

Thermodynamic Stability of Mn(II) Complexes with Aminocarboxylate Ligands Analyzed Using Structural Descriptors

Rocío Uzal-Varela, Francisco Pérez-Fernández, Laura Valencia, Aurora Rodríguez-Rodríguez,* Carlos Platas-Iglesias,* Peter Caravan, and David Esteban-Gómez



Cite This: *Inorg. Chem.* 2022, 61, 14173–14186



Read Online

ACCESS |



Metrics & More

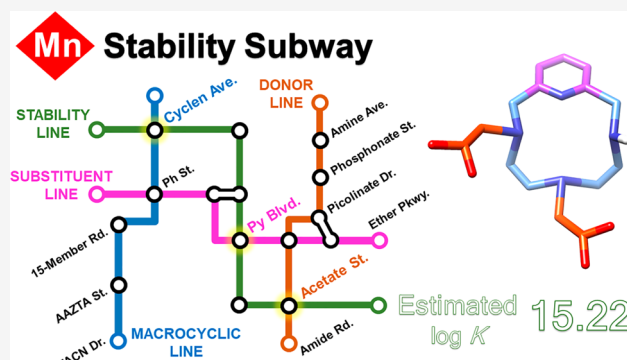


Article Recommendations



Supporting Information

ABSTRACT: We present a quantitative analysis of the thermodynamic stabilities of Mn(II) complexes, defined by the equilibrium constants ($\log K_{\text{MnL}}$ values) and the values of pMn obtained as $-\log[\text{Mn}]_{\text{free}}$ for total metal and ligand concentrations of 1 and 10 μM , respectively. We used structural descriptors to analyze the contributions to complex stability of different structural motifs in a quantitative way. The experimental $\log K_{\text{MnL}}$ and pMn values can be predicted to a good accuracy by adding the contributions of the different motifs present in the ligand structure. This allowed for the identification of features that provide larger contributions to complex stability, which will be very helpful for the design of efficient chelators for Mn(II) complexation. This issue is particularly important to develop Mn(II) complexes for medical applications, for instance, as magnetic resonance imaging (MRI) contrast agents. The analysis performed here also indicates that coordination number eight is more common for Mn(II) than is generally assumed, with the highest $\log K_{\text{MnL}}$ values generally observed for hepta- and octadentate ligands. The X-ray crystal structure of $[\text{Mn}_2(\text{DOTA})(\text{H}_2\text{O})_2]$, in which eight-coordinate $[\text{Mn}(\text{DOTA})]^{2-}$ units are bridged by six-coordinate exocyclic Mn(II) ions, is also reported.



The analysis performed here also indicates that coordination number eight is more common for Mn(II) than is generally assumed, with the highest $\log K_{\text{MnL}}$ values generally observed for hepta- and octadentate ligands. The X-ray crystal structure of $[\text{Mn}_2(\text{DOTA})(\text{H}_2\text{O})_2]$, in which eight-coordinate $[\text{Mn}(\text{DOTA})]^{2-}$ units are bridged by six-coordinate exocyclic Mn(II) ions, is also reported.

INTRODUCTION

Magnetic resonance imaging (MRI) often uses contrast-enhanced procedures to attain a more accurate diagnosis of different malignancies.^{1–4} The contrast agents (CAs) that are currently used in clinics are complexes with the paramagnetic metal ion Gd(III),^{5,6} which are very efficient relaxation agents of water proton nuclei in their vicinity. As a result, CAs shorten significantly the water longitudinal relaxation times, providing an enhanced signal of the tissues in which they are distributed, as fast T_1 relaxation allows for the accumulation of more signal intensity using short repetition times.⁷ The efficacy of Gd(III) as a T_1 relaxation agent is related to the dipolar interaction between the nuclear and electron spins, which is particularly efficient due to the presence of seven unpaired electrons and the long electronic relaxation time.^{8,9} While the use of Gd(III) CAs is regarded to be safe, a few cases of adverse effects have been reported.¹⁰ Long-term accumulation of Gd(III) in patients that received multiple doses was also described, which stimulated the search for alternative CAs.¹¹

High-spin Mn(II) possesses five unpaired electrons that originate a symmetrical ^6S ground state term for the free ion. This leads to a slow electronic relaxation, making Mn(II) an efficient relaxation agent.¹² Thus, it is not surprising that Mn(II) complexes were considered as CA candidates with the advent of MRI in the 1970s and 1980s.^{13,14} The Mn(II)-based

CA $[\text{Mn}(\text{DPDP})]^{4-}$ ($\text{DPDP}^{6-} = N,N'$ -dipyridoxyethylenediamine- N,N' -diacetate- $5,5'$ -bis(phosphate), **L11**, Chart 1)¹⁵ was also introduced in clinical practice for liver imaging,^{16,17} though its use has been discontinued due to poor sales.¹⁸ Nevertheless, the problems associated with Gd(III) toxicity and deposition have provoked a renewed interest in Mn(II)-based MRI contrast agents.¹⁹ One of the main aims of the research in this field is to develop stable and inert complexes endowed with high relaxation efficiencies.^{20–29}

In a recent paper, we proposed an empirical correlation to predict and rationalize the thermodynamic stabilities of Gd(III) complexes with polyaminopolycarboxylate ligands.³⁰ We showed that the thermodynamic stability constants and pGd values can be approximated to a good accuracy using structural descriptors. The contribution of each structural descriptor (ligand motif) to complex stability was obtained by a least-squares fitting procedure to stability data reported in the literature. The stability constants and pGd values were

Received: July 6, 2022

Published: August 22, 2022



Chart 1. Nonmacrocyclic Ligands Discussed in the Text

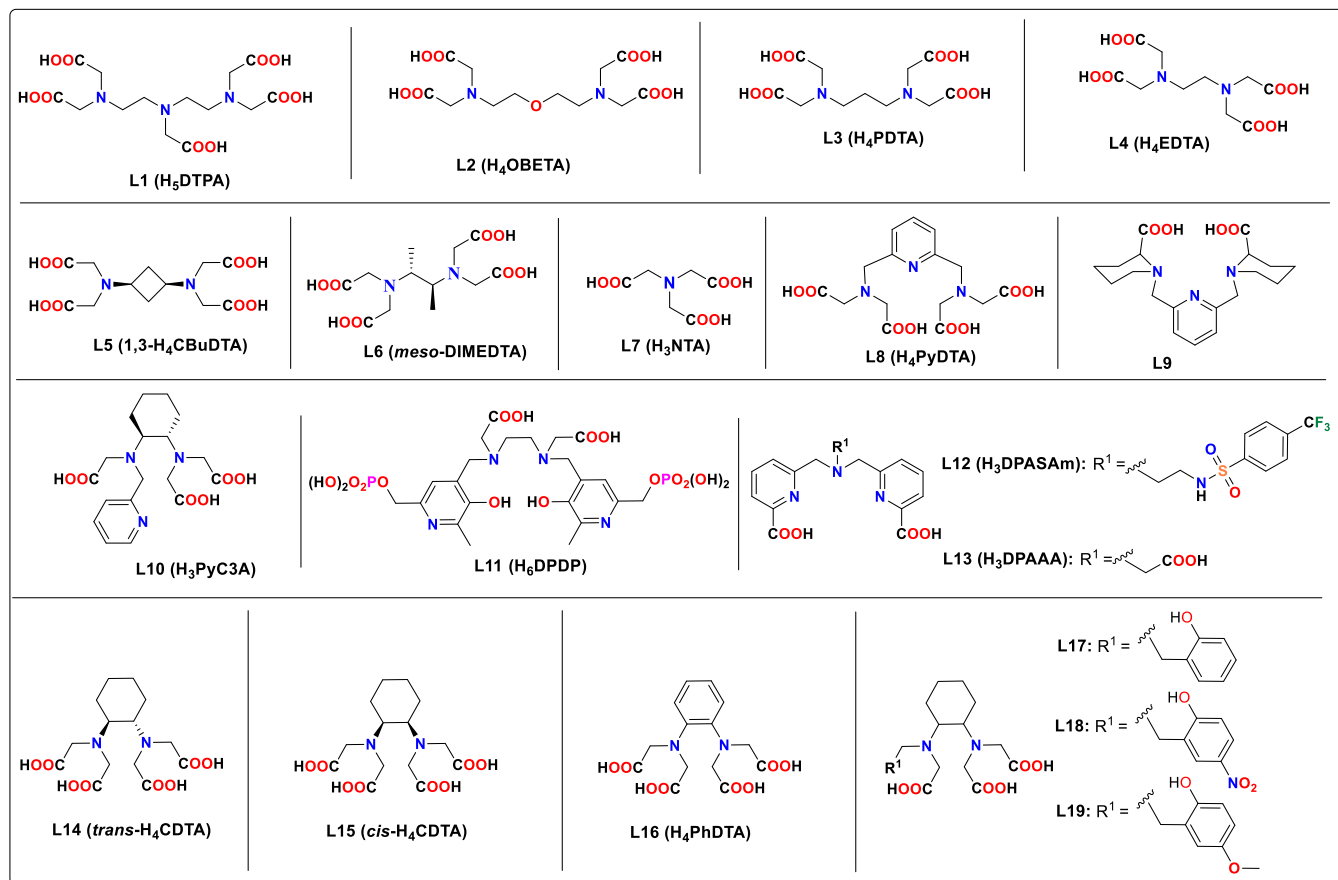
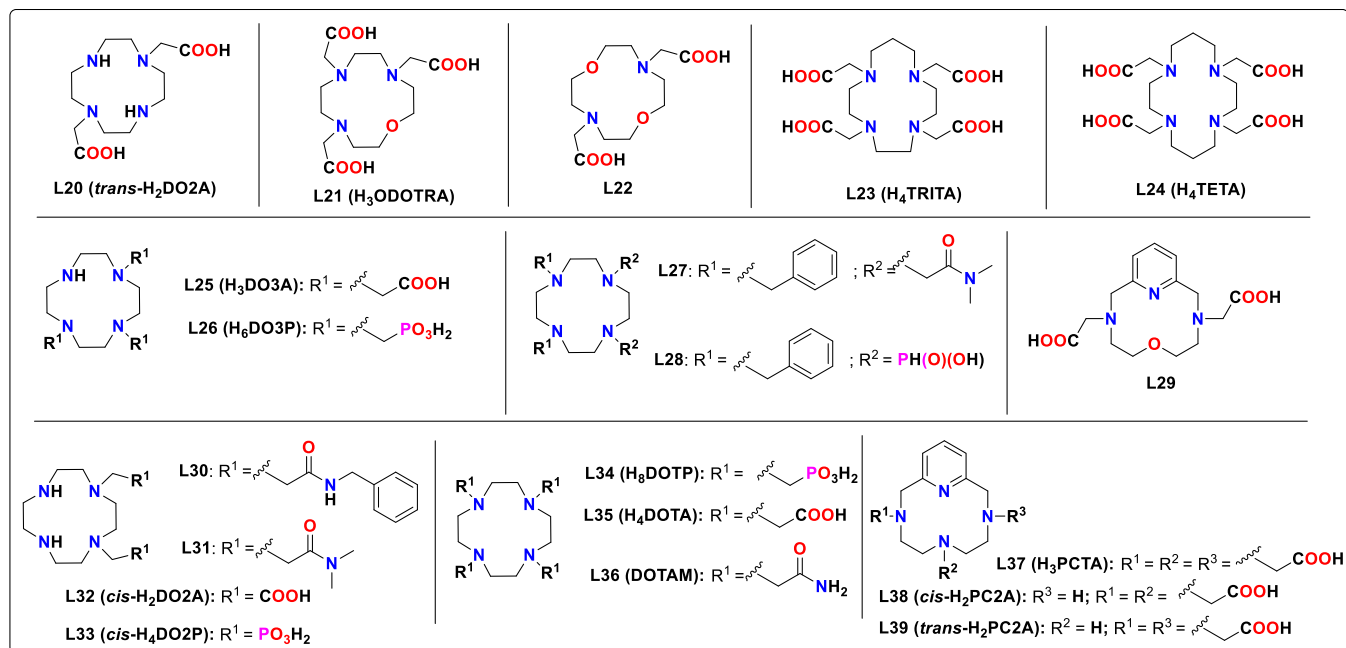


Chart 2. Macrocyclic Ligands Discussed in the Text



subsequently obtained by adding the contributions of each structural descriptor. We validated the predictive character of the model by determining the stability constants of a test set of complexes. The prediction of Mn(II) complex stability is of great interest to aid ligand design and reduce synthetic efforts.

Thus, we envisaged to extend to Mn(II) the methodology developed to predict Gd(III) complex stabilities.

This paper presents an overview of the stability constants of Mn(II) complexes reported in the literature, which are subsequently used to develop an empirical correlation

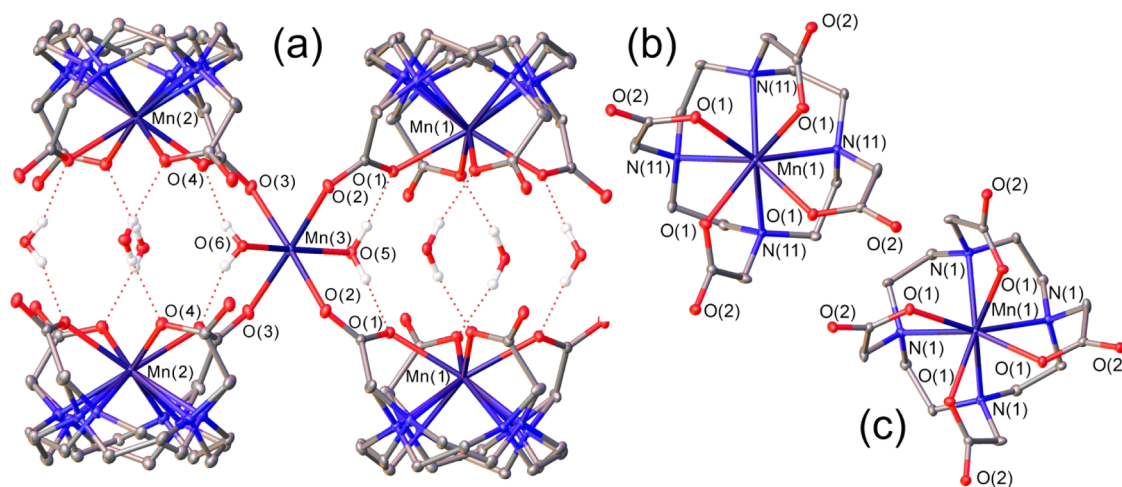


Figure 1. View of the crystal structure of $[\text{Mn}_2(\text{DOTA})(\text{H}_2\text{O})_2]$ showing the $[\text{Mn}(\text{DOTA})]^{2-}$ units joined by six-coordinate exocyclic Mn(II) (a), and views of the square antiprismatic (SAP) (b) and twisted square antiprismatic (TSAP) (c) isomers of $[\text{Mn}(\text{DOTA})]^{2-}$ complexes. Oak Ridge thermal-ellipsoid plot (ORTEP)⁶⁹ plots are at the 30% probability level.

mentioned above. The contributions of the different structural motifs are then discussed and compared with those reported previously for Gd(III). To support our analysis, we also report here the X-ray structure of the Mn(II) complex of DOTA^{4-} , which displays eight-coordinate Mn(II) ions.

RESULTS AND DISCUSSION

Coordination Numbers in Mn(II) Complexes. The analysis described in this work assumes that the thermodynamic stability of Mn(II) complexes can be predicted by adding the contributions of the different donor groups present in the ligand structure. However, the number of donor groups that contribute to complex stability is limited by the coordination number of the metal ion. Once the coordination sphere is saturated, the incorporation of additional donor groups into the ligand scaffold is not expected to contribute to an increased stability. The metal ion in Mn(II) complexes with polyaminopolycarboxylate ligands generally displays coordination numbers 6 or 7. Depending on the denticity of the ligand, water molecules present in the first coordination sphere may complete the metal coordination environment. Heptacoordinated metal complexes are relatively rare within the first-row transition-metal series but are more abundant for Mn(II) than for any other metal ion within the series.^{31,32} Typical seven-coordinate Mn(II) complexes are those with EDTA^{4-} (L4, Chart 1) and its derivatives,³³ in which a coordinated water molecule completes the metal coordination sphere.^{34–37} A remarkable example of this class is $\text{H}_3\text{PyC3A}$ (L10, Chart 1), which forms a very stable Mn(II) complex that generates excellent MRI contrast.^{38,39} Seven coordination is also favored by 15-membered macrocyclic ligands containing five donor atoms, which generally provide pentagonal bipyramidal coordination in which the equatorial positions are occupied by the donor atoms of the macrocyclic unit.^{40,41} The Mn(II) complex with the triethyl ester derivative of the 12-membered macrocycle H_3PCTA (L37, Chart 2) was also found to be seven-coordinated in the solid state.⁴² However, complexes with 1,4,7-triazacyclononane derivatives such as H_3NOTA and related ligands generally form six-coordinate complexes in solution.^{43–45}

Eight-coordinate Mn(II) complexes are rare.^{46,47} Eight-coordinate bispidine derivatives providing exceptionally high

thermodynamic stabilities ($\log K_{\text{MnL}} > 24$) have been reported recently.⁴⁸ Remarkable examples in the context of MRI contrast agents are the cyclen derivatives $[\text{Mn}(\text{DOTAM})]^{2+}$ (DOTAM = L36, Chart 2) and the analogue containing four 2-pyridyl pendants, which were both found to exhibit eight-coordinate metal ions in the solid state.^{49,50} However, it is not clear whether eight coordination is favored for these complexes by the presence of charge-neutral pendant arms. The charge-neutral $[\text{Mn}(\text{cis-DO2A})]$ complex (see L32, Chart 2) was found to display seven-coordinate Mn(II) ions in solution, thanks to the presence of a coordinated water molecule.⁵¹ A seven-coordinate structure was also observed in the solid state.⁵² The incorporation of a third acetic acid pendant into the cyclen structure to give $\text{H}_3\text{DO3A}$ (L25, Chart 2) yields a seven-coordinate Mn(II) complex, as demonstrated by relaxometric studies.⁵¹ The coordination number of the metal ion in $[\text{Mn}(\text{DOTA})]^{2-}$ (see L35, Chart 2) was never ascertained, though the small zero field splitting evidenced by electron paramagnetic resonance (EPR) measurements is compatible with a rigid and symmetrical coordination environment.^{53,54}

Crystal Structure of $[\text{Mn}_2(\text{DOTA})(\text{H}_2\text{O})_2]$. Crystals with formula $[\text{Mn}_2(\text{DOTA})(\text{H}_2\text{O})_2]$ were obtained from an aqueous solution of the $[\text{Mn}(\text{DOTA})]^{2-}$ complex in the presence of 1 equiv of Mn(II). The compound crystallizes in the tetragonal $P4/m$ space group. Crystals contain $[\text{Mn}(\text{DOTA})]^{2-}$ entities joined by exocyclic Mn(II) ions with octahedral coordination, provided by four oxygen atoms of bridging $\mu_2-\eta^1:\eta^1$ -carboxylate groups⁵⁵ and two coordinated water molecules (Figure 1). The Mn–O distances involving the coordinated water molecules (Table 1) are close to those observed in the solid state for $(\text{NH}_4)_2[\text{Mn}(\text{H}_2\text{O})_6](\text{SO}_4)$.⁵⁶ The coordinated water molecules are involved in hydrogen bonds with oxygen atoms of the carboxylate groups.

The macrocyclic fragment in each of the $[\text{Mn}(\text{DOTA})]^{2-}$ entities is disordered into two positions, which correspond to the two square [3333] conformations⁵⁷ of cyclen that are usually denoted as $(\delta\delta\delta\delta)$ and $(\lambda\lambda\lambda\lambda)$.⁵⁸ Interestingly, the position of the pendant arms is not disordered, adopting either a Δ or a Λ conformation for each $[\text{Mn}(\text{DOTA})]^{2-}$ entity.⁵⁹ As a result, the two disordered macrocyclic units generate the $\Delta(\lambda\lambda\lambda\lambda)$ and $\Delta(\delta\delta\delta\delta)$ isomers, which provide square

Table 1. Bond Distances (Å) and Angles (deg) Observed in the Crystal Structure of [Mn₂(DOTA)(H₂O)₂]

Mn(1)–O(1)	2.2867(17)	Mn(2)–O(4)	2.2845(18)
Mn(1)–N(1)	2.462(4)	Mn(2)–N(3)	2.449(3)
Mn(1)–N(11)	2.416(7)	Mn(2)–N(31)	2.398(14)
Mn(1)–N ₄ SAP ^a	1.218	Mn(2)–N ₄ SAP ^a	1.184
Mn(1)–N ₄ TSAP ^a	1.357	Mn(2)–N ₄ TSAP ^a	1.308
ϕ , SAP ^b	43.0	ϕ , SAP ^b	41.2
ϕ , TSAP ^b	26.6	ϕ , TSAP ^b	25.5
Mn(3)–O(2)	2.2154(18)	Mn(3)–O(3)	2.2153(19)
Mn(3)–O(5)	2.198(2)	Mn(3)–O(6)	2.186(3)

^aDistance between the metal ion and the plane defined by the four N atoms of the macrocycle (N₄). ^bTwist angle of the O₄ and N₄ planes.

antiprismatic (SAP) and twisted square antiprismatic (TSAP) coordination polyhedra, respectively.^{60,61} An inversion center relates the $\Delta(\lambda\lambda\lambda\lambda)/\Lambda(\delta\delta\delta\delta)$ and $\Delta(\delta\delta\delta\delta)/\Lambda(\lambda\lambda\lambda\lambda)$ enantiomeric pairs of [Mn(DOTA)]²⁻. Thus, all four stereoisomers of the complex are present in the crystal lattice. The crystal lattice contains two [Mn(DOTA)]²⁻ entities with slightly different bond distances (Table 1). The twist angles (ϕ) of the O₄ and N₄ planes that define the square planes of the coordination polyhedra are close to those expected for SAP (45°) or TSAP (22.5°) coordination. The Mn–N distances in TSAP isomers are ~0.05 Å longer than in the corresponding SAP isomers. As a result, the Mn(II) ions reside closer to the N₄ plane in the SAP isomers than in the TSAP counterparts (see Mn–N₄ distances in Table 1). All of these structural features parallel those observed for DOTA-type complexes of the lanthanide ions and Sc(III).^{62–64} The Mn–O distances are longer than those involving carboxylate oxygen atoms in seven-coordinate Mn(II) complexes (ca. 2.17–2.27 Å),^{24,52,65–67} as a result of the higher coordination number of the metal ion in [Mn(DOTA)]²⁻. A similar situation is observed for the Mn–N bonds, which fall within the range 2.33–2.45 Å for seven-coordinate complexes containing amine N atoms.^{52,67,68}

Stability Constants of Mn(II) Complexes. The literature reports a wide collection of thermodynamic stability constants determined for Mn(II) complexes with a wide variety of ligands. In this study, we aimed at predicting the stability constants of Mn(II) complexes, relevant as MRI contrast agents, using structural descriptors. Thus, we included in our study ligands that contain structural motifs present in the ligands used for this purpose. Among the nonmacrocylic ligands, two families have been widely investigated for Mn(II) complexation. The first class comprises H₄EDTA (L4, Chart 1) and its derivatives, including: (1) EDTA analogues in which the ethyl spacer is replaced by propyl (H₄PDTA, L3),⁷⁰ cyclohexyl (H₄CDTA, L14),⁷¹ phenyl (H₄PhDTA, L16),⁷² 1,2-cyclobutyl,⁷³ or 1,3-cyclobutyl⁷⁴ (H₄CBDTA, L5) groups. (2) Ligands bearing one of these spacers in which some of the acetic acid arms are replaced by donor groups such as phenols (i.e., L17–L19) or pyridine groups (i.e. H₃PyC3A, L10). (3) Extended H₄EDTA structures such as those of H₄OBETA (L2),⁷⁵ H₄PyDTA (L8),⁷⁶ and H₄EGTA.⁷⁵ (4) Ligands related to the latter three classes in which some donor groups are absent.⁷⁶ A more exhaustive list of ligands and their protonation and stability constants is provided in Table S1 (Supporting Information).

A second ligand family comprises tripodal ligands, in which the different donor groups, up to three, are appended on an amine nitrogen atom. This family includes H₃NTA⁷⁷ (L7) and

derivatives in which acetic acid groups are replaced by different donors like picolinic acid (i.e. H₃DPAAA, L13),²¹ sulphonamide (i.e. H₃DPASAm, L12)⁷⁸ or methylphosphonic acid⁷⁹ groups. Table 2 presents the stability constants reported for Mn(II) complexes of selected nonmacrocylic ligands.

The class of macrocylic ligands that have been more extensively investigated for Mn(II) complexation is certainly the family of tetraazamacrocycles (Chart 2), more commonly cyclen (1,4,7,10-tetraazacyclododecane),^{80–82} cyclam (1,4,8,11-tetraazacyclotetradecane),^{83,84} or pycen (3,6,9,15-tetraazabicyclo[9.3.1]pentadeca-1(15),11,13-triene)^{26,85,86} functionalized with different pendant arms, typically acetic acid, primary or N-substituted acetamides, methylphosphonic, methylphosphinic, or picolinic acid groups, among others. Some of these macrocycles incorporate ether oxygen atoms into the macrocylic structure replacing some of the amine N atoms.⁸⁷ Alternatively, macrocylic ligands derived from 1,4,7-triazacyclononane (TACN) functionalized with different pendant arms, and often with mixed N/O donor sets in the macrocylic scaffold, can be used as ligands for Mn(II) (i.e. L43, Chart 3).^{88–90} The structurally related 15-membered macrocylic ligands containing mixed N/O donor sets form rather stable complexes as well (i.e. L47–L50, Chart 3).^{40,41} Macrocylic ligands of this family incorporating acetic acid pendant arms were also investigated for Mn(II) complexation.⁹¹ Finally, the stability of Mn(II) complexes with a few mesocyclic ligands derived from AAZTA was also explored (i.e. L44–L46, Chart 3).⁹²

The thermodynamic stability of Mn(II) complexes depends on several factors, as illustrated in Figure 2. Stability constants generally increase with increasing ligand denticity, as would be expected. The highest log *K*_{MnL} values are observed for complexes with hepta- and particularly octadentate ligands. This suggests that several Mn(II) complexes display coordination number eight in solution, as, for instance, the bispidines reported recently by Comba⁴⁸ and some cyclen derivatives such as [Mn(DOTA)]²⁻. In the latter case, stability constants of log *K*_{MnL} = 20.2 and 19.9 were determined using ionic strengths of 0.1 M Me₄N(NO₃)⁹³ and Me₄NCl,⁵² while that reported for [Mn(DO3A)]⁻ in 0.1 M Me₄NCl is slightly lower (log *K*_{MnL} = 19.4).⁵² The log *K*_{MnL} values determined for a given ligand denticity spread over several orders of magnitude, highlighting the critical effect of ligand topology and the nature of the donor groups incorporated into the ligand scaffold. The data shown in Figure 2 also evidence that macrocylic ligands derived from the 12-membered macrocycles pycen, and particularly cyclen, tend to form Mn(II) complexes with higher stabilities than other ligand classes, with the exception of the bispidine ligands described recently,⁴⁸ which form extraordinary stable Mn(II) complexes. Taken together, the stability constants reported for Mn(II) complexes span more than 20 orders of magnitude, from log *K*_{MnL} values of ca. 1–3 for simple bidentate ligands (i.e. picolinic acid)⁹⁴ to log *K*_{MnL} ~ 24 for the mentioned bispidine complexes.

Structural Descriptors. The descriptors used to predict the thermodynamic stability of Mn(II) complexes are essentially those used previously for Gd(III)³⁰ and are listed in Table 3. Linear polyaminopolycarboxylate ligands are described by the corresponding number of amine N atoms, denoted as N, and the number of acetic acid groups C. For instance, H₄EDTA is described as nN = 2 + nC = 4, where n indicates the number of groups of a given class. In the case of macrocylic ligands, the macrocylic unit as a whole, including

Table 2. Stability Constants ($\log K_{\text{MnL}}$ Values, 25 °C), Values of pMn, Structural Descriptors, and Calculated $\log K_{\text{MnL}}$ and pMn Values for Mn(II) Complexes

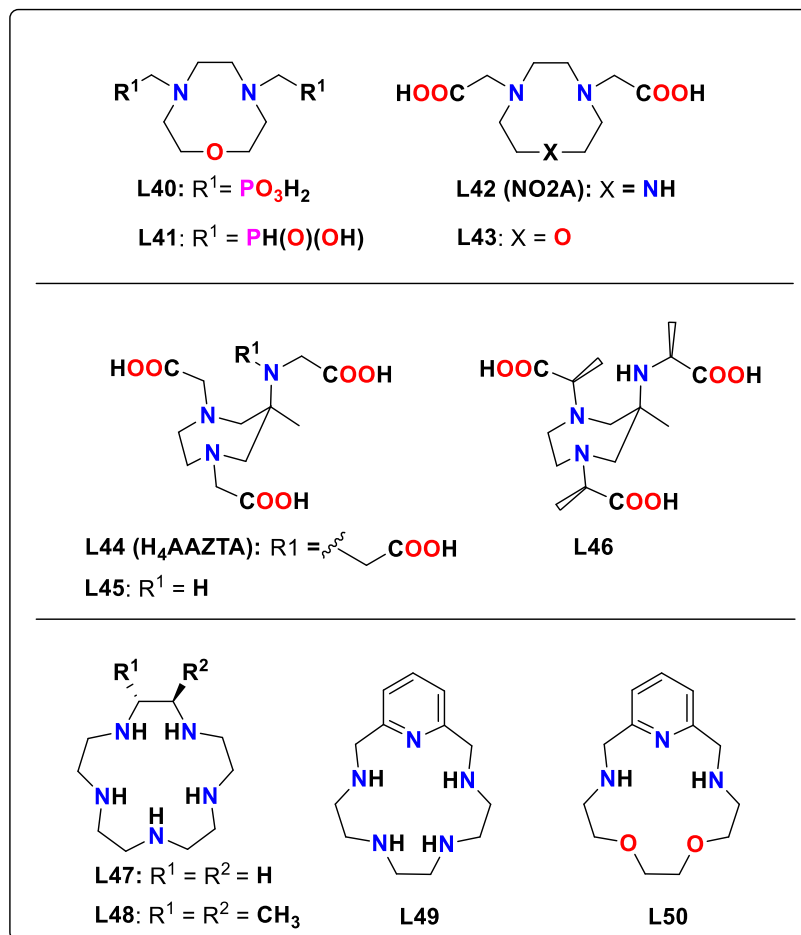
	$\log K_{\text{MnL}}$	pMn	descriptors ^e	ref	$\log K_{\text{MnL}}^{\text{calc}}$	pMn ^{calc}
L1 (H ₃ DTPA)	15.50	12.07	3N + 5C	102	17.12 (14.47 ^d)	11.97 ^d
	14.54	11.88		71		
L2 (H ₄ OBETA)	13.57	11.00	3N + 4C + SOe	75	12.39	11.12
L3 (H ₄ PDTA)	10.01	7.44	2N + 4C + SProp	103	10.41	8.66
L4 (H ₄ EDTA)	12.46	11.62	2N + 4C	71	13.18	11.50
L5 (1,3-H ₄ CBuDTA)	10.78	9.44	2N + 4C + SCBu	74	11.34	9.95
L6 (meso-DIMEDTA)	14.10	11.20	2N + 4C + 2SCalk	70	13.38	11.76
L7 (H ₃ NTA)	7.44	6.36	N + 3C	79	9.24	8.39
L8 (H ₄ PyDTA)	14.13	13.39	3N + 4C + SPy	76	14.74	13.98
L9	11.37	10.83	3N + 2C + 2SCyhx + SPy	99	11.88	10.14
L10 (H ₃ PyC3A)	14.14	12.29	2N + 3C + Py + SCyhx	20	13.26	11.87
L11 (H ₆ DPDP)	15.10	10.34	2N + 2C + 2Phe	15	14.16	8.46
L12 (H ₃ DPASAm)	13.53	11.55	N + 2Pic + Sulph	78	13.76	11.48
L13 (H ₃ DPAAA)	13.19	13.91	N + C + 2Pic	21	13.18	13.07
L14 (trans-H ₄ CDTA)	14.32	13.59	2N + 4C + SCyhx	71	14.40	12.22
L15 (cis-H ₄ CDTA)	14.19	11.54	2N + 4C + SCyhx	98	14.40	12.22
L16 (H ₄ PhDTA)	11.79	12.67	2N + 4C + SPh	72	11.79	11.77
L17	14.16	9.21	2N + 3C + Phe	95	13.67	9.98
L18	13.66	11.06	2N + 3C + PheNO ₂	95	11.86	10.17
L19	14.61	8.41	2N + 3C + PheOMe	95	13.35	8.24
L20 (trans-H ₂ DO2A)	14.64	8.95	A ₁₂ + 2C	80	14.95	10.05
L21 (H ₃ ODOTRA)	13.88	13.09	A ₁₂ + 3C + SOe	81	15.52	11.92
L22	9.38	9.32	A ₁₂ + 2C + 2SOe	87	10.79	8.51
L23 (H ₄ TRITA)	16.74	11.41	A ₁₂ + 4C + Spropyl	93	17.48	12.49
L24 (H ₄ TETA)	11.27	6.51	A ₁₂ + 4C + 2Spropyl	93	14.71 (12.06 ^b)	7.01 ^b
L25 (H ₃ DO3A)	19.43	13.68	A ₁₂ + 3C	104	17.60	12.69
L26 (H ₆ DO3P)	17.45	8.82	A ₁₂ + 3Pho	81	19.91 (16.49 ^c)	
L27	11.54	7.91	A ₁₂ + 2A _{NR2}	96	12.19	9.29
L28	9.39	6.58	A ₁₂ + 2Phosphi	82	10.13	6.59
L29	13.03	10.99	A ₁₂ + 2C + SOe + SPy	105	13.14	11.57
L30	10.72	9.34	A ₁₂ + 2 A _{NHR}	96	11.49	8.91
L31	12.64	9.83	A ₁₂ + 2 A _{NR2}	96	12.19	9.29
L32 (cis-H ₂ DO2A)	15.22	9.99	A ₁₂ + 2C	52	14.95	10.05
L33 (cis-H ₄ DO2P)	15.41	7.41	A ₁₂ + 2Pho	82	16.49	9.15
L34 (H ₈ DOTP)	18.98	8.64	A ₁₂ + 4Pho	81	23.33 (16.49 ^c)	
L35 (H ₄ DOTA)	19.44	13.95	A ₁₂ + 4C	104	20.25	15.33
L36 (DOTAM)	11.96	12.65	A ₁₂ + 4A _{NH2}	81	12.01	12.01
L37 (H ₃ PCTA)	16.83	15.13	A ₁₂ + 3C + SPy	81	17.87	14.70
L38 (cis-H ₂ PC2A)	15.53	12.15	A ₁₂ + 2C + SPy	86	15.22	12.06
L39 (trans-H ₂ PC2A)	17.09	13.18	A ₁₂ + 2C + SPy	86	15.22	12.06
L40	10.61	6.35	A ₉ + 2Pho + SOe	88	10.55	6.39
L41	4.30	6.09	A ₉ + 2Phosphi + SOe	88	4.19	^d
L42 (H ₂ NO2A)	11.56	8.02	A ₉ + 2C	106	11.09	8.06
L43	7.73	6.13	A ₉ + 2C + SOe	107	9.01	7.29
L44 (H ₄ AAZTA)	14.19	12.52	A _{AAZTA} + 4C	92	13.89	12.01
L45	11.00	9.10	A _{AAZTA} + 3C	92	11.24	9.37
L46	10.67	8.72	A _{AAZTA} + 3C α	92	10.91	8.50
L47	10.85	7.03	A ₁₅	101	10.88	6.55
L48	11.09	6.92	A ₁₅ + 2SCalk	101	11.08	6.77
L49	10.89	8.67	A ₁₅ + SPy	40	11.15	8.56
L50	7.18	6.40	A ₁₅ + 2SOe + SPy	40	6.99	7.02

^aCalculated for 3N + 4C. ^bValue calculated for A₁₂ + 4C + 2Spropyl. ^cCalculated for A₁₂ + 2Pho. ^dExcluded from the fit because the complex is nearly fully dissociated under the conditions used to define pMn. ^eDescriptors detailed in Table 3.

the donor atoms, is represented by a single descriptor denoted as A₉, A₁₂, and A₁₅ for triaza-, tetraaza-, and pentaaza-macrocycles, respectively. These descriptors are intended to catch the peculiarities of each macrocyclic unit in terms of not only the number of donor atoms but also the match between

the size of the cavity and that of the Mn(II) ion. Thus, H₄DOTA is described as A₁₂ + 4C, while H₂NO₂A is defined as A₉ + 2C. Similarly, we used a descriptor A_{AAZTA} to account for the ligand 6-amino-6-methylperhydro-1,4-diazepine fragment. Thus, H₄AAZTA (L44, Chart 3) is described as A_{AAZTA}

Chart 3. Representative Examples of Ligands Derived from TACN, AAZTA, and 15-Membered Macrocycles



+ 4C. Bispidine derivatives were excluded from the analysis presented below due to the scarce thermodynamic data reported for this family of complexes.

The different donor groups incorporated into linear or macrocyclic structures are associated with the following descriptors: methylphosphonic acid (Pho), methylphosphinic acid (Phosphi), hydroxyethyl (HE), 2-methylpyridine (Py), primary acetamide (A_{NH_2}), secondary acetamide (A_{NHR}), tertiary acetamide (A_{NR_2}), 2-propionic acid or α -substituted acetic acid (C_α), 2-methylpicolinic acid (Pic), and ethylsulphonamide (Sulph). The stability constants reported for 2-methylphenol derivatives appear to be very sensitive to the nature of the substituent at position 4. Indeed, the stability constants of derivatives containing the electron withdrawing $-\text{NO}_2$ substituent (i.e. L18, Chart 1) are lower than those of unsubstituted derivatives (L17), which in turn are lower than the stability of derivatives containing an electron-donating $-\text{OMe}$ substituent (L19, see stability constants in Table 2).⁹⁵ We thus used three structural descriptors for 2-methylphenol groups, denoted as Phe, PheNO₂, and PheOMe. Similarly, the use of different descriptors for primary (A_{NH_2}), secondary (A_{NHR}), and tertiary (A_{NR_2}) acetamides is justified by the stability constants of complexes with ligands such as L30, L31,^{96,97} and pyclen derivatives with amide groups,⁸¹ which indicate that complex stability increases upon increasing the number of alkyl substituents. Ligands containing α -substituted acetic acid arms generally provide Mn(II) complexes with slightly lower stability than the parent derivatives (i.e. L45 and

L46),⁹² and thus we used two different descriptors to consider the effect of α -substitution.

The comparison of the stability constants reported for H_4EDTA derivatives bearing different spacers evidences that this structural modification has an important impact on the thermodynamic stability constants. The incorporation of a cyclohexyl ring (i.e. H_4CDTA , L14) results in increased stability, while all remaining modifications result in lower $\log K_{\text{MnL}}$ values than for the parent complex. We note that the use of 1,2-cyclobutyl or *cis*-1,3-cyclobutyl spacers yields complexes with very similar stabilities.^{73,74} Thus, these structural modifications, consisting in replacing an ethyl group such as that in H_4EDTA by a cyclobutyl ring, are described by the same structural descriptor SCYbu. Similarly, the complexes of *cis*- and *trans*- H_4CDTA give also very similar values of $\log K_{\text{MnL}}$ ⁹⁸ and thus these structural modifications are described by a single structural descriptor SCYhx. Additionally, the same descriptor was used to account for the incorporation of a piperidine ring into the ligand scaffold (see L9, Chart 1).⁹⁹ Similarly, the incorporation of a phenyl ring (i.e. H_4PhDTA , L16) and a propyl chain (i.e. H_4PDTA , L3) are denoted as SPh and Spropyl, respectively. The same descriptors are employed to account for the introduction of phenyl or propyl groups into macrocyclic units, for instance, the propyl chains in H_4TRITA (L23) and H_4TETA (L24).⁹³

A rather common structural modification introduced to macrocyclic systems consists in replacing amine N atoms by ether oxygen atoms (L21,⁸¹ L22,⁸⁷ L40,⁸⁸ L41,⁸⁸ L50⁴⁰) or

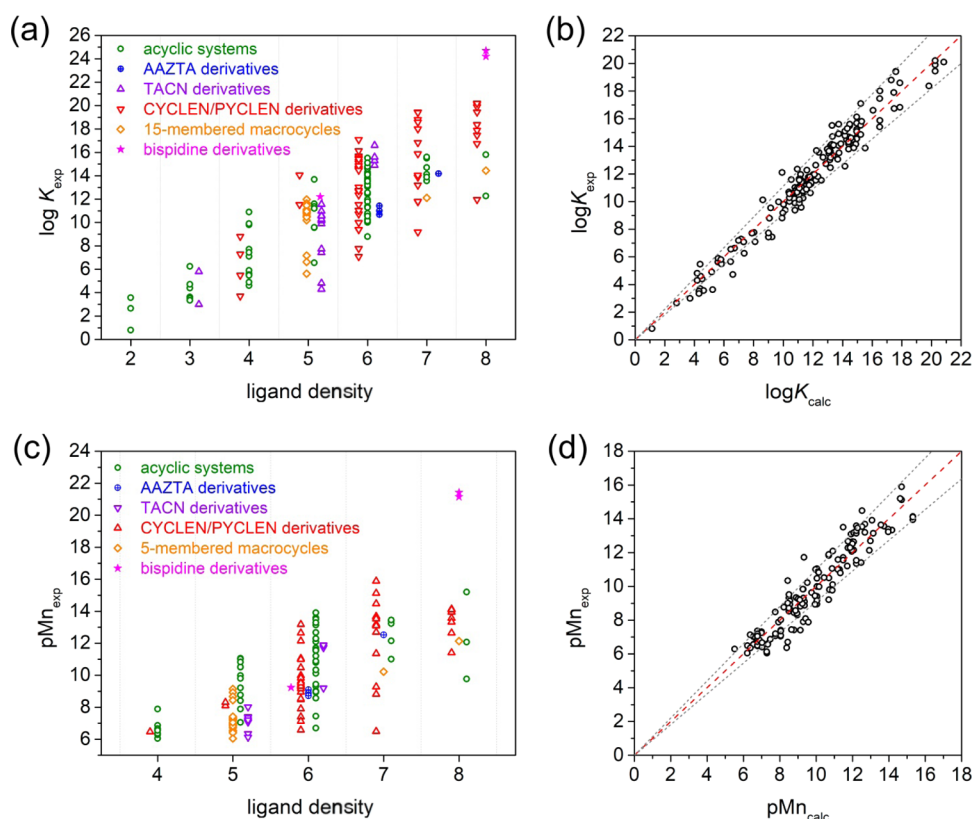


Figure 2. (a) Stability constants ($\log K_{\text{MnL}}$ values) and (c) pMn values of Mn(II) complexes classified according to ligand denticity for different structural families. (b) Plot of the $\log K_{\text{MnL}}$ values reported in the literature (168 values) versus those calculated using eq 1 and (d) plot of pMn values (141 values) versus those calculated using eq 2. Dashed lines represent the lines of identity, while the area within gray dotted lines corresponds to deviations $\pm 10\%$ between experimental and calculated values.

Table 3. Structural Descriptors Used for the Prediction of Mn(II) Complex Stability

N	amine N atom
Pho	methylphosphonic acid
Phosphi	methylphosphinic acid
C	acetic acid
HE	hydroxyethyl
C_α	α -alkyl acetic acid
A_{NH_2}	primary acetamide
A_{NHR}	secondary acetamide
A_{NR_2}	tertiary acetamide
Pic	2-methylpicolinic acid
Ph	2-methylphenol
PhNO ₂	2-methyl-4-nitrophenol
PhOMe	2-methyl-4-methoxyphenol
Sulph	ethylsulfonamide
Py	2-methylpyridine
SCalk	C-alkyl substituent
SOe	ether O atom
Spropyl	propyl group
SCyhx	cyclohexyl ring
SPh	phenyl ring
SPy	pyridyl ring
SCybu	cyclobutyl ring
A_9	triazacyclononane ring
A_{12}	tetraazacyclododecane ring
A_{15}	pentaazacyclopentadecane ring
A_{AAZTA}	6-amino-6-methylperhydro-1,4-diazepine moiety

pyridyl rings (i.e. all pycen derivatives, L49 and L50).⁴⁰ These structural modifications are considered by structural descriptors SOe and SPy. Some nonmacrocylic ligands also incorporate these structural motifs, for instance, H₄OBETA (L2)^{75,100} and H₄PyDTA (L8).⁷⁶

The $\log K_{\text{MnL}}$ values of C-alkylated linear and macrocylic complexes, such as *meso*-H₄DIMEDTA (L6)⁷⁰ and L48,¹⁰¹ are slightly higher than the parent nonsubstituted derivatives. Thus, this alteration was considered with an additional descriptor (SCalk).

Prediction of Stability Constants and Conditional Stability. The structural descriptors presented in the previous section were used to estimate the $\log K_{\text{MnL}}$ values of Mn(II) complexes using the following expression

$$\log K_{\text{MnL}} = \sum_i n_i \Delta \log K_i + \sum_j n_j \Delta \log K_j + A_{\text{MnL}} \quad (1)$$

In this expression, n_i is the number of structural descriptors of type i , while $\Delta \log K_i$ represents the contribution to the stability constant of this donor group. The second term accounts for the different structural modifications present in the ligand structure, where n_j is the number modifications of a given type and $\Delta \log K_j$ its contribution to complex stability. For the complexes with acyclic ligands, A_{MnL} was set to zero.

Similarly, we used a similar expression to analyze complex stabilities at pH 7.4 using pMn values (eq 2), which are defined here as $-\log[\text{Mn}]_{\text{free}}$ for a total Mn(II) concentration of $1 \mu\text{M}$ and a total ligand concentration of $[\text{L}]_{\text{tot}} = 10 \mu\text{M}$.

$$pMn = \sum_i n_i \Delta pMn_i + \sum_j n_j \Delta pMn_j + A_{pMn} \quad (2)$$

The values of pMn allow for a comparison of the stabilities of complexes with different ligands at physiological pH, which depend not only on $\log K_{MnL}$ values but also on ligand basicity as well. In principle, one could define pMn using different conditions, for instance, taking equimolar concentrations of ligand and metal ions. We have chosen here the conditions proposed by Raymond,¹¹⁵ which imply using a 10-fold ligand excess. This results in higher pMn values, increasing the number of ligands that provide a pMn value >6, as pMn = 6 corresponds to a fully dissociated system. Figure 2 shows that the pMn values vary in the range of ca. 6–21, and tend to increase with ligand denticity. Bidentate and tridentate ligands generally provide pMn values of 6 for the definition used here, and thus these ligands were excluded in Figure 2c.

A least-squares fit of 168 $\log K_{MnL}$ values reported in the literature to eq 1 provided the contribution to the complex stability of the different structural descriptors, which were used as fitting parameters. The results of the analysis are provided in Table 4. The agreement between the experimental $\log K_{MnL}$ values and those predicted by eq 1 is very good, as shown in Figure 2. The linear fit of the data provides a slope very close to 1 [0.997(5)], as would be expected, with a Pearson's correlation coefficient of 0.9978. The mean deviation of the calculated data with respect to the experimental values is only

Table 4. Contributions of the Different Structural Descriptors to $\log K_{MnL}$ and pMn Obtained from the Least-Squares Fit of the Stability Data to Equations 1 and 2 and Total Number of Structural Descriptors of Each Type ($\sum n_{ij}$)^a

	$\Delta \log K_{ij}$	$\sum n_{ij}$	ΔpMn_{ij}	$\sum n_{ij}$
N	1.29(0.10)	136	0.47(0.18)	110
Pho	3.42(0.15)	23	2.19(0.18)	16
Phosphi	0.24(0.28)	6	0.91(0.48)	2
C	2.65(0.06)	286	2.64(0.10)	256
HE	-0.15(0.32)	6	0.46(0.66)	2
C α	2.54(0.16)	13	2.35(0.19)	10
A _{NH2}	0.59(0.19)	7	1.81(0.20)	7
A _{NHR}	0.92(0.26)	5	2.07(0.27)	5
A _{NR2}	1.27(0.21)	9	2.26(0.23)	9
Pic	4.62(0.19)	19	4.98(0.21)	17
Phe	3.14(0.24)	9	1.12(0.26)	9
PheNO2	1.33(0.65)	2	1.31(0.67)	2
PheOMe	2.82(0.65)	2	-0.62(0.67)	2
Sulph	3.23(0.54)	3	1.05(0.56)	3
Py	1.51(0.12)	24	2.29(0.16)	15
SCalk	0.10(0.14)	39	0.11(0.15)	36
SOe	-2.08(0.14)	34	-0.77(0.16)	27
Spropyl	-2.77(0.18)	25	-2.84(0.26)	9
SCyhx	1.22(0.27)	12	0.72(0.29)	12
SPh	-1.39(0.47)	4	0.27(0.49)	4
SPy	0.27(0.22)	30	2.01(0.24)	27
SCybu	-1.84(0.44)	5	-1.55(0.46)	5
A ₉	5.79(0.26)	17	2.78(0.34)	12
A ₁₂	9.65(0.22)	59	4.77(0.30)	47
A ₁₅	10.88(0.31)	24	6.55(0.34)	23
A _{AAZTA}	3.29(0.50)	4	1.45(0.57)	4

^aStructural descriptors detailed in Table 3.

0.63. The agreement between the experimental and calculated data is remarkable, as the mean deviation is comparable to the differences in stability constants reported for the same system by independent groups, often using different ionic strengths. For instance, $\log K_{MnL}$ values differing by more than 1.3 and 0.7 log *K* units were reported for H₅DTPA (L1)¹⁰⁸ and H₄EGTA.^{71,75} Similar differences in stability constants were observed for macrocyclic ligands when using different ionic strengths.⁸⁰ Furthermore, the structural descriptors presented above predict identical $\log K_{MnL}$ values for regioisomeric ligands such as *cis*-H₂PC2A (L38) and *trans*-H₂PC2A (L39), which were found to differ by ~ 1.5 log *K* units.⁸⁶ Thus, it is obvious that the stability constants of a given complex are affected not only by the nature of the donor groups present in the ligand scaffold but also by the arrangement of these donor groups in the ligand structure.

The analysis of the $\log K_{MnL}$ values using eq 1 allows us to infer the coordination number of the metal ion in certain complexes, for which the assignment of a given coordination number is ambiguous. For instance, a $\log K_{MnL}$ value of 17.12 is calculated for H₅DTPA (L1) assuming octadentate binding of the ligand to the Mn(II) ion (3N + 5C). However, the value calculated for a seven-coordinate complex (3N + 4C) of 14.47 is in much better agreement with the experimental values of 15.50¹⁰² and 14.54.⁷¹ Thus, this complex is very likely heptacoordinated in aqueous solution, as observed in the solid state for bis(amide) derivatives of H₅DTPA.¹⁰⁹ A similar situation is observed for H₄TETA (L24), for which the $\log K_{MnL}$ value estimated assuming eight coordination (14.71), differs considerably from the experimental value of $\log K_{MnL} = 11.27$.⁹³ A considerably better agreement is observed by assuming the formation of a heptacoordinated complex (Table 2). Most likely the propyl chains present in the ligand structure introduce some steric hindrance around the metal ion, favoring a lower coordination number in comparison with H₄DOTA, as observed for the corresponding Gd(III) complexes.¹¹⁰ The presence of bulky methylphosphonic acid groups in H₆DO3P (L26) and H₈DOTP (L34) appears to favor the formation of six-coordinate complexes in solution (Table 2).⁸¹

The empirical expression obtained here may be useful to aid experimental stability constant determination, as the predicted $\log K_{MnL}$ value can help anticipate the pH range in which complex dissociation is expected to occur. Furthermore, eq 1 can be used to identify stability constant values that are likely to be incorrect. For instance, a stability constant of $\log K_{MnL} = 14.29$ was reported for a pentadentate ligand containing a piperazine ring functionalized with a picolinic acid and an acetic acid function.²⁵ The stability constant predicted with eq 1 using 2N + 1C + 1Pic + 1SCyhx is 11.07. The very large discrepancy between the experimental and calculated values suggests that the experimental stability constant may be incorrect and should be taken with some caution.

The pMn values obtained from 141 complexes were fitted to eq 2 following the same strategy used for stability constants. The number of data points used in this analysis is lower than for $\log K_{MnL}$, as ligands with pMn ~ 6 had to be excluded from the analysis, and for a few systems, ligand protonation constants were not reported together with stability constants. The agreement between the experimental and calculated pMn data is reasonably good (Figure 2), though not as good as for $\log K_{MnL}$ values. The linear fit of the data gives a slope of 0.993(7) and a Pearson's correlation coefficient of 0.9967. The mean deviation of calculated versus experimental data amounts

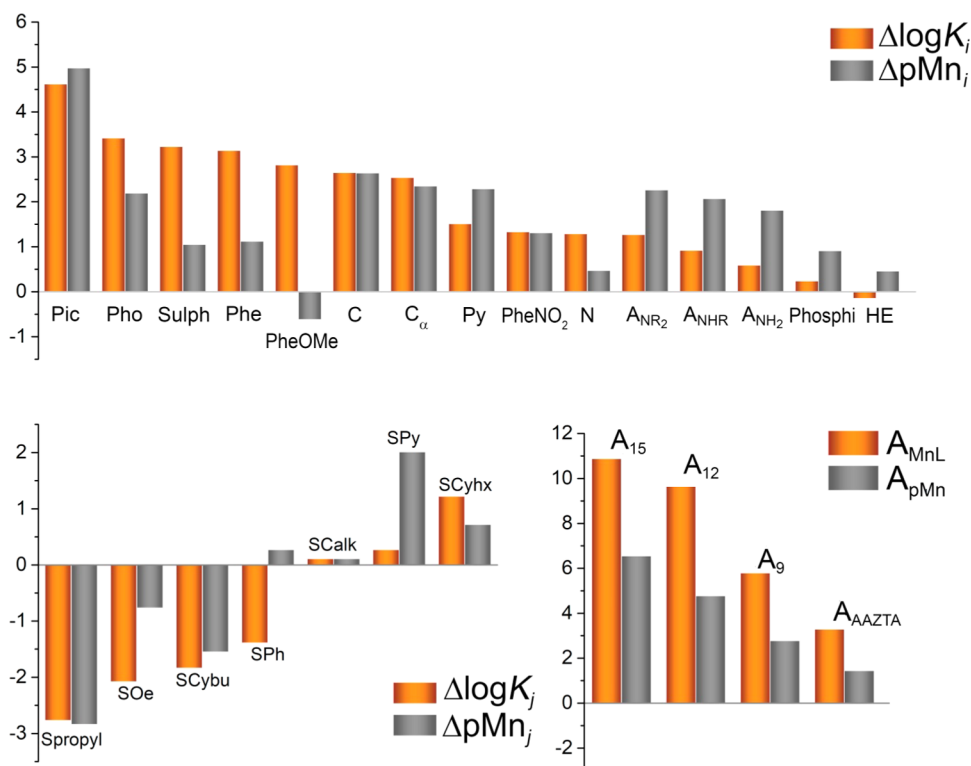


Figure 3. Comparison of the contributions of the different structural descriptors to $\log K_{\text{MnL}}$ and pMn, obtained from the least-squares fit of the stability data to eqs 1 and 2. Structural descriptors detailed in Table 3.

to 0.66 pMn units. We note that the $\Delta \log K_{ij}$ and $\Delta p\text{Mn}_{ij}$ contributions characterizing some structural descriptors were obtained with rather large standard deviations (Table 4). This situation is generally associated with structural motifs that have been seldom incorporated into ligand structures (low $\sum n_{ij}$ values in Table 4).

Analysis of the Structural Descriptors. The contributions of the different structural descriptors to $\log K_{\text{MnL}}$ and pMn provide valuable information that can be used for ligand design. Figure 3 shows the contributions of the different motifs to $\log K_{\text{MnL}}$ and pMn. The group with the highest contribution to $\log K_{\text{MnL}}$ is the picolinic acid moiety (Pic), which is characterized by a $\Delta \log K_i$ value of 4.65. The latter value is significantly higher than the sum of the contributions of an acetic acid (C) and a pyridyl group (Py), which amounts to 4.16. Thus, picolinate units are particularly well suited for stable Mn(II) complexation. Other groups characterized by large $\Delta \log K_i$ contributions are methylphosphonic acid (Pho), ethylsulphonamide (Sulph), and 2-methylphenol groups, either unsubstituted at position 4 (Phe) or bearing a methoxy substituent (PheOMe). However, the high basicity of these groups results in a very significant decrease in their contribution to pMn compared with $\log K_{\text{MnL}}$. In other words, these groups provide a large contribution to the stability constant, but they are also prone to protonation at pH 7.4, which has a very negative impact on the stability of the complex close to physiological pH. Indeed, very high protonation constants were determined for Mn(II) complexes containing methylphosphonate,⁸² ethylsulphonamide,⁷⁸ and phenolate groups,⁹⁵ with $\log K_{\text{MnL}}$ typically >5.5 . This effect is particularly dramatic for PheOMe, which is characterized by $\Delta \log K_i = 2.82$ and $\Delta p\text{Mn}_i = -0.62$. The lower basicity of the phenol group functionalized with a $-\text{NO}_2$ substituent at

position 4 results however in very similar $\Delta \log K_i$ and $\Delta p\text{Mn}_i$ values. The high basicity of amine N atoms (N) also justifies the fact that $\Delta \log K_i > \Delta p\text{Mn}_i$.

The low basicity of acetate groups (C) results in very similar $\Delta \log K_i$ and $\Delta p\text{Mn}_i$ contributions. The introduction of α -alkyl groups (C_α) has a very minor impact in terms of $\Delta \log K_i$, but decreases slightly $\Delta p\text{Mn}_i$, likely because of an enhanced basicity associated with the electron-donating effect of the alkyl substituent. Picolinate groups are known to decrease the overall ligand basicity compared with similar ligands containing acetate groups, explaining that $\Delta \log K_i < \Delta p\text{Mn}_i$ in the latter case.¹¹¹

Donor groups with low basicities are generally characterized by $\Delta \log K_i < \Delta p\text{Mn}_i$, and thus are well suited to increase complex stability at physiological pH. As a result, donor groups such as 2-methylpyridine (Py) and tertiary (A_{NR_2}) and secondary (A_{NHR}) acetamides provide contributions to $\Delta p\text{Mn}_i$ approaching that of carboxylates (C). The contribution of a picolinate group (4.98) is nearly identical to the sum of the contributions of acetate (2.64) and pyridine (2.29).

Concerning the effect of structural modifications, the incorporation of propyl groups (Spropyl), ether oxygen atoms (SOe), or cyclobutyl (SCybu) groups has a very negative impact on both $\log K$ and $\Delta p\text{Mn}$, as evidenced by their negative contributions. Replacing ethylene groups of the ligand backbone by phenyl groups has a negative impact in terms of $\log K$, but results in a slight positive contribution to pMn. The incorporation of a pyridyl group into the ligand scaffold results in improved stability, with a particularly positive effect on pMn. This can be attributed to the lower basicity of pyridine with respect to amine N atoms. Examples of ligands that exploit this effect for stable Mn(II) complexation are H_4PyDTA (L8) derivatives⁷⁶ and $\text{H}_2\text{PC}_2\text{A}$

derivatives L38 and L39.⁸⁶ Cyclohexyl rings have also a beneficial impact on complex stability when replacing ethyl groups of polyaminopolycarboxylate ligands, an effect exploited in the well-known H₃PyC3A ligand (L10), which affords a stable Mn(II) complex with appealing properties as an MRI contrast agent.^{38,39}

The terms describing the contributions of macrocyclic and mesocyclic platforms indicate that 15-membered macrocycles provide the largest contribution to both complex stability and pMn, followed by 12-membered macrocycles, TACN and AAZTA derivatives. The same trend is observed for both log *K* and ΔpMn values. However, one has to consider that these structural motifs contain a different number of donor groups, and thus impose some limitations to the number of additional donor atoms that can be incorporated into the Mn(II) coordination sphere. 15-Membered macrocycles generally favor seven-coordinate complexes, where two additional donor atoms coordinate to the metal ion from different sides of the macrocyclic mean plane. As a result, only one additional donor atom can be incorporated into the ligand scaffold if an inner-sphere water molecule should be present. The TACN unit contains three donor atoms, but Mn(II) complexes based on this platform do not exceed coordination number six, which greatly limits the stability that can be achieved (Figure 2). Thus, 12-membered macrocycles appear to be the best choice among those analyzed here, as they combine rather large *A*_{MnL} and *A*_{pMn} contributions and coordination numbers of seven or even eight in the case of cyclen derivatives, as demonstrated here for [Mn(DOTA)]²⁻.

Figure 4 provides a comparison of the contributions of the different structural descriptors to Mn(II) and Gd(III)³⁰

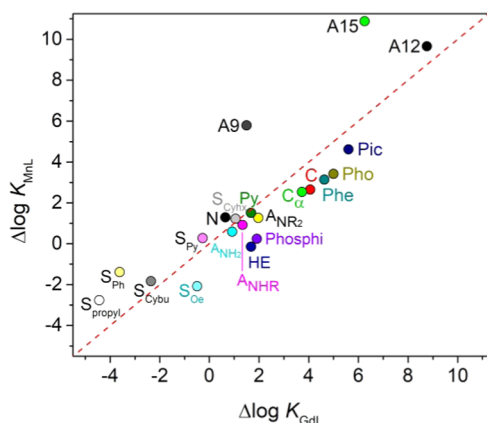


Figure 4. Comparison of the contributions of the different structural descriptors to the stability constants of Gd(III) and Mn(II) complexes. Data above the dashed line provide more favorable contributions to Mn(II) complex stability than to Gd(III).

complex stabilities. The Δlog *K* values of most structural descriptors fall close to the line of identity, indicating that they contribute to a similar extent to Gd(III) and Mn(II) complex stability. However, Gd(III) complexes with high denticity ligands (8–10) are often characterized by higher stability constants than the Mn(II) analogues, due to the higher coordination numbers that the former achieve. 9-Membered and 15-membered macrocyclic units appear to be better suited for Mn(II) than Gd(III) complexation, while 12-membered macrocycles provide similar contributions to the stabilities of complexes with the two metal ions. Hydroxyethylmethyl and

phosphinic acid arms are not adequate for stable Mn(II) complexation. On the contrary, phenyl and propyl spacers are more detrimental to Gd(III) complex stability compared with Mn(II). In the case of Spropyl term, this is related to the strong preference of large metal ions to form five-membered chelate rings.¹¹²

CONCLUSIONS

The reemergence of Mn(II) complexes as MRI contrast agents has stimulated a great amount of work to determine their thermodynamic stabilities.^{48,86,95} Compared to Gd(III)-based MRI contrast agents, the stability constants of Mn(II) complexes are typically lower. High-spin Mn(II) has no ligand field stabilization energy (LFSE), is generally kinetically labile, and forms less stable complexes than biologically important divalent cations like Fe(II), Zn(II), and Cu(II). Thus, for in vivo applications, it is critical to achieve as high a stability as possible to minimize the risk of Mn(II) dissociation in the body. This is also true for emerging applications involving Mn-S2 PET imaging.^{113,114} Concurrently, Mn-based MRI contrast agents require the presence of an inner-sphere water ligand to increase the relaxivity of the complex, and leaving a coordination site vacant for water access may result in lower stability. Finally, the large size of the Mn(II) ion and lack of LFSE results in coordination numbers of 6, 7, and 8 for complexes in aqueous solution, and coordination number is difficult to predict a priori.

Here, we utilized the large body of published stability constant data to establish quantitative structure–stability correlations to predict stability constants, employing the methodology we initially applied to Gd(III). The empirical relations presented here are remarkably accurate despite known differences in experimental stability constants arising from the use of different ionic media and ionic strengths. An interesting result from this study was the prediction of the denticity of the coordinated ligand when multidentate ligands are used, e.g., [Mn(DOTA)]²⁻ has eight-coordinate Mn(II) (confirmed by X-ray crystallography), while the predicted stability constant for [Mn(DTPA)]³⁻ is consistent with seven-coordinate Mn(II). Our approach also serves to identify published stability constant data that may be incorrect. For the development of novel Mn(II) complexes, this method allows the prediction of stability constants and the ability to rule out the synthesis of likely inferior complexes. Finally, the quantification of different structural parameters like donor atom or chelate ring type allows the design of new complexes that might have optimal stability.

Having now applied this methodology to Gd(III) and Mn(II) complexes, it is apparent that this work can be extended to other metal ions for which exist a large body of stability constant data. A limitation of this work is that the method requires stability constant data for specific donors or ligand archetypes. As such, it does not anticipate novel chelators like bispindines.^{27,48} However, these types of ligands can be added to the model once stability constant data are collected for a few examples.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.inorgchem.2c02364>.

Ligand protonation constants and stability constants of Mn(II) complexes and crystallographic data (PDF)

Accession Codes

CCDC 2184025 contains the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

CCDC 2184025 contains the supporting crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, U.K.; fax: +44 1223 336033.

AUTHOR INFORMATION

Corresponding Authors

Aurora Rodríguez-Rodríguez – *Centro de Investigaciones Científicas Avanzadas (CICA) and Departamento de Química, Facultade de Ciencias, Universidade da Coruña, 15071 A Coruña, Galicia, Spain*; orcid.org/0000-0002-4951-4470; Email: aurora.rodriguez@udc.es

Carlos Platas-Iglesias – *Centro de Investigaciones Científicas Avanzadas (CICA) and Departamento de Química, Facultade de Ciencias, Universidade da Coruña, 15071 A Coruña, Galicia, Spain*; orcid.org/0000-0002-6989-9654; Email: carlos.platas.iglesias@udc.es

Authors

Rocío Uzal-Varela – *Centro de Investigaciones Científicas Avanzadas (CICA) and Departamento de Química, Facultade de Ciencias, Universidade da Coruña, 15071 A Coruña, Galicia, Spain*

Francisco Pérez-Fernández – *Centro de Investigaciones Científicas Avanzadas (CICA) and Departamento de Química, Facultade de Ciencias, Universidade da Coruña, 15071 A Coruña, Galicia, Spain*

Laura Valencia – *Departamento de Química Inorgánica, Facultad de Ciencias, Universidade de Vigo, 36310 Pontevedra, Spain*

Peter Caravan – *The Institute for Innovation in Imaging and the A. A. Martinos Center for Biomedical Imaging, Massachusetts General Hospital, Harvard Medical School, Charlestown, Massachusetts 02129, United States*; orcid.org/0000-0002-3179-6537

David Esteban-Gómez – *Centro de Investigaciones Científicas Avanzadas (CICA) and Departamento de Química, Facultade de Ciencias, Universidade da Coruña, 15071 A Coruña, Galicia, Spain*; orcid.org/0000-0001-6270-1660

Complete contact information is available at: <https://pubs.acs.org/10.1021/acs.inorgchem.2c02364>

Author Contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

C.P.-I. and D.E.-G. (PID2019-104626GB-I00) and A.R.-R. (Grant PID2019-108352RJ-I00) thank Ministerio de Ciencia e Innovación and Xunta de Galicia (Grant ED431B 2020/52) for generous financial support. R.U.-V. thanks the Xunta de Galicia (Grant ED481A-2018/314) for funding her Ph.D. contract. L.V. is indebted to CACTI (Universidade de Vigo) for X-Ray measurements. P.C. acknowledges funding from the National Institute of Diabetes and Digestive and Kidney Diseases (R01DK121789).

REFERENCES

- Wahsner, J.; Gale, E. M.; Rodríguez-Rodríguez, A.; Caravan, P. Chemistry of MRI Contrast Agents: Current Challenges and New Frontiers. *Chem. Rev.* **2019**, *119*, 957–1057.
- Li, H.; Meade, T. J. Molecular Magnetic Resonance Imaging with Gd(III)-Based Contrast Agents: Challenges and Key Advances. *J. Am. Chem. Soc.* **2019**, *141*, 17025–17041.
- Lux, J.; Sherry, A. D. Advances in Gadolinium-Based MRI Contrast Agent Designs for Monitoring Biological Processes in Vivo. *Curr. Opin. Chem. Biol.* **2018**, *45*, 121–130.
- Bonnet, C. S.; Tóth, É. Molecular Magnetic Resonance Imaging Probes Based on Ln³⁺ Complexes. In *Advances in Inorganic Chemistry*, Elsevier, 2016; Vol. 68, pp 43–96.
- Aime, S.; Botta, M.; Terreno, E. Gd(III)-Based Contrast Agents for MRI. In *Advances in Inorganic Chemistry*, Elsevier, 2005; Vol. 57, pp 173–237.
- Tircsó, G.; Molnár, E.; Csupász, T.; Garda, Z.; Botár, R.; Kálmán, F. K.; Kovács, Z.; Brücher, E.; Tóth, I. Gadolinium(III)-Based Contrast Agents for Magnetic Resonance Imaging. A Re-Appraisal. In *Metal Ions in Bio-Imaging Techniques*; Sigel, A.; Freisinger, E.; Sigel, R. K. O., Eds.; De Gruyter, 2021; pp 39–70.
- Doan, B.-T.; Meme, S.; Beloeil, J.-C. General Principles of MRI. In *The Chemistry of Contrast Agents in Medical Magnetic Resonance Imaging*; Merbach, A.; Helm, L.; Tóth, É., Eds.; John Wiley & Sons, Ltd.: Chichester, U.K., 2013; pp 1–23.
- Helm, L. Relaxivity in Paramagnetic Systems: Theory and Mechanisms. *Prog. Nucl. Magn. Reson. Spectrosc.* **2006**, *49*, 45–64.
- Luchinat, C. Relaxometry and Paramagnetic Metal Ions in Biological Systems. *Magn. Reson. Chem.* **1993**, *31*, S145–S153.
- Rogosnitzky, M.; Branch, S. Gadolinium-Based Contrast Agent Toxicity: A Review of Known and Proposed Mechanisms. *BioMetals* **2016**, *29*, 365–376.
- Le Fur, M.; Caravan, P. The Biological Fate of Gadolinium-Based MRI Contrast Agents: A Call to Action for Bioinorganic Chemists. *Metallomics* **2019**, *11*, 240–254.
- Botta, M.; Carniato, F.; Esteban-Gómez, D.; Platas-Iglesias, C.; Tei, L. Mn(II) Compounds as an Alternative to Gd-Based MRI Probes. *Future Med. Chem.* **2019**, *11*, 1461–1483.
- Lauffer, R. B. Paramagnetic Metal Complexes as Water Proton Relaxation Agents for NMR Imaging: Theory and Design. *Chem. Rev.* **1987**, *87*, 901–927.
- Koenig, S. H.; Brown, R. D.; Goldstein, E. J.; Burnett, K. R.; Wolf, G. L. Magnetic Field Dependence of Proton Relaxation Rates in Tissue with Added Mn²⁺: Rabbit Liver and Kidney. *Magn. Reson. Med.* **1985**, *2*, 159–168.
- Rocklage, S. M.; Cacheris, W. P.; Quay, S. C.; Hahn, F. E.; Raymond, K. N. Manganese(II) *N,N'*-Dipyridoxylethylenediamine-*N,N'*-Diacetate 5,5'-Bis(Phosphate). Synthesis and Characterization of a Paramagnetic Chelate for Magnetic Resonance Imaging Enhancement. *Inorg. Chem.* **1989**, *28*, 477–485.
- Chung, J.-J.; Kim, M.-J.; Kim, K. W. Mangafodipir Trisodium-Enhanced MRI for the Detection and Characterization of Focal Hepatic Lesions: Is Delayed Imaging Useful? *J. Magn. Reson. Imaging* **2006**, *23*, 706–711.

- (17) Regge, D.; Cirillo, S.; Macera, A.; Galatola, G. Mangafodipir Trisodium: Review of Its Use as an Injectable Contrast Medium for Magnetic Resonance Imaging. *Rep. Med. Imaging* **2009**, *2*, 55–68.
- (18) Cloyd, R. A.; Koren, S. A.; Abisambra, J. F. Manganese-Enhanced Magnetic Resonance Imaging: Overview and Central Nervous System Applications With a Focus on Neurodegeneration. *Front. Aging Neurosci.* **2018**, *10*, No. 403.
- (19) Pan, D.; Caruthers, S. D.; Senpan, A.; Schmieder, A. H.; Wickline, S. A.; Lanza, G. M. Revisiting an Old Friend: Manganese-Based MRI Contrast Agents. *Wiley Interdiscip. Rev.: Nanomed. Nanobiotechnol.* **2011**, *3*, 162–173.
- (20) Gale, E. M.; Atanasova, I. P.; Blasi, F.; Ay, I.; Caravan, P. A Manganese Alternative to Gadolinium for MRI Contrast. *J. Am. Chem. Soc.* **2015**, *137*, 15548–15557.
- (21) Forgács, A.; Pujales-Paradela, R.; Regueiro-Figueroa, M.; Valencia, L.; Esteban-Gómez, D.; Botta, M.; Platas-Iglesias, C. Developing the Family of Picolinate Ligands for Mn²⁺ Complexation. *Dalton Trans.* **2017**, *46*, 1546–1558.
- (22) Regueiro-Figueroa, M.; Rolla, G. A.; Esteban-Gómez, D.; de Blas, A.; Rodríguez-Blas, T.; Botta, M.; Platas-Iglesias, C. High Relaxivity Mn²⁺-Based MRI Contrast Agents. *Chem.—Eur. J.* **2014**, *20*, 17300–17305.
- (23) Loving, G. S.; Mukherjee, S.; Caravan, P. Redox-Activated Manganese-Based MR Contrast Agent. *J. Am. Chem. Soc.* **2013**, *135*, 4620–4623.
- (24) Khannam, M.; Weyhermüller, T.; Goswami, U.; Mukherjee, C. A Highly Stable L-Alanine-Based Mono(Aquated) Mn(II) Complex as a T₁-Weighted MRI Contrast Agent. *Dalton Trans.* **2017**, *46*, 10426–10432.
- (25) Phukan, B.; Mukherjee, C.; Goswami, U.; Sarmah, A.; Mukherjee, S.; Sahoo, S. K.; Moi, S. C. A New Bis(Aquated) High Relaxivity Mn(II) Complex as an Alternative to Gd(III)-Based MRI Contrast Agent. *Inorg. Chem.* **2018**, *57*, 2631–2638.
- (26) Drahoš, B.; Kotek, J.; Císařová, I.; Hermann, P.; Helm, L.; Lukeš, I.; Tóth, É. Mn²⁺ Complexes with 12-Membered Pyridine Based Macrocycles Bearing Carboxylate or Phosphonate Pendant Arm: Crystallographic, Thermodynamic, Kinetic, Redox, and ¹H/ ¹⁷O Relaxation Studies. *Inorg. Chem.* **2011**, *50*, 12785–12801.
- (27) Ndiaye, D.; Sy, M.; Pallier, A.; Mème, S.; Silva, I.; Lacerda, S.; Nonat, A. M.; Charbonnière, L. J.; Tóth, É. Unprecedented Kinetic Inertness for a Mn²⁺-Bispidine Chelate: A Novel Structural Entry for Mn²⁺-Based Imaging Agents. *Angew. Chem., Int. Ed.* **2020**, *59*, 11958–11963.
- (28) Gale, E. M.; Caravan, P. Gadolinium-Free Contrast Agents for Magnetic Resonance Imaging of the Central Nervous System. *ACS Chem. Neurosci.* **2018**, *9*, 395–397.
- (29) Anbu, S.; Hoffmann, S. H. L.; Carniato, F.; Kenning, L.; Price, T. W.; Prior, T. J.; Botta, M.; Martins, A. F.; Stasiuk, G. J. A Single-Pot Template Reaction Towards a Manganese-Based T₁ Contrast Agent. *Angew. Chem., Int. Ed.* **2021**, *60*, 10736–10744.
- (30) Uzal-Varela, R.; Rodríguez-Rodríguez, A.; Wang, H.; Esteban-Gómez, D.; Brandariz, I.; Gale, E. M.; Caravan, P.; Platas-Iglesias, C. Prediction of Gd(III) Complex Thermodynamic Stability. *Coord. Chem. Rev.* **2022**, *467*, No. 214606.
- (31) Casanova, D.; Alemany, P.; Bofill, J. M.; Alvarez, S. Shape and Symmetry of Heptacoordinate Transition-Metal Complexes: Structural Trends. *Chem.—Eur. J.* **2003**, *9*, 1281–1295.
- (32) Regueiro-Figueroa, M.; Lima, L. M. P.; Blanco, V.; Esteban-Gómez, D.; de Blas, A.; Rodríguez-Blas, T.; Delgado, R.; Platas-Iglesias, C. Reasons behind the Relative Abundances of Heptacoordinate Complexes along the Late First-Row Transition Metal Series. *Inorg. Chem.* **2014**, *53*, 12859–12869.
- (33) Wang, X. F.; Gao, J.; Wang, J.; Zhang, Z. H.; Wang, Y. F.; Chen, L. J.; Sun, W.; Zhang, X. D. Crystal Structures of Seven-Coordinate (NH₄)₂[Mn^{II}(Edta)(H₂O)]·3H₂O, (NH₄)₂[Mn^{II}(Cydta)(H₂O)]·4H₂O and K₂[Mn^{II}(Hdtpa)]·3.5H₂O Complexes. *J. Struct. Chem.* **2008**, *49*, 724–731.
- (34) Maigut, J.; Meier, R.; Zahl, A.; van Eldik, R. Effect of Chelate Dynamics on Water Exchange Reactions of Paramagnetic Aminopolycarboxylate Complexes. *Inorg. Chem.* **2008**, *47*, 5702–5719.
- (35) Maigut, J.; Meier, R.; Zahl, A.; van Eldik, R. Triggering Water Exchange Mechanisms via Chelate Architecture. Shielding of Transition Metal Centers by Aminopolycarboxylate Spectator Ligands. *J. Am. Chem. Soc.* **2008**, *130*, 14556–14569.
- (36) Aime, S.; Anelli, P.; Botta, M.; Brocchetta, M.; Canton, S.; Fedeli, F.; Gianolio, E.; Terreno, E. Relaxometric Evaluation of Novel Manganese(II) Complexes for Application as Contrast Agents in Magnetic Resonance Imaging. *JBIC, J. Biol. Inorg. Chem.* **2002**, *7*, 58–67.
- (37) Gale, E. M.; Zhu, J.; Caravan, P. Direct Measurement of the Mn(II) Hydration State in Metal Complexes and Metalloproteins through ¹⁷O NMR Line Widths. *J. Am. Chem. Soc.* **2013**, *135*, 18600–18608.
- (38) Erstad, D. J.; Ramsay, I. A.; Jordan, V. C.; Sojoodi, M.; Fuchs, B. C.; Tanabe, K. K.; Caravan, P.; Gale, E. M. Tumor Contrast Enhancement and Whole-Body Elimination of the Manganese-Based Magnetic Resonance Imaging Contrast Agent Mn-PyC3A. *Invest. Radiol.* **2019**, *54*, 697–703.
- (39) Wang, J.; Wang, H.; Ramsay, I. A.; Erstad, D. J.; Fuchs, B. C.; Tanabe, K. K.; Caravan, P.; Gale, E. M. Manganese-Based Contrast Agents for Magnetic Resonance Imaging of Liver Tumors: Structure–Activity Relationships and Lead Candidate Evaluation. *J. Med. Chem.* **2018**, *61*, 8811–8824.
- (40) Drahoš, B.; Kotek, J.; Hermann, P.; Lukeš, I.; Tóth, É. Mn²⁺ Complexes with Pyridine-Containing 15-Membered Macrocycles: Thermodynamic, Kinetic, Crystallographic, and ¹H/ ¹⁷O Relaxation Studies. *Inorg. Chem.* **2010**, *49*, 3224–3238.
- (41) Pota, K.; Molnár, E.; Kálmán, F. K.; Freire, D. M.; Tirscó, G.; Green, K. N. Manganese Complex of a Rigidified 15-Membered Macrocyclic: A Comprehensive Study. *Inorg. Chem.* **2020**, *59*, 11366–11376.
- (42) Wen, J.; Geng, Z.; Yin, Y.; Wang, Z. A Mononuclear Mn²⁺ Complex Based on a Novel Tris-(Ethyl Acetate) Pendant-Armed Tetraazamacrocyclic: Effect of Pyridine on Self-Assembly and Weak Interactions. *Inorg. Chem. Commun.* **2012**, *21*, 16–20.
- (43) Uzal-Varela, R.; Valencia, L.; Lalli, D.; Maneiro, M.; Esteban-Gómez, D.; Platas-Iglesias, C.; Botta, M.; Rodríguez-Rodríguez, A. Understanding the Effect of the Electron Spin Relaxation on the Relaxivities of Mn(II) Complexes with Triazacyclononane Derivatives. *Inorg. Chem.* **2021**, *60*, 15055–15068.
- (44) de Sá, A.; Bonnet, C. S.; Geraldine, C. F. G. C.; Tóth, É.; Ferreira, P. M. T.; André, J. P. Thermodynamic Stability and Relaxation Studies of Small, Triaza-Macrocyclic Mn(II) Chelates. *Dalton Trans.* **2013**, *42*, 4522.
- (45) Balogh, E.; He, Z.; Hsieh, W.; Liu, S.; Tóth, É. Dinuclear Complexes Formed with the Triazacyclononane Derivative ENOTA⁴⁻: High-Pressure ¹⁷O NMR Evidence of an Associative Water Exchange on [Mn₂^{II}(ENOTA)(H₂O)₂]. *Inorg. Chem.* **2007**, *46*, 238–250.
- (46) Casanova, D.; Lluell, M.; Alemany, P.; Alvarez, S. The Rich Stereochemistry of Eight-Vertex Polyhedra: A Continuous Shape Measures Study. *Chem.—Eur. J.* **2005**, *11*, 1479–1494.
- (47) Dube, K. S.; Harrop, T. C. Structure and Properties of an Eight-Coordinate Mn(II) Complex That Demonstrates a High Water Relaxivity. *Dalton Trans.* **2011**, *40*, 7496.
- (48) Cieslik, P.; Comba, P.; Dittmar, B.; Ndiaye, D.; Tóth, É.; Velmurugan, G.; Wadepohl, H. Exceptional Manganese(II) Stability and Manganese(II)/Zinc(II) Selectivity with Rigid Polydentate Ligands**. *Angew. Chem., Int. Ed.* **2022**, *61*, No. e202115580.
- (49) Bu, X.-H.; Chen, W.; Mu, L.-J.; Zhang, Z.-H.; Zhang, R.-H.; Clifford, T. Syntheses, Crystal Structures and Properties of New Manganese(II) Complexes with Macrocyclic Polyamine Ligands Bearing Pyridyl Donor Pendant. *Polyhedron* **2000**, *19*, 2095–2100.
- (50) Wang, S.; Westmoreland, T. D. Correlation of Relaxivity with Coordination Number in Six-, Seven-, and Eight-Coordinate Mn(II)

- Complexes of Pendant-Arm Cyclen Derivatives. *Inorg. Chem.* **2009**, *48*, 719–727.
- (51) Rolla, G. A.; Platas-Iglesias, C.; Botta, M.; Tei, L.; Helm, L. ¹H and ¹⁷O NMR Relaxometric and Computational Study on Macrocyclic Mn(II) Complexes. *Inorg. Chem.* **2013**, *52*, 3268–3279.
- (52) Bianchi, A.; Calabi, L.; Giorgi, C.; Losi, P.; Mariani, P.; Palano, D.; Paoli, P.; Rossi, P.; Valtancoli, B. Thermodynamic and Structural Aspects of Manganese(II) Complexes with Polyaminopolycarboxylic Ligands Based upon 1,4,7,10-Tetraazacyclododecane (Cyclen). Crystal Structure of Dimeric [MnL]₂·2CH₃OH Containing the New Ligand 1,4,7,10-Tetraazacyclododecane-1,4-Diacetate. *J. Chem. Soc., Dalton Trans.* **2001**, 917–922.
- (53) Corzilius, B.; Smith, A. A.; Barnes, A. B.; Luchinat, C.; Bertini, I.; Griffin, R. G. High-Field Dynamic Nuclear Polarization with High-Spin Transition Metal Ions. *J. Am. Chem. Soc.* **2011**, *133*, 5648–5651.
- (54) Akhmetzhanov, D.; Ching, H. Y. V.; Denysenkov, V.; Demay-Drouhard, P.; Bertrand, H. C.; Tabares, L. C.; Policar, C.; Prisner, T. F.; Un, S. RIDME Spectroscopy on High-Spin Mn²⁺ Centers. *Phys. Chem. Chem. Phys.* **2016**, *18*, 30857–30866.
- (55) Gao, J.; Ye, K.; He, M.; Xiong, W.-W.; Cao, W.; Lee, Z. Y.; Wang, Y.; Wu, T.; Huo, F.; Liu, X.; Zhang, Q. Tuning Metal–Carboxylate Coordination in Crystalline Metal–Organic Frameworks through Surfactant Media. *J. Solid State Chem.* **2013**, *206*, 27–31.
- (56) Cotton, F. A.; Daniels, L. M.; Murillo, C. A.; Quesada, J. F. Hexaaqua Dipositive Ions of the First Transition Series: New and Accurate Structures; Expected and Unexpected Trends. *Inorg. Chem.* **1993**, *32*, 4861–4867.
- (57) Dale, J. Exploratory Calculations of Medium and Large Rings. *Acta Chem. Scand.* **1973**, *27*, 1115–1129.
- (58) Beattie, J. K. Conformational Analysis of Tris-(Ethylenediamine) Complexes. *Acc. Chem. Res.* **1971**, *4*, 253–259.
- (59) Aime, S.; Botta, M.; Fasano, M.; Marques, M. P. M.; Geraldes, C. F. G. C.; Pubanz, D.; Merbach, A. E. Conformational and Coordination Equilibria on DOTA Complexes of Lanthanide Metal Ions in Aqueous Solution Studied by ¹H-NMR Spectroscopy. *Inorg. Chem.* **1997**, *36*, 2059–2068.
- (60) Parker, D.; Dickins, R. S.; Puschmann, H.; Crossland, C.; Howard, J. A. K. Being Excited by Lanthanide Coordination Complexes: Aqua Species, Chirality, Excited-State Chemistry, and Exchange Dynamics. *Chem. Rev.* **2002**, *102*, 1977.
- (61) Platas-Iglesias, C. The Solution Structure and Dynamics of MRI Probes Based on Lanthanide(III) DOTA as Investigated by DFT and NMR Spectroscopy. *Eur. J. Inorg. Chem.* **2012**, *2012*, 2023–2033.
- (62) Kotek, J.; Rudovský, J.; Hermann, P.; Lukeš, I. Three in One: TSA, TSA', and SA Units in One Crystal Structure of a Yttrium(III) Complex with a Monophosphinated H₄Dota Analogue. *Inorg. Chem.* **2006**, *45*, 3097–3102.
- (63) Urbanovský, P.; Kotek, J.; Císarová, I.; Hermann, P. The Solid-State Structures and Ligand Cavity Evaluation of Lanthanide(III) Complexes of a DOTA Analogue with a (Dibenzylamino)-Methylphosphinate Pendant Arm. *Dalton Trans.* **2020**, *49*, 1555–1569.
- (64) Benetollo, F.; Bombieri, G.; Calabi, L.; Aime, S.; Botta, M. Structural Variations Across the Lanthanide Series of Macrocyclic DOTA Complexes: Insights into the Design of Contrast Agents for Magnetic Resonance Imaging. *Inorg. Chem.* **2003**, *42*, 148–157.
- (65) Nakasuka, N.; Azuma, S.; Katayama, C.; Honda, M.; Tanaka, J.; Tanaka, M. Structure of a Binuclear Manganese(II) Complex: 1,1,1,1,1,2-Hexaaqua-μ-(o-Phenylenediaminetetraacetato-O¹:N,N',O²,O³,O⁴,O⁵)-Dimanganese(II), [Mn₂(C₁₄H₁₂N₂O₈)(H₂O)₆]. *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.* **1985**, *41*, 1176–1179.
- (66) Phukan, B.; Patel, A. B.; Mukherjee, C. A Water-Soluble and Water-Coordinated Mn(II) Complex: Synthesis, Characterization and Phantom MRI Image Study. *Dalton Trans.* **2015**, *44*, 12990–12994.
- (67) Richards, S.; Pedersen, B.; Silvertson, J. V.; Hoard, J. L. Stereochemistry of Ethylenediaminetetraacetato Complexes. I. The Structure of Crystalline Mn₂(HY)₂·10H₂O and the Configuration of the Seven-Coordinate Mn(OH₂)Y⁻² Ion. *Inorg. Chem.* **1964**, *3*, 27–33.
- (68) Aston, K.; Rath, N.; Naik, A.; Slomczynska, U.; Schall, O. F.; Ribley, D. P. Computer-Aided Design (CAD) of Mn(II) Complexes: Superoxide Dismutase Mimetics with Catalytic Activity Exceeding the Native Enzyme. *Inorg. Chem.* **2001**, *40*, 1779–1789.
- (69) Farrugia, L. J. WinGX and ORTEP for Windows: An Update. *J. Appl. Crystallogr.* **2012**, *45*, 849–854.
- (70) Forgács, A.; Giovenzana, G. B.; Botta, M.; Brücher, E.; Tóth, I.; Baranyai, Z. Influence of Gem-Dimethyl Substitution on the Stability, Kinetics and Relaxometric Properties of PDTA Complexes. *Eur. J. Inorg. Chem.* **2012**, *2012*, 2074–2086.
- (71) Kálmán, F. K.; Tircsó, G. Kinetic Inertness of the Mn²⁺ Complexes Formed with AAZTA and Some Open-Chain EDTA Derivatives. *Inorg. Chem.* **2012**, *51*, 10065–10067.
- (72) Pota, K.; Garda, Z.; Kálmán, F. K.; Barriada, J. L.; Esteban-Gómez, D.; Platas-Iglesias, C.; Tóth, I.; Brücher, E.; Tircsó, G. Taking the next Step toward Inert Mn²⁺ Complexes of Open-Chain Ligands: The Case of the Rigid PhDTA Ligand. *New J. Chem.* **2018**, *42*, 8001–8011.
- (73) Porcar-Tost, O.; Pallier, A.; Esteban-Gómez, D.; Illa, O.; Platas-Iglesias, C.; Tóth, E.; Ortuño, R. M. Stability, Relaxometric and Computational Studies on Mn²⁺ Complexes with Ligands Containing a Cyclobutane Scaffold. *Dalton Trans.* **2021**, *50*, 1076–1085.
- (74) Uzal-Varela, R.; Lalli, D.; Brandariz, I.; Rodríguez-Rodríguez, A.; Platas-Iglesias, C.; Botta, M.; Esteban-Gómez, D. Rigid Versions of PDTA⁴⁻ Incorporating a 1,3-Diaminocyclobutyl Spacer for Mn²⁺ Complexation: Stability, Water Exchange Dynamics and Relaxivity. *Dalton Trans.* **2021**, *50*, 16290–16303.
- (75) Negri, R.; Baranyai, Z.; Tei, L.; Giovenzana, G. B.; Platas-Iglesias, C.; Bényei, A. C.; Bodnár, J.; Vágner, A.; Botta, M. Lower Denticity Leading to Higher Stability: Structural and Solution Studies of Ln(III)–OBETA Complexes. *Inorg. Chem.* **2014**, *53*, 12499–12511.
- (76) Laine, S.; Bonnet, C. S.; Kálmán, F. K.; Garda, Z.; Pallier, A.; Caille, F.; Suzenet, F.; Tircsó, G.; Tóth, E. Mn²⁺ Complexes of Open-Chain Ligands with a Pyridine Backbone: Less Donor Atoms Lead to Higher Kinetic Inertness. *New J. Chem.* **2018**, *42*, 8012–8020.
- (77) Anderegg, G. Critical Survey of Stability Constants of NTA Complexes. *Pure Appl. Chem.* **1982**, *54*, 2693–2758.
- (78) Uzal-Varela, R.; Rodríguez-Rodríguez, A.; Martínez-Calvo, M.; Carniato, F.; Lalli, D.; Esteban-Gómez, D.; Brandariz, I.; Pérez-Lourido, P.; Botta, M.; Platas-Iglesias, C. Mn²⁺ Complexes Containing Sulfonamide Groups with PH-Responsive Relaxivity. *Inorg. Chem.* **2020**, *59*, 14306–14317.
- (79) Sawada, K.; Duan, W.; Ono, M.; Satoh, K. Stability and Structure of Nitrilo(Acetate–Methylphosphonate) Complexes of the Alkaline-Earth and Divalent Transition Metal Ions in Aqueous Solution. *J. Chem. Soc., Dalton Trans.* **2000**, 919–924.
- (80) Garda, Z.; Forgács, A.; Do, Q. N.; Kálmán, F. K.; Timári, S.; Baranyai, Z.; Tei, L.; Tóth, I.; Kovács, Z.; Tircsó, G. Physico-Chemical Properties of Mn^{II} Complexes Formed with Cis- and Trans-DO2A: Thermodynamic, Electrochemical and Kinetic Studies. *J. Inorg. Biochem.* **2016**, *163*, 206–213.
- (81) Garda, Z.; Molnár, E.; Kálmán, F. K.; Botár, R.; Nagy, V.; Baranyai, Z.; Brücher, E.; Kovács, Z.; Tóth, I.; Tircsó, G. Effect of the Nature of Donor Atoms on the Thermodynamic, Kinetic and Relaxation Properties of Mn(II) Complexes Formed With Some Trisubstituted 12-Membered Macrocyclic Ligands. *Front. Chem.* **2018**, *6*, No. 232.
- (82) Bárta, J.; Hermann, P.; Kotek, J. Coordination Behavior of 1,4-Disubstituted Cyclen Endowed with Phosphonate, Phosphonate Monoethylester, and H-Phosphinate Pendant Arms. *Molecules* **2019**, *24*, No. 3324.
- (83) Molnár, E.; Camus, N.; Patinec, V.; Rolla, G. A.; Botta, M.; Tircsó, G.; Kálmán, F. K.; Fodor, T.; Tripier, R.; Platas-Iglesias, C. Picolinate-Containing Macrocyclic Mn²⁺ Complexes as Potential MRI Contrast Agents. *Inorg. Chem.* **2014**, *53*, 5136–5149.

- (84) Svobodová, I.; Lubal, P.; Plutnar, J.; Havlíčková, J.; Kotek, J.; Hermann, P.; Lukeš, I. Thermodynamic, Kinetic and Solid-State Study of Divalent Metal Complexes of 1,4,8,11-Tetraazacyclotetradecane (Cyclam) Bearing Two *Trans* (1,8-)Methylphosphonic Acid Pendant Arms. *Dalton Trans.* **2006**, 5184–5197.
- (85) Devreux, M.; Henoumont, C.; Dioury, F.; Boutry, S.; Vacher, O.; Elst, L. V.; Port, M.; Muller, R. N.; Sandre, O.; Laurent, S. Mn^{2+} Complexes with Pyclyen-Based Derivatives as Contrast Agents for Magnetic Resonance Imaging: Synthesis and Relaxometry Characterization. *Inorg. Chem.* **2021**, *60*, 3604–3619.
- (86) Garda, Z.; Molnár, E.; Hamon, N.; Barriada, J. L.; Esteban-Gómez, D.; Várad, B.; Nagy, V.; Pota, K.; Kálmán, F. K.; Tóth, I.; Lih, N.; Platas-Iglesias, C.; Tóth, É.; Tripier, R.; Tircsó, G. Complexation of Mn(II) by Rigid Pyclyen Diacetates: Equilibrium, Kinetic, Relaxometric, Density Functional Theory, and Superoxide Dismutase Activity Studies. *Inorg. Chem.* **2021**, *60*, 1133–1148.
- (87) Kálmán, F. K.; Nagy, V.; Uzal-Varela, R.; Pérez-Lourido, P.; Esteban-Gómez, D.; Garda, Z.; Pota, K.; Mezei, R.; Pallier, A.; Tóth, É.; Platas-Iglesias, C.; Tircsó, G. Expanding the Ligand Classes Used for Mn(II) Complexation: Oxa-Aza Macrocycles Make the Difference. *Molecules* **2021**, *26*, No. 1524.
- (88) Drahoš, B.; Pniok, M.; Havlíčková, J.; Kotek, J.; Cisařová, I.; Hermann, P.; Lukeš, I.; Tóth, É. Mn^{2+} Complexes of 1-Oxa-4,7-Diazacyclononane Based Ligands with Acetic, Phosphonic and Phosphinic Acid Pendant Arms: Stability and Relaxation Studies. *Dalton Trans.* **2011**, *40*, 10131.
- (89) Pujales-Paradela, R.; Carniato, F.; Esteban-Gómez, D.; Botta, M.; Platas-Iglesias, C. Controlling Water Exchange Rates in Potential Mn^{2+} -Based MRI Agents Derived from NO_2A^{2-} . *Dalton Trans.* **2019**, *48*, 3962–3972.
- (90) Drahoš, B.; Kubiček, V.; Bonnet, C. S.; Hermann, P.; Lukeš, I.; Tóth, É. Dissociation Kinetics of Mn^{2+} Complexes of NOTA and DOTA. *Dalton Trans.* **2011**, *40*, 1945.
- (91) Cabral, M. F.; Delgado, R. 4,7,10,13-Tetrakis(Carboxymethyl)-1-Oxa-4,7,10,13-Tetraazacyclotetradecane and Properties of Its Metal Complexes. *Polyhedron* **1999**, *18*, 3479–3489.
- (92) Tei, L.; Gugliotta, G.; Fekete, M.; Kálmán, F. K.; Botta, M. Mn(II) Complexes of Novel Hexadentate AAZTA-like Chelators: A Solution Thermodynamics and Relaxometric Study. *Dalton Trans.* **2011**, *40*, 2025.
- (93) Chaves, S.; Delgado, R.; Da Silva, J. J. R. F. The Stability of the Metal Complexes of Cyclic Tetra-Aza Tetra-Acetic Acids. *Talanta* **1992**, *39*, 249–254.
- (94) Anderegg, G. Pyridinderivate als Komplexbildner I. Pyridin-carbonsäuren. *Helv. Chim. Acta* **1960**, *43*, 414–424.
- (95) Gale, E. M.; Mukherjee, S.; Liu, C.; Loving, G. S.; Caravan, P. Structure–Redox–Relaxivity Relationships for Redox Responsive Manganese-Based Magnetic Resonance Imaging Probes. *Inorg. Chem.* **2014**, *53*, 10748–10761.
- (96) Forgács, A.; Tei, L.; Baranyai, Z.; Esteban-Gómez, D.; Platas-Iglesias, C.; Botta, M. Optimising the Relaxivities of Mn^{2+} Complexes by Targeting Human Serum Albumin (HSA). *Dalton Trans.* **2017**, *46*, 8494–8504.
- (97) Forgács, A.; Tei, L.; Baranyai, Z.; Tóth, I.; Zékány, L.; Botta, M. A Bisamide Derivative of $[Mn(1,4-DO_2A)]$ - Solution Thermodynamic, Kinetic, and NMR Relaxometric Studies. *Eur. J. Inorg. Chem.* **2016**, *2016*, 1165–1174.
- (98) Molnár, E.; Várad, B.; Garda, Z.; Botár, R.; Kálmán, F. K.; Tóth, É.; Platas-Iglesias, C.; Tóth, I.; Brücher, E.; Tircsó, G. Remarkable Differences and Similarities between the Isomeric Mn(II)-*Cis*- and *Trans*-1,2-Diaminocyclohexane-*N,N,N',N'*-Tetraacetate Complexes. *Inorg. Chim. Acta* **2018**, *472*, 254–263.
- (99) Su, H.; Wu, C.; Zhu, J.; Miao, T.; Wang, D.; Xia, C.; Zhao, X.; Gong, Q.; Song, B.; Ai, H. Rigid Mn(II) Chelate as Efficient MRI Contrast Agent for Vascular Imaging. *Dalton Trans.* **2012**, *41*, 14480.
- (100) Baranyai, Z.; Botta, M.; Fekete, M.; Giovenzana, G. B.; Negri, R.; Tei, L.; Platas-Iglesias, C. Lower Ligand Denticity Leading to Improved Thermodynamic and Kinetic Stability of the Gd^{3+} Complex: The Strange Case of OBETA. *Chem.—Eur. J.* **2012**, *18*, 7680–7685.
- (101) Riley, D. P.; Henke, S. L.; Lennon, P. J.; Weiss, R. H.; Neumann, W. L.; Rivers, W. J.; Aston, K. W.; Sample, K. R.; Rahman, H.; Ling, C.-S.; Shieh, J.-J.; Busch, D. H. Witold Szulbinski. Synthesis, Characterization, and Stability of Manganese(II) C-Substituted 1,4,7,10,13-Pentaazacyclotetradecane Complexes Exhibiting Superoxide Dismutase Activity. *Inorg. Chem.* **1996**, *35*, 5213–5231.
- (102) Reilly, C. N.; Good, B. W.; Allendoerfer, R. D. Separation of Contact and Dipolar Lanthanide Induced Nuclear Magnetic Resonance Shifts: Evaluation and Application of Some Structure Independent Methods. *Anal. Chem.* **1976**, *48*, 1446–1458.
- (103) Ogino, H. The Stability Constants of Ethylenediaminetetraacetato, Trimethylenediaminetetraacetato and Propylenediaminetetraacetato Complexes of Some Divalent Metal Ions. *Bull. Chem. Soc. Jpn.* **1965**, *38*, 771–777.
- (104) Takács, A.; Napolitano, R.; Purgel, M.; Bényei, A. C.; Zékány, L.; Brücher, E.; Tóth, I.; Baranyai, Z.; Aime, S. Solution Structures, Stabilities, Kinetics, and Dynamics of DO3A and DO3A–Sulphonamide Complexes. *Inorg. Chem.* **2014**, *53*, 2858–2872.
- (105) Csupász, T.; Szűcs, D.; Kálmán, F. K.; Hollóczki, O.; Fekete, A.; Szikra, D.; Tóth, É.; Tóth, I.; Tircsó, G. A New Oxygen Containing Pyclyen-Type Ligand as a Manganese(II) Binder for MRI and ^{52}Mn PET Applications: Equilibrium, Kinetic, Relaxometric, Structural and Radiochemical Studies. *Molecules* **2022**, *27*, No. 371.
- (106) Lázár, I.; Király, R.; Takács, Z. Synthesis, Potentiometric and 1H NMR Study of Protonation and Complex Formation of 1,4,7-Triazacyclononane-1,4-Diacetate. *J. Coord. Chem.* **2000**, *51*, 293–304.
- (107) Cabral, M. F.; Costa, J.; Delgado, R.; Da Silva, J. J. R. F.; Vilhena, M. F. Protonation and Metal Complexation Studies on Some Oxa-Diaza Macrocyclic Ligands. *Polyhedron* **1990**, *9*, 2847–2857.
- (108) Anderegg, G.; Arnaud-Neu, F.; Delgado, R.; Felcman, J.; Popov, K. Critical Evaluation of Stability Constants of Metal Complexes of Complexones for Biomedical and Environmental Applications. *Pure Appl. Chem.* **2005**, *77*, 1445–1495.
- (109) Liu, Y.-C.; Ma, S.-L.; Guo, Q.-L.; Zhang, J.; Xu, M.-Q.; Zhu, W.-X. Syntheses, Crystal Structures and Properties of Two Mn(II) Complexes of DTPA-Bisamide Derivative. *Inorg. Chem. Commun.* **2005**, *8*, 574–577.
- (110) Laus, S.; Ruloff, R.; Tóth, É.; Merbach, A. E. Gd^{III} Complexes with Fast Water Exchange and High Thermodynamic Stability: Potential Building Blocks for High-Relaxivity MRI Contrast Agents. *Chem.—Eur. J.* **2003**, *9*, 3555–3566.
- (111) Chatterton, N.; Gateau, C.; Mazzanti, M.; Pécaut, J.; Borel, A.; Helm, L.; Merbach, A. The Effect of Pyridinecarboxylate Chelating Groups on the Stability and Electronic Relaxation of Gadolinium Complexes. *Dalton Trans.* **2005**, 1129–1135.
- (112) Hancock, R. D.; Martell, A. E. Ligand Design for Selective Complexation of Metal Ions in Aqueous Solution. *Chem. Rev.* **1989**, *89*, 1875–1914.
- (113) Graves, S. A.; Hernandez, R.; Fonslet, J.; England, C. G.; Valdovinos, H. F.; Ellison, P. A.; Barnhart, T. E.; Elema, D. R.; Theuer, C. P.; Cai, W.; Nickles, R. J.; Severin, G. W. Novel Preparation Methods of ^{52}Mn for ImmunoPET Imaging. *Bioconjugate Chem.* **2015**, *26*, 2118–2124.
- (114) Zhou, I. Y.; Ramsay, I. A.; Ay, I.; Pantazopoulos, P.; Ratile, N. J.; Wong, A.; Caravan, P.; Gale, E. M. Positron Emission Tomography–Magnetic Resonance Imaging Pharmacokinetics, In Vivo Biodistribution, and Whole-Body Elimination of Mn-PyC3A. *Invest. Radiol.* **2021**, *56*, 261–270.
- (115) Xu, J.; Franklin, S. J.; Whisenhunt, D. W.; Raymond, K. N. Gadolinium complex of tris[(3-hydroxy-1-methyl-2-oxo-1,2-dihydro-4-pyridine-4-carboxamido)ethyl]-amine: A New Class of gadolinium magnetic resonance relaxation agents. *J. Am. Chem. Soc.* **1995**, *117* (27), 7245–7246.