beyond the scope of the present study, the results obtained here suggest that M062X/TZVP calculations may be useful in predicting hydration numbers of Mn^{2+} complexes relevant as MRI contrast agents.

Electrochemical measurements

The redox stability of the $[\text{Mn}(\text{PhDTA})]^{2+}$ complex was investigated using cyclic voltammetry experiments in aqueous 0.15 M NaCl. The cyclic voltammograms obtained using different scan rates are characteristic of an irreversible system. The cyclic voltammogram recorded at a scan rate of 50 mV/s showed an oxidation peak at $E_{\text{ox}} = +813$ mV and a reduction peak at $E_{\text{red}} = +552$ mV ($\Delta E_{1/2} = 683$ mV vs Ag/AgCl, Fig 8). The cyclic voltammogram obtained for $[\text{Mn}(\text{EDTA})]^{2-}$ using the same conditions shows both the anodic ($E_{\text{ox}} = +769$ mV) and cathodic ($E_{\text{ox}} = +510$ mV) peaks shifted to lower potentials, which shows that PhDTA presents a lower ability to stabilize Mn^{3+} compared to EDTA. This might be related to the harder nature of the amine nitrogen atoms of EDTA compared to the aniline donor atoms of PhDTA (according to Pearson's HSAB principle). This may be also related to the differences in the geometries as evidenced for $[\text{Mn}(\text{PhDTA})]^{2+}$ by X-ray crystallography when compared with that of the $[\text{Mn}(\text{trans-CDTA})]^{2-}$ complex,^[37, 40] as any axial distortion of the Mn(II) coordination environment should favour Mn(III) over Mn(II), thereby leading to a decrease in potential.^[41, 42] The half-cell potential of $[\text{Mn}(\text{EDTA})]^{2-}$ ($\Delta E_{1/2} = 640$ mV vs Ag/AgCl, Figure S3, ESI[†]) is very similar to that reported by Caravan using 0.5 M KNO_3 as supporting electrolyte (633 mV).⁸

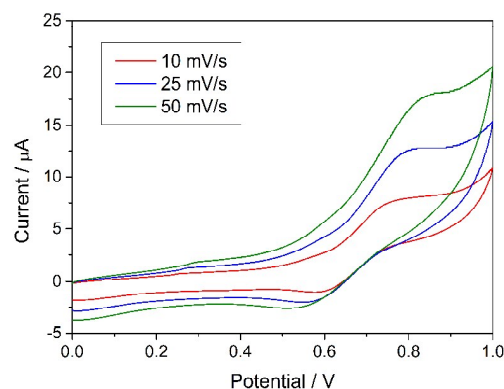


Figure 8. Cyclic voltammograms recorded using different scan rates for 3 mM $[\text{Mn}(\text{PhDTA})]^{2+}$ (pH = 7.1, 0.15 NaCl, potentials reported vs Ag/AgCl).

Conclusions

The replacement of the ethylenediamine bridging unit in EDTA by the *ortho*-phenylenediamine backbone to give PhDTA decreases the basicity of the ligand owing to the electron withdrawing effect of the aromatic unit, yet the PhDTA complexes formed with Ca^{2+} , Mg^{2+} , Mn^{2+} , Cu^{2+} and Zn^{2+} possess very similar conditional stabilities to those of the corresponding EDTA complexes. The kinetic studies approached by studying the metal exchange reactions occurring between the Cu^{2+} ion and the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex revealed an improved inertness compared to $[\text{Mn}(\text{trans-CDTA})]^{2-}$. The complex possesses a half-life of 19.1 hours at physiological pH and 25 °C, thus only about 8% of the injected amount is expected to dissociate in the body. It has to be emphasized however that at 37 °C the extent of the dissociation will likely be slightly higher, and thus further ligand design is needed to find even better ligand candidates for Mn^{2+} ion complexation. DFT calculations confirmed that the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex possesses one metal bound water molecule (in agreement with the solid state structure of the complex published previously³⁷). As a result, we found a relaxivity ($r_1 = 3.72 \text{ mM}^{-1} \text{ s}^{-1}$ at 20 MHz and 25 °C) very similar to those determined earlier for the Mn^{2+} complexes formed with EDTA as well as *cis*- or *trans*-CDTA ligands (Table 3). The most important physicochemical parameters of the $[\text{Mn}(\text{EDTA})]^{2-}$, $[\text{Mn}(\text{cis-CDTA})]^{2-}$, $[\text{Mn}(\text{trans-CDTA})]^{2-}$ and $[\text{Mn}(\text{PhDTA})]^{2-}$ complexes are collected and compared in Table 3. As it can be seen from the data compared in Table 3 the *ortho*-phenylenediamine building block represents an attractive alternative to the *trans*-1,2-cyclohexanediamine unit when tailoring inert open-chain based Mn^{2+} complexes for safe MRI applications.

Table 3. Rate and stability constants characterizing the dissociation of the $[\text{Mn}(\text{EDTA})]^{2-}$, $[\text{Mn}(\text{cis-CDTA})]^{2-}$, $[\text{Mn}(\text{trans-CDTA})]^{2-}$ and $[\text{Mn}(\text{PhDTA})]^{2-}$ complexes ($I = 0.15 \text{ M NaCl}$, $T = 25 \text{ °C}$).

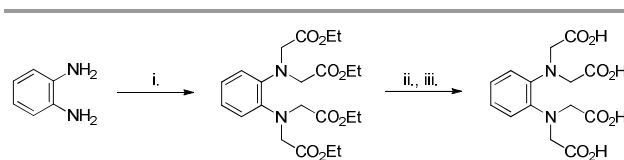
	EDTA ^a	<i>cis</i> -CDTA ^b	<i>trans</i> -CDTA ^a	PhDTA
$\log K_{[\text{Mn(L)]}$	12.46	14.19	14.32	11.79
pMn ^c	7.83	7.82	8.68	8.38
$k_{\text{ex}}^{298} (\times 10^7 \text{ s}^{-1})$	41 ^d	22.5	14.0 ^e	35.0 ^f
$r_{1p}^{298} (\text{mM}^{-1} \text{ s}^{-1})^g$	3.20	3.85	3.62	3.72
$t_{1/2} (\text{h})^h$	0.076	0.47	12.2	19.1

^a Ref. 23; ^b Ref. 27; ^c pMn values were calculated at pH=7.4 by using 0.01 mM Mn^{2+} and ligand concentration as suggested by É. Tóth and o-workers, Ref. 15; ^d From Ref. 43; ^e From Ref. 44; ^f From Ref. 45; ^g at 25 °C and 20 MHz; ^h The half-lives (h) of dissociation were extrapolated to pH=7.4 by using 0.01 mM Cu^{2+} ion concentration

Experimental Section

Synthesis

The ligand PhDTA was synthesized as shown in Scheme 3 using a slight modification of the previously published procedure.³³



Scheme 3. Synthesis of PhDTA. Reagents and conditions: i) $\text{BrCH}_2\text{COOEt}$, DIPEA, NaI, MeCN; ii) NaOH, EtOH; iii) HCl/H₂O and crystallization.

***ortho*-Phenylenediamine-*N,N,N',N'*-tetraacetate tetraethyl ester (1).** 1,2-Diaminobenzene (1.10 g, 10.2 mmol), ethyl-bromoacetate (6.70 mL, 10.1 g, 60.5 mmol), sodium iodide (1.30 g, 8.67 mmol) and *N,N*-diisopropylethylamine (8.30 mL, 6.16 g, 47.7 mmol) were mixed in acetonitrile (10 mL) and the mixture was refluxed under N_2 atmosphere for 7 hours. After cooling to room temperature, the reaction mixture was evaporated under reduced pressure. Water (50 mL) was added to the residue and the mixture extracted with chloroform (3×40 mL). The combined organic extracts were dried with Na_2SO_4 and evaporated under reduced pressure resulting in a brown oil, which was purified with flash silica gel chromatography (petroleum ether:ethyl-acetate 10:1). Yield 2.90 g (62%).

***ortho*-Phenylenediamine-*N,N,N',N'*-tetraacetic acid (PhDTA).** The ethyl ester precursor **1** (2.90 g, 6.41 mmol) was dissolved in ethanol (40 mL) and NaOH (1.28 g, 32 mmol) dissolved in distilled water (5 mL) was added dropwise to the stirred reaction mixture. A white precipitate was formed and the mixture was refluxed for 18 hours. The white crystalline product was filtered with G3 filter crucible and dissolved in water (2-3 mL). The pH was adjusted to pH=2 with concentrated HCl and the precipitated white solid was isolated by filtration and washed with cold distilled water (3×5 mL). Finally the product was dried to constant mass. Yield 1.78 g (52%). ¹H NMR [360 MHz, D₂O] δ 4.05 (8H, s, NCH₂), 7.2-7.4 (4H, m, CH aromatic); ¹³C NMR [100 MHz, D₂O] δ 57.7 (4C NCH₂), 123.2 (2C CH ring), 127.6 (2C CH ring), 141.0 (2C C ring), 174.7 (4C CO); MS (ESI) m/z 481,300 (M+H)⁺ 35%; 503,282 (M+Na)⁺ 100%; 519,250 (M+K)⁺ 13%.

Equilibrium studies

The MnCl_2 , CaCl_2 , MgCl_2 , ZnCl_2 and CuCl_2 stock solutions were prepared from the highest analytical grade chemicals, and their concentrations were determined by complexometric titration with standardized $\text{Na}_2\text{H}_2\text{EDTA}$ and eriochrome black T indicator in the presence of ascorbic acid and potassium hydrogen tartrate for MnCl_2 , murexide indicator for CaCl_2 and CuCl_2 , eriochrome black T for MgCl_2 and xilenolorange in the presence of hexamethylenetetramine for the ZnCl_2 solution. The concentration of the ligand stock solution was determined by pH-potentiometric titrations. For determining the protonation constants of the ligand, pH potentiometric titrations were carried out with 0.2 M NaOH, using 0.002 M ligand solutions. The ionic strength was set to 0.15 M by using NaCl. The titrated samples (starting volume of 6 mL) were stirred mechanically and thermostated at 25 °C by a circulating water bath (± 0.1 °C). The protonation constants of the ligand ($\log K_i^{\text{H}}$) are defined as follows:

$$K_i^H = \frac{[H_iL]}{[H_{i-1}L][H^+]} \quad (3)$$

where $i = 1, 2, \dots, 5$ and $[H_iL]$ and $[H^+]$ are the equilibrium concentrations of the ligand ($i = 1$), protonated forms of the ligand ($i = 2, \dots, 5$), and hydrogen ions respectively. To avoid the effect of CO_2 , N_2 gas was bubbled through the solutions during the titrations process. The pH-potentiometric titrations were performed with a *Metrohm 785 DMP Titrino* titration workstation with the use of a *Metrohm 6.0234.100* combined electrode in the pH range of 1.8–12.0. For the calibration of the pH meter, KH-phthalate (pH = 4.005) and borax (pH = 9.177) buffers were used, and the H^+ concentrations were calculated from the measured pH values by applying the method proposed by Irving et al.⁴⁶ A solution of approximately 0.01 M HCl was titrated with a 0.2 M NaOH solution (0.15 M NaCl), and the differences between the measured and calculated pH values (for the samples with pH < 2.4) were used to calculate the $[H^+]$ from the pH values measured in the titration experiments. The measured points with pH > 11.0 of the acid-base titration were used to calculate the ionic product of water which was found to be 13.867 under our experimental conditions. For the calculation of the equilibrium constants, the PSEQUAD program was used.⁴⁷ The protonation constants of the PhDTA ligand were determined by titrating ligand solutions (acidified with a known volume of a standard HCl solution) with 0.2 M NaOH at 0.15 M ionic strength in the 1.8–12.0 pH range. The $\log K_i^H$ values were calculated from 260 V(mL)-pH data pairs. The stability constants of the metal complexes were determined using the direct pH-potentiometric method by titrating samples with 1:1 and 2:1 metal-to-ligand ratios (the number of data pairs were between 150–200), allowing 1 min for the sample equilibration to occur. In the presence of metal excess the titration was continued until the appearance of the precipitate (pH = 7–9). The stability constant of the Cu^{2+} complexes was too high to be determined by pH-potentiometry, hence a direct UV-vis spectrophotometric method was used. The spectrophotometric measurements were performed with a *Cary 100 Bio* spectrophotometer at 25 °C, using semimicro 1.0 cm cells. For the determination of the stability constant the absorbance was measured at 12 different acid concentrations (0.01–3.00 M) at 7 wavelengths between 650 and 800 nm (the concentration of the complex was 4.0 mM). A pH-titration of $[Cu(PhDTA)]^{2-}$ was also performed in the pH range 1.50–12.0 with a starting volume of 6 mL, $C_{Cu} = C_{PhDTA} = 2.0$ mM.

Kinetic Studies

The dissociation rates of the $[Mn(PhDTA)]^{2-}$ complex were investigated at 25 °C and 0.15 M NaCl ionic strength by spectrophotometry, in the presence of a high (10–40-fold) excess of exchanging Cu^{2+} ion to ensure pseudo-first-order conditions, using a *Cary 100 Bio* spectrophotometer at 300 nm (using semimicro 1.0 cm cells). The concentration of the complex was set to 2×10^{-4} M, and the kinetic studies were performed by using a noncoordinating buffer, N'-methyl piperazine (NMP, $\log K_2^H = 4.92(0.02)$) at 0.04 M concentration to maintain constant pH in the samples. Before the kinetic runs

only the solution of the complex was buffered, while in the solution of exchanging Cu^{2+} the ionic strength was set. Under these conditions, the rate of the reaction can be expressed as follows: $-d[Mn(PhDTA)]^{2-}_{tot}/dt = k_{obs}[Mn(PhDTA)]^{2-}_{tot}$, where k_{obs} is the pseudo-first-order rate constant and $[Mn(PhDTA)]^{2-}_{tot}$ is the total concentration of the Mn^{2+} complex. The k_{obs} values were determined using the following equation: $A_t = (A_r - A_p)e^{(-k_{obs}t)} + A_v$, where A_t is the absorbance at time t , A_r is the absorbance of the reactants, A_p is the absorbance of the products.

Determination of the relaxivity of the Mn^{2+} Complex

The relaxation times of water protons were measured at 20 MHz with a Bruker Minispec MQ-20 NMR Analyzer. The temperature of the sample holder was set to 25.0 ± 0.2 °C and controlled with the use of circulating water bath. The longitudinal relaxation times (T_1) were measured by using the inversion recovery method ($180^\circ - \tau - 90^\circ$) by averaging 5–6 data points obtained at 8 different τ values. During the studies, the concentration of the $[Mn(PhDTA)]^{2-}$ complex was 1.0 mM ($I = 0.15$ M NaCl) and the pH was varied in the range of 1.6–10.6.

Cyclic voltammetry Experiments

Cyclic voltammograms were recorded using a 797 VA Computrace potentiostat/galvanostat from Metrohm (Herisau, Switzerland) connected to a typical three electrode cell. A glassy carbon rotating disk electrode (stirring rate of 2000 rpm) was used as working electrode, while the counter electrode was a platinum rod. Potentials were measured using a Ag/AgCl reference electrode filled with 3 mol·L⁻¹ KCl. Solutions were purged with high purity (99.999%) nitrogen during 30 seconds prior recording the voltammograms. The starting and end potentials were 0.0 V, while the first vertex potential was set to +1.0 V. Sweep rates of 10–100 mV s⁻¹ were used.

Computational Details

The $[Mn(PhDTA)(H_2O)]^{2-} \cdot 2H_2O$ and $[Mn(EDTA)(H_2O)]^{2-} \cdot 2H_2O$ systems were fully optimized by using the hybrid GGA B3LYP⁴⁸ functional and the hybrid meta-GGA M062X⁴⁹ functional along with the TZVP⁴⁷ basis set. The nature of optimized geometries as true energy minima was confirmed by frequency analysis. Hyperfine coupling constants were calculated on the optimized structures by using the EPR-III basis set for the ligand atoms⁵¹ and the aug-cc-pVTZ-J basis set for Mn.⁵² Throughout this study bulk solvent effects were considered by using the integral equation formalism variant of the polarizable continuum model (IEFPCM).⁵³ All calculations were performed using the Gaussian 09 program package (revision D.01).⁵⁴

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TOC Graphic

Equilibrium, dissociation kinetics, relaxometric and electrochemical properties of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex were investigated and the structure of the $[\text{Mn}(\text{PhDTA})]^{2-}$ complex was studied by using DFT calculations.

INERTNESS OF Mn(II) COMPLEXES

