

Thermodynamic Stability and Speciation of Ga(III) and Zr(IV) Complexes with High-Denticity Hydroxamate Chelators

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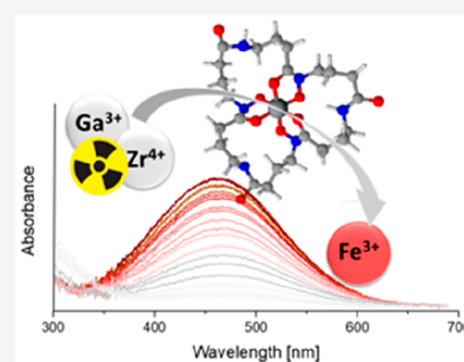


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ABSTRACT: Increasing attention has been recently devoted to $^{89}\text{Zr(IV)}$ and $^{68}\text{Ga(III)}$ radionuclides, due to their favorable decay characteristics for positron emission tomography (PET). In the present paper, a deep investigation is presented on Ga(III) and Zr(IV) complexes with a series of tri- ($\text{H}_3\text{L1}$, $\text{H}_3\text{L3}$, $\text{H}_3\text{L4}$ and desferrioxamine E, DFOE) and tetrahydroxamate ($\text{H}_4\text{L2}$) ligands. Herein, we describe the rational design and synthesis of two cyclic complexing agents ($\text{H}_3\text{L1}$ and $\text{H}_4\text{L2}$) bearing three and four hydroxamate chelating groups, respectively. The ligand structures allow us to take advantage of the macrocyclic effect; the $\text{H}_4\text{L2}$ chelator contains an additional side amino group available for a possible further conjugation with a biomolecule. The thermodynamic stability of Ga(III) and Zr(IV) complexes in solution has been measured using a combination of potentiometric and pH-dependent UV–vis titrations, on the basis of metal–metal competition. The Zr(IV)- $\text{H}_4\text{L2}$ complex is characterized by one of the highest formation constants reported to date for a tetrahydroxamate zirconium chelate ($\log \beta = 45.9$, $pZr = 37.0$), although the complex-stability increase derived from the introduction of the fourth hydroxamate binding unit is lower than that predicted by theoretical calculations. Solution studies on Ga(III) complexes revealed that $\text{H}_3\text{L1}$ and $\text{H}_4\text{L2}$ are stronger chelators in comparison to DFOB. The complex stability obtained with the new ligands is also compared with that previously reported for other hydroxamate ligands. In addition to increasing the library of the thermodynamic stability data of Ga(III) and Zr(IV) complexes, the present work allows new insights into Ga(III) and Zr(IV) coordination chemistry and thermodynamics and broadens the selection of available chelators for $^{68}\text{Ga(III)}$ and $^{89}\text{Zr(IV)}$.



INTRODUCTION

Recent research confirms an increasing interest in the use of gallium and zirconium radioisotopes for medical diagnostic techniques such as PET or single-photon emission computed tomography (SPECT).^{1–8}

The interest in the use of ^{68}Ga ($t_{1/2} = 1.13$ h, $E_{\beta+\text{avg}} = 830$ keV, 89%) for clinical PET comes from the accessibility of its production via an easily portable and long-lived $^{68}\text{Ge}/^{68}\text{Ga}$ generator system.⁹ The favorable decay characteristics of ^{89}Zr ($t_{1/2} = 78.4$ h, $E_{\beta+\text{avg}} = 395.5$ keV, 23%) allow high PET image resolution to be obtained, since the sufficiently long half-life is an optimal match for the pharmacokinetics of most monoclonal antibodies.¹⁰

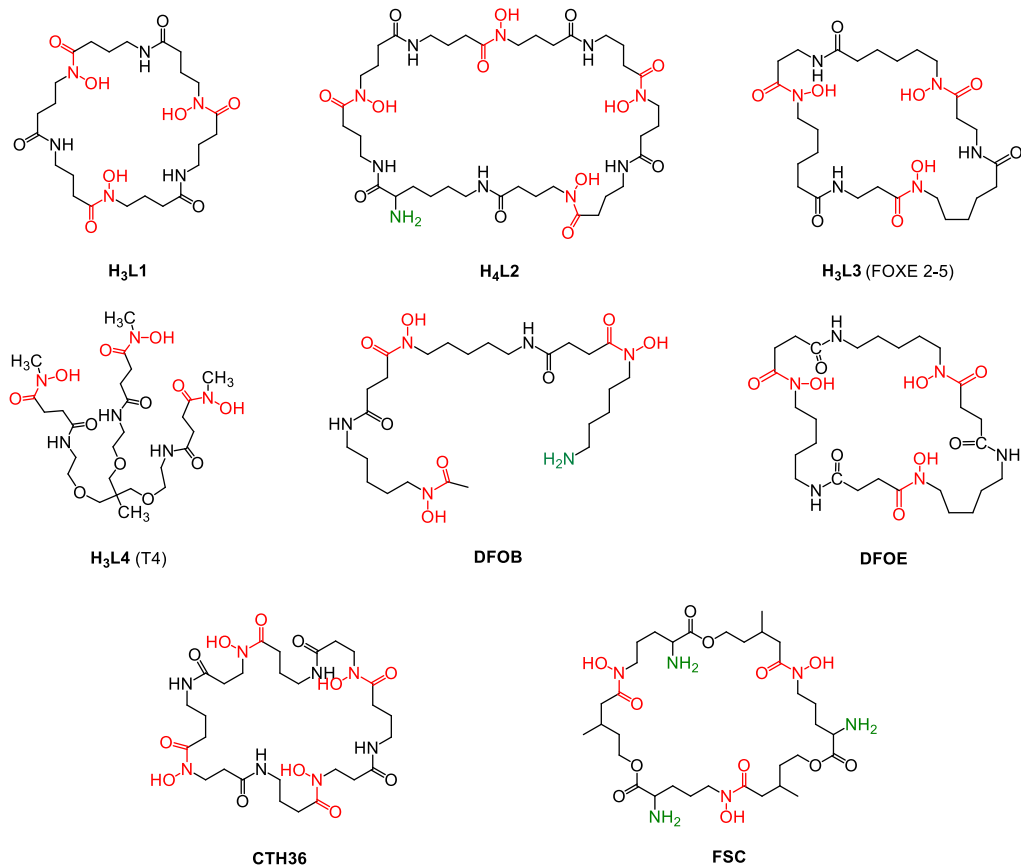
To be applied to molecular imaging, the metal isotope must be converted into a radiolabeled probe that can specifically reach the target of interest *in vivo* and remain there long enough to be detected. Therefore, the metal ion must be bound by an efficient chelator to overcome metal hydrolysis and transchelation, and linked to a biologically active targeting molecule, to be properly directed to the desirable molecular target *in vivo*.

To the best of our knowledge, the most widely used chelator for ^{68}Ga is 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid (DOTA).¹¹ In 2016 the FDA approved a ^{68}Ga -DOTATATE (NETSPOT) kit for the preparation of a ^{68}Ga -DOTATATE injection;¹² clinical trials revealed the superiority of ^{68}Ga -DOTATATE with respect to ^{111}In -pentetreotide in imaging neuroendocrine tumors.¹³ Currently, the most successfully used $^{89}\text{Zr(IV)}$ chelator is DFOB, but some decomposition has been observed over time *in vivo*, and ^{89}Zr slowly accumulates in bones.^{14–16}

Many chelators have been already suggested for ^{68}Ga and ^{89}Zr on the basis of *in vivo* assays.^{14,17–23} While scientists are currently devoting considerable efforts toward the design of

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Scheme 1. Structures of the Ligands Investigated and Discussed in This Paper



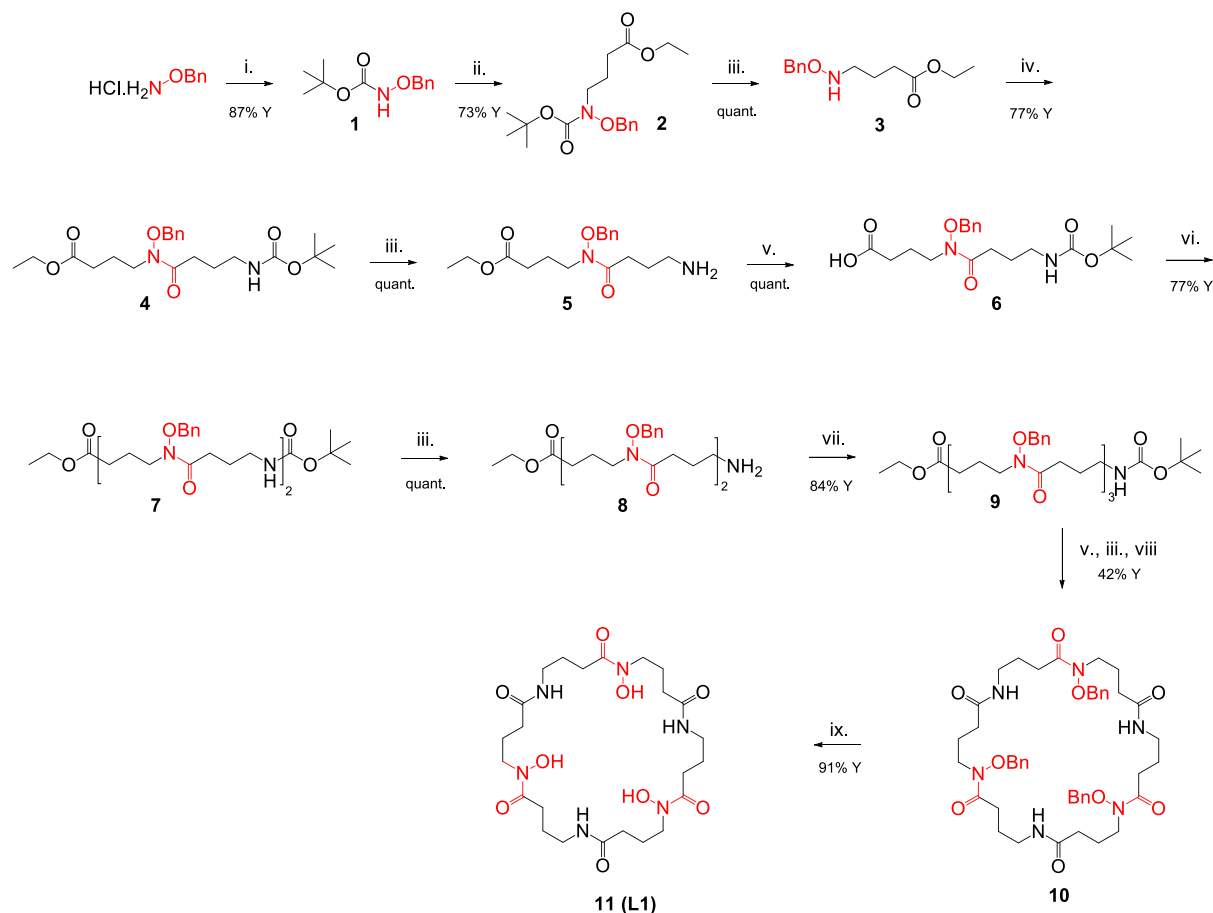
more efficient ^{89}Zr and ^{68}Ga chelators by increasing the *in vivo* stability of the corresponding complexes, the solution equilibrium chemistry, especially of Zr(IV) systems, has still rarely been investigated.^{2,24–27} Solution studies on the coordination chemistry and formation constants of Zr(IV) complexes are not trivial for several reasons: an extremely high thermodynamic stability (requiring application of indirect competition methods with the use of other strong ligands with known stability constants), strong hydrolysis (occurring over almost the entire pH range), and the lack of spectral activity of the complexes. On the other hand, knowledge of the speciation of such complexes, especially at physiological pH, could provide information concerning the actual chemical form of the complex in biological media, and this can contribute to both a better understanding of the *in vivo* speciation and an explanation of the differences in the biological activity.

Our laboratory has recently reported the thermodynamic properties of Zr(IV)-DFOB complexes, suggesting the formation of three mononuclear complexes, i.e. $[\text{ZrHL}]^{2+}$, $[\text{ZrL}]^+$, and $[\text{ZrLH}_2]$, over the pH range 1–11.²⁴ The stability constants and pZr value determined for the Zr(IV)-DFOB system place DFOB among the best Zr(IV) chelators, although the formation of six-coordinate unsaturated complexes (i.e., with the coordination sphere of Zr(IV) completed by two water molecules²⁸ or hydroxide ligands²⁹) and the susceptibility of coordinated water molecules to deprotonation were suggested to be the reason for the *in vivo* lability of ^{89}Zr (IV)-DFOB complexes. The thermodynamic stability of Zr(IV)-DFOB complexes is in line with *in vivo* research^{15,16,30,31} and also with Holland's recent DFT calcu-

lations,³² indicating that our experimental approach was appropriate.

By capitalizing upon our earlier works on siderophore mimics,^{33–36} in this work we have designed, synthesized, and fully characterized tri- and tetrahydroxamic **H₃L1** and **H₄L2** chelators (Scheme 1), analogues of **DFOE**,³⁷ and their Ga(III) and Zr(IV) complexes. In addition, the physicochemical properties of Zr(IV) complexes with three further trihydroxamic ligands, i.e. **H₃L3** (FOX E 2-5),³⁸ **H₃L4** (T4),³⁵ and **DFOE** (Scheme 1), were also investigated for the sake of comparison. The ligands employed here were selected to investigate the influence of some structural elements on the physicochemical properties of Zr(IV) complexes: i.e., (i) the number of binding groups required to complete the coordination sphere of the metal cation; (ii) the possible advantage of the macrocyclic effect; (iii) the size of the ligand cavity, which should be large enough to minimize ring strain; (iv) the symmetry of the ligand, which could limit the probability of a complex challenge. Ga(III) (and Fe(III)) binding to all of these ligands was also performed and will be discussed here, as the relation between the ligand structure and the stability of the complexes has been much more studied and is better understood for trivalent metal ions. Moreover, Fe(III) complexes are used as a tool to determine the thermodynamic stability of Ga(III) and Zr(IV) analogues.

According to DFT studies, to minimize the ring strain, the cyclic tetrahydroxamic ligand should consist of a minimum of 36 atoms, which gives at least 7 chemical groups or 8 bonds.^{39,40} The tetrahydroxamic ligand **H₄L2**, designed to completely saturate the oxophilic coordination sphere of Zr(IV), meets this criterion, possessing at least 9 bonds

Scheme 2. Synthetic Pathway of the Ligand H₃L1^a

^aReagents and conditions: (i) (Boc)₂O, K₂CO₃, H₂O/dioxane; (ii) ethyl 4-bromobutyrate, NaH, DMF; (iii) TFA; (iv) Boc- γ -aminobutyric acid, HATU, DIPEA, DMF; (v) LiOH, H₂O/dioxane; (vi) **5**, HATU, DIPEA, DMF; (vii) **6**, HATU, DIPEA, DMF; (viii) HATU, DIPEA, DMF; (ix) H₂, Pd/C, MeOH.

between binding groups. Of note, **H₄L2** has been designed with the aim of introducing in the chelator a primary amine group, useful to improve the solubility of the ligand but, primarily, to allow the easy conjugation of the chelator to a targeting molecule for future *in vivo* studies. Trihydroxamic **H₃L1** is a symmetrical, macrocyclic ligand, comprising 9 bonds between binding groups; it allows the determination and direct comparison of the influence of the fourth binding group of **H₄L2** on Zr(IV) stability. This ligand also reveals the effect of the shortening of the chain in comparison to **DFOE**, **H₃L3**, and **DFOB**, all with 10 bonds (Scheme 1). Tripodal **H₃L4**, used earlier as a good mimic of a ferrichrome siderophore, was investigated here in order to compare its Zr(IV) binding capacity to those of other tri- and tetrahydroxamic ligands.³⁵ In **H₃L1**, **H₄L2**, and **H₃L3**³⁸ we have used retrohydroxamic units, with a reversed order with respect to the native hydroxamic moiety.

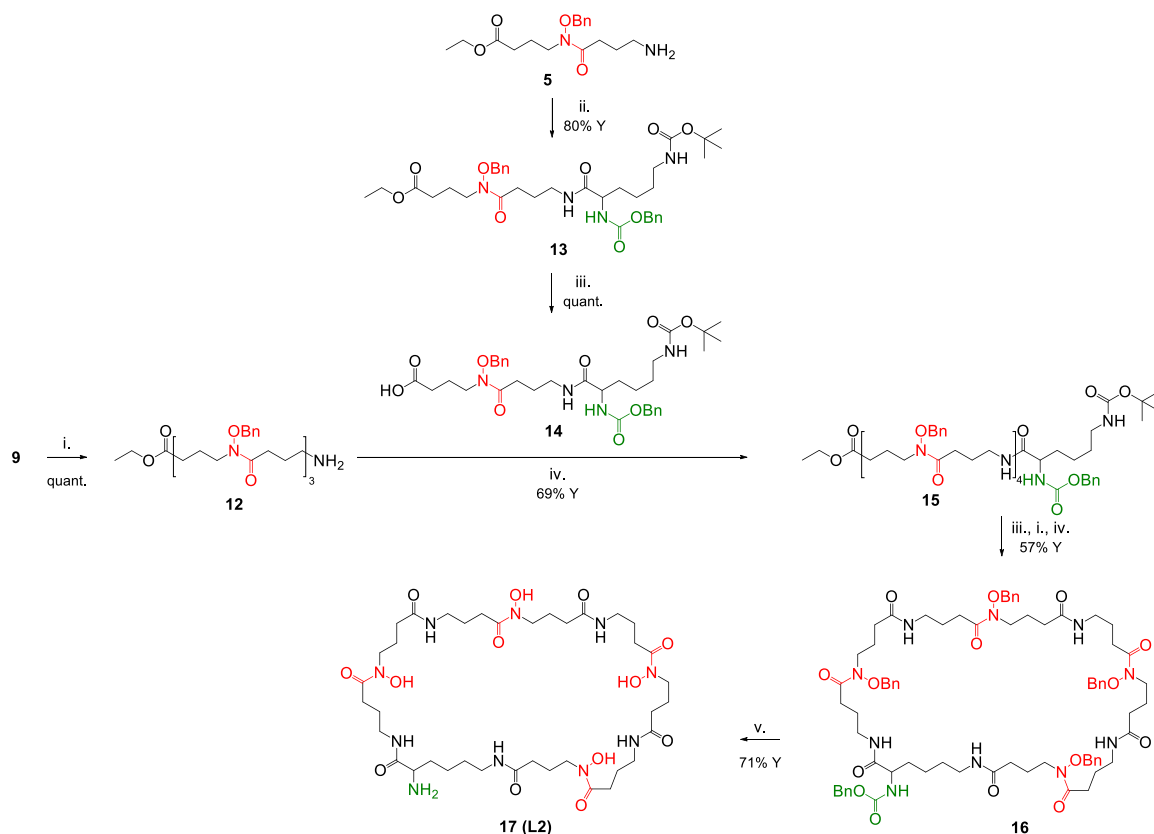
Until now, the stability constants for tetrahydroxamate Zr(IV) complexes have only been estimated from computational calculations³² for linear DFO* ($\log \beta_{[\text{Zr}(\text{DFO}^*)]} = 51.56$) and cyclic CTH36 ($\log \beta_{[\text{Zr}(\text{CTH36})]} = 52.84$). DFO*⁴¹ and CTH36³⁹ possess 10 and 8 bonds between two hydroxamic groups, respectively.

Therefore, an experimental verification of the order of the stability increase between tri- and tetrahydroxamic chelators should be very useful. The same is true for examining the

effects of a number of other variable structural elements, such as the symmetry of the structure and the length of the chain between the hydroxamate binding units. Although the thermodynamic formation constants do not predict *in vivo* stability and kinetic inertness, they are very useful and interesting parameters that may allow for a better design of efficient chelators for Zr(IV) ions.

RESULTS AND DISCUSSION

Design and Synthesis of the Ligands. The synthetic approaches for the preparation of **H₃L1** and **H₄L2** are depicted in Schemes 2 and 3, respectively. The hydroxamate-based monomers protected either at the carboxylic group as an ethyl ester (**5**) or at the amino function with Boc (**6**) were employed as common synthetic precursors of **H₃L1** and **H₄L2**. The building blocks **5** and **6** were synthesized by starting from *O*-benzylhydroxylamine hydrochloride that was first reacted with di-*tert*-butyl dicarbonate to give compound **1** and then alkylated with ethyl 4-bromobutyrate in the presence of NaH to provide compound **2** (Scheme 2). Boc removal with TFA furnished the intermediate **3**, which was coupled with Boc-protected γ -aminobutyric acid using HATU as a coupling reagent. This allowed us to obtain the orthogonally protected **4** as a suitable precursor of both **5** and **6**, which were alternatively isolated after acidic and basic treatments, respectively.

Scheme 3. Synthetic Pathway of the Ligand H_4L2^a 

^aReagents and conditions: (i) TFA; (ii) Z-Lys(Boc)-OH, HATU, DIPEA, DMF; (iii) LiOH, H₂O/dioxane; (iv) HATU, DIPEA, DMF; (v) H₂, Pd/C, MeOH.

For the synthesis of H_3L1 , 5 and 6 were linked together via a standard amide coupling followed by TFA treatment to give 8. The reaction with another unit of 6 gave 9 as the linear protected precursor of H_3L1 . A head–tail HATU-mediated cyclization of 9 was realized under dilute conditions (0.5 mg/mL) after removal of the protection at the C and N terminal positions. A final Pd-catalyzed hydrogenolysis afforded the desired macrocycle H_3L1 in good yields.

For the synthesis of the tetrahydroxamic derivative H_4L2 (Scheme 3), an appropriate hydroxamate-bearing lysine derivative was first prepared as a building block (14). This was obtained by a coupling reaction between 5 and a residue of Z-Lys(Boc)-OH followed by saponification of the ester function. The monomeric unit 14 was coupled to 12, which resulted from TFA-mediated Boc deprotection of 9. The resulting intermediate 15 underwent head–tail HATU-mediated cyclization after removal of the protections at the C and N terminal positions as described above, leading to 16. In addition in this case, the final macrocycle was obtained after Pd-catalyzed removal of the benzyl functions from the hydroxamic groups. These conditions led also to CBz cleavage, leaving the free amino group suitable for future bioconjugation strategies.

H_3L1 , H_4L2 , and their precursors were fully characterized by ¹H and ¹³C NMR (see the Supporting Information). The degree of purity of the final product was evaluated by analytical HPLC assays (see the Supporting Information), showing a purity of higher than 95%.

Thermodynamic Solution Studies. Ligand Protonation Constants. The metal affinity of a ligand depends on its acid–base properties; therefore, the protonation equilibria of H_3L1 and H_4L2 were first investigated. The proton-dissociation processes of the ligands were followed by potentiometric titrations in the pH range 2–11. The hydrolytic stability of the ligands was monitored by a second titration of the same sample with NaOH, following back-acidification of the initially titrated sample. The recorded titration curves were almost exactly superimposed; consequently, the protonation constants calculated from the two consecutive titrations were found to be equal within 0.05 log unit, which indicated that no decomposition occurred.

Data analysis allowed the determination of three protonation constants for H_3L1 and four protonation constants for H_4L2 ; all of them fall in the pH range 8–10 and can be attributed to the hydroxamate groups (Table 1). For each ligand, the protonation constants were assigned by comparing them with the known protonation constants of hydroxamate ligands.^{24,35,37,42} When the changes in temperature, ionic strength, and ligand structures are allowed for, the protonation constants of H_3L1 and H_4L2 are in excellent agreement with the literature values of the cyclic hydroxamate siderophore DFOE (log K_1 = 9.89, log K_2 = 9.42, and log K_3 = 8.65) reported by Anderegg et al.³⁷ The pH-dependent UV–vis titrations of H_3L1 and H_4L2 (Figure S1) revealed the development of a strong band with λ_{max} = 230 nm, when the pH was increased from 7 to 11, which is usually observed for a hydroxamic group deprotonation process.^{24,43} The amino group protonation of H_4L2 was not detectable in the

Table 1. Protonation Constants of Ligands and log β Values of Complexes Formed with Fe(III), Ga(III), and Zr(IV)^a Ions

| assign | H ₃ L1 | | H ₄ L2 | | H ₃ L3 (FOXE 2-S) ^{38,b} | | H ₃ L4 (T4) ^{35,c} | | DFOE ^{36,38,b} | |
|-----------------------|---|---|--|---|--|--|--|---|-------------------------|-----------------|
| | log β | log K | log β | log K | log β | log K | log β | log K | log β | log K |
| LH | 9.89(1) ^d | 9.89 ^d | 10.06(1) ^d | 10.06 ^d | 9.89 | 9.89 | 9.50 | 9.50 | 9.89 | 9.89 |
| LH ₂ | 19.13(1) ^d | 9.24 ^d | 19.65(1) ^d | 9.59 ^d | 19.31 | 19.31 | 18.47 | 8.97 | 19.31 | 9.42 |
| LH ₃ | 27.44(1) ^d | 8.31 ^d | 28.59(1) ^d | 8.94 ^d | 27.96 | 27.96 | 26.73 | 8.26 | 27.96 | 8.65 |
| LH ₄ | | | 36.77(1) ^d | 8.18 ^d | | | | | | |
| [FeH ₂ L] | H ₃ L1 | | H ₄ L2 | | H ₃ L3 (FOXE 2-S) ^{38,b} | | H ₃ L4 (T4) ^{35,c} | | DFOE ^{36,38,b} | |
| | log β | pK _a | log β | pK _a | log β | pK _a | log β | pK _a | log β | pK _a |
| [FeHL] | 31.80(3) ^e | 3.21 ^d 3.1 ^d | 36.96(7) ^e 36.82(6) ^d | 3.00 ^d 3.12 ^d | | | 29.82 | 2.48 | | |
| [FeL] | 28.59(2) ^e 28.7(1) ^d | | | | 32.43 | 32.43 | 27.34 | | 32.43 | |
| [GaH ₂ L] | | | 38.11(3) ^f | 2.20 ^d 2.15(4) ^e | | | | | | |
| [GaHL] | 29.44(7) ^f | 2.65 ^d 2.53(2) ^e | 35.91(7) ^d | 8.13 ^d | | | 27.92(3) ^f | 2.36 ^d 2.24(1) ^e | | |
| [GaL] | 26.79(2) ^d | 9.79 ^d | 27.60(6) ^d | | 29.79 | 29.79 | 25.56(1) ^d | | 29.79 | |
| [ZrHL] | | | 45.9(3) ^f | 2.6 ^d 2.2(1) ^e | | | 36.4(5) ^f | 2.15 ^d 2.2(2) ^e | | |
| [ZrL] | 34.8(2) ^f | 5.48 ^d 5.34(5) ^e | 43.3(1) ^d | | 35.46(5) ^f | 7.03 ^d 7.2(1) ^e | 34.25(5) ^d | 5.43 ^d 5.5(2) ^e | 35.54(9) ^f | |
| [ZrLH ₋₁] | 29.32(8) ^d | | | | 28.31(7) ^d | | 28.8(1) ^d | | | |

^aCharges omitted for clarity. ^bThe protonation constants of the ligands together and the stability constants of Fe(III) and Ga(III) complexes were taken from the literature. ^cThe protonation constants of the ligands and the stability constants of Fe(III) complexes were taken from the literature. ^dDetermined by potentiometric titrations. ^eDetermined by pH-dependent UV–vis titrations. ^fDetermined by metal–metal competition titrations; all measurements performed at 25 °C and $I = 0.1$ M NaClO₄.

experimental pH range. We are aware that the electron-withdrawing character of both the $-\text{NH}_3^+$ and $-\text{NHOH}$ groups affects the acidity of the other group in comparison with that in the related nonsubstituted compound H₃L1. As was previously reported in the example of α - and β -alanine hydroxamic acids, the amino group may be more acidic than the hydroxamic group, or vice versa. According to the literature, the amino group may be more acidic than hydroxamic one, or vice versa.^{44,45} On the basis of the protonation constant of the amino group of DFOB (log $K_{\text{amine}} = 10.97^{24}$), we assume for H₄L2 that it is >11. However, we keep in mind that the deprotonation processes of amino and hydroxamate groups overlap and cannot be distinguished by potentiometry. To elucidate the protonation microequilibria of H₄L2, ¹H NMR titrations should be carried out. However, such a precise analysis is not needed for the determination of the stability of H₄L2-metal complexes and therefore was not performed. The species distribution diagrams of H₃L1 and H₄L2 are presented in Figure S2. The protonation constants of H₃L3 and H₃L4 were reported elsewhere (Table 1).^{35,38}

ESI-MS: Stoichiometry Evaluation. The stoichiometry of the complexes was evaluated by ESI-MS, frequently used as the first step in the determination of metal complex stoichiometry and already previously employed.^{35,46,47} When the fact that ESI-MS is not able to distinguish the ionizable protons in the species is taken into account, this method can be successfully applied to evaluate the metal to ligand stoichiometry directly from the m/z values. For all of the investigated systems, an analysis of the ESI-MS data (collected for various metal to ligand molar ratios) revealed only mononuclear complexes (for

details see Figure S3 and Table S1 in the Supporting Information).

Determination of Complex Stability. To evaluate the thermodynamic stability of Ga(III) and Zr(IV) complexes of the investigated ligands, the binding properties and speciation of Fe(III) complexes have first been determined (all the details are given in the Supporting Information). There are several reasons for this protocol. First, (i) the electron configuration of Ga(III) (d^{10}) and Zr(IV) (d^0) hinders the attainment of spectral information for most of the complexes. Furthermore, (ii) both metal ions are highly acidic and they are readily hydrolyzed over almost the entire pH range. In addition, (iii) the high charge to size ratio of the Zr(IV) ion implies the formation of complexes with an exceptional thermodynamic stability (already at very low pH); as a consequence, the stability constants cannot be directly determined using standard potentiometric titrations. Thus, the thermodynamic stability constants of the Fe(III)-H₃L1 and Fe(III)-H₄L2 systems were first determined (using a combination of potentiometric and pH-dependent UV–vis titrations), followed by Fe(III)–Ga(III) and Fe(III)–Zr(IV) metal–metal competition experiments. Of importance, in order to get accurate results, an experiment where two metal ions compete for a ligand must fulfill two basic requirements: (i) one of the metal chelates should have a strong absorption band in either the visible or ultraviolet region of the spectrum, with an extinction coefficient much different from that of the free metal ion, while the second metal complex should not absorb in the same region of the spectrum; (ii) the equilibrium constant for the competition reaction must not be too small or too large.

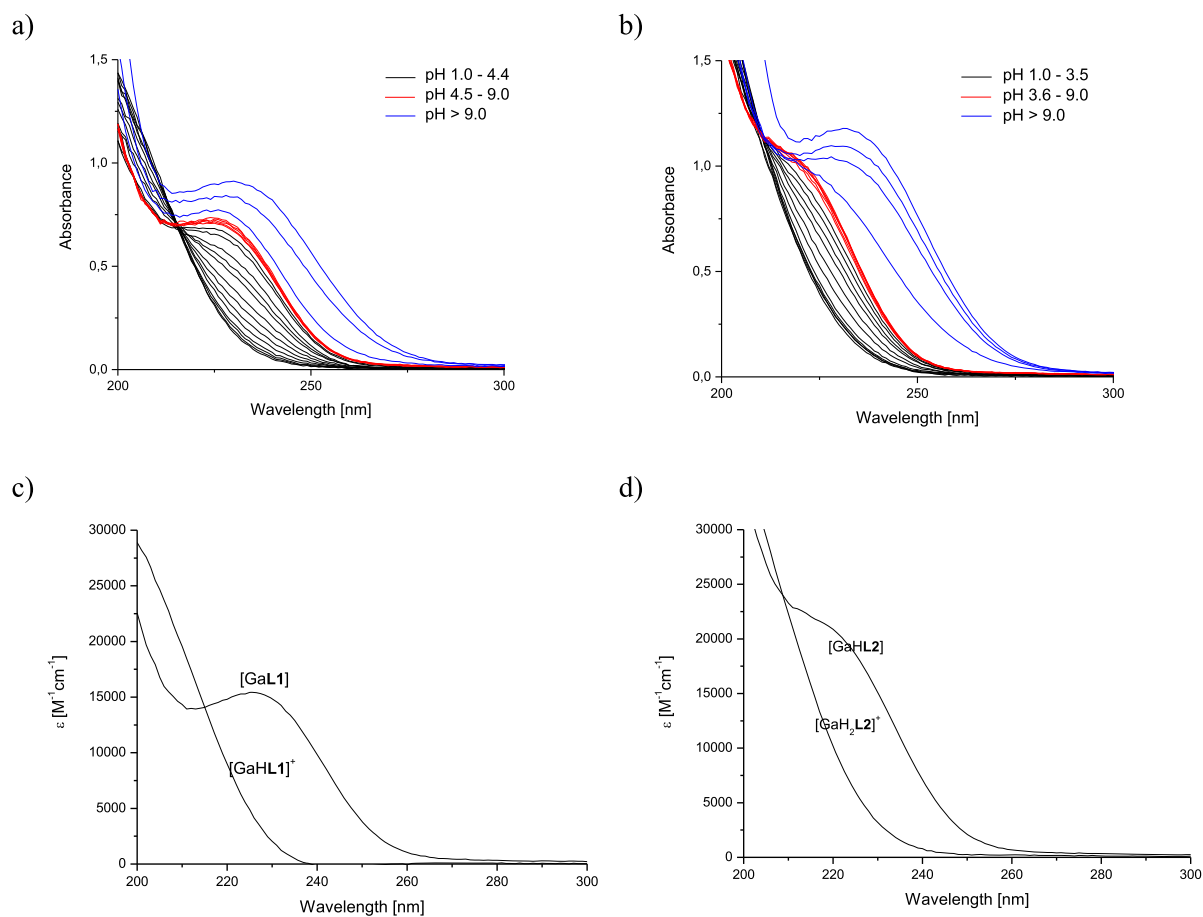


Figure 1. UV spectra of Ga(III)- $\text{H}_3\text{L1}$ (a, c) and Ga(III)- $\text{H}_4\text{L2}$ (b, d) systems at a metal to ligand molar ratio of 1:1 in the pH range 1.0–11.0. Conditions: $c_{\text{L1}} = 0.05 \text{ mM}$, $c_{\text{L2}} = 0.05 \text{ mM}$, 0.1 M NaClO_4 , $T = 25 \text{ }^\circ\text{C}$.

These requirements are fulfilled in the competition experiments between Fe(III) and Ga(III) for both ligands. In the case of Fe(III)–Zr(IV) competition, the difference between the stability constants of Fe(III) and Zr(IV) complexes was too high; thus, an additional competitor ligand—nitrilotriacetic acid (NTA)—was used in the titrations.

NTA is one of the most widely investigated and often used chelating agents.^{48,49} It was selected for the current studies, as both Zr(IV)-NTA and Fe(III)-NTA complexes remain stable until pH 4, even at a metal to ligand molar ratio of 1:1.^{27,48} Moreover, as an additional competing agent, NTA prevents the hydrolysis of the metal ions present in solution and weakens the transchelation observed in the case of $\text{H}_3\text{L1}$ and $\text{H}_4\text{L2}$ Fe(III) complexes titrated directly by Zr(IV) ions. The accuracy of the metal–metal competition titration with NTA was checked on the Zr(IV)-DFOB system, for which experimental data gave $\log \beta_{[\text{ZrHDFOB}]} = 46.1(2)$ (see the Supporting Information), in very good agreement with our recently reported data ($\log \beta_{[\text{ZrHDFOB}]} = 47.7$, allowing for changes in the ionic strength).²⁴ Similar competition procedures are widely used for the evaluation of the stability constants of spectroscopically blind metal complexes.^{25,50}

Ga(III) Complex Formation Equilibria. The evaluation of thermodynamic stability constants of Ga(III) and Zr(IV) complexes with $\text{H}_3\text{L1}$, $\text{H}_4\text{L2}$, and $\text{H}_3\text{L4}$ started from pH-dependent UV–vis spectrophotometric titrations in the pH range 1–11 (Figure 1 and Figure S7, respectively). The spectral changes in the 200–300 nm range, corresponding to

the hydroxamate group protonation state, were monitored.⁵¹ The appearance of a band with a maximum at 225–230 nm with an increase in pH was observed and associated with complex formation.

The pH-dependent UV–vis titration experiments for the Ga(III)- $\text{H}_3\text{L1}$ system revealed an increase in a 230 nm transition band starting from pH 1 up to pH 4.4, with $\text{p}K_{\text{a}} = 2.53(2)$ (Figure 1 and Table 1). For the Ga(III)- $\text{H}_4\text{L2}$ system, the 225 nm band development was observed starting from pH 1 up to pH 3.5, with $\text{p}K_{\text{a}} = 2.15(4)$ (Figure 1 and Table 1), while for Ga(III)- $\text{H}_3\text{L4}$, it continued to increase up to pH 4.6 with $\text{p}K_{\text{a}} = 2.36(1)$ (Figure S7 and Table 1). Similar behavior was observed in the acidic range for Ga(III)-DFOB (and was assigned to two protonation constants of the complex, $\text{p}K_{\text{a}1} = 0.78$ and $\text{p}K_{\text{a}2} = 1.10$ ⁴³) and Th(IV)-DFOB (with $\text{p}K_{\text{a}} = 1.9$ ⁴²) complexes. For the three investigated systems, UV spectra did not reveal any additional changes up to pH 9, indicating that the fully coordinated complex $[\text{GaL}]$, in the case of the Ga(III)- $\text{H}_3\text{L1}$ and Ga(III)- $\text{H}_3\text{L4}$ systems, and the monoprotonated $[\text{GaHL2}]^+$ complex, in the Ga(III)- $\text{H}_4\text{L2}$ system, are the dominant species in solution. When the pH was increased to above 9, an increase in absorbance below 240 nm was observed (Figure 1 and Figure S7). Considering that the spectrophotometric titrations of the free ligands showed the same absorption curves at higher pH, we can suppose that the sharp band at 230–240 nm arises from unbound deprotonated hydroxamate chromophores. Most probably, at higher pH the complex is dissociated, yielding the hydrolyzed gallium species

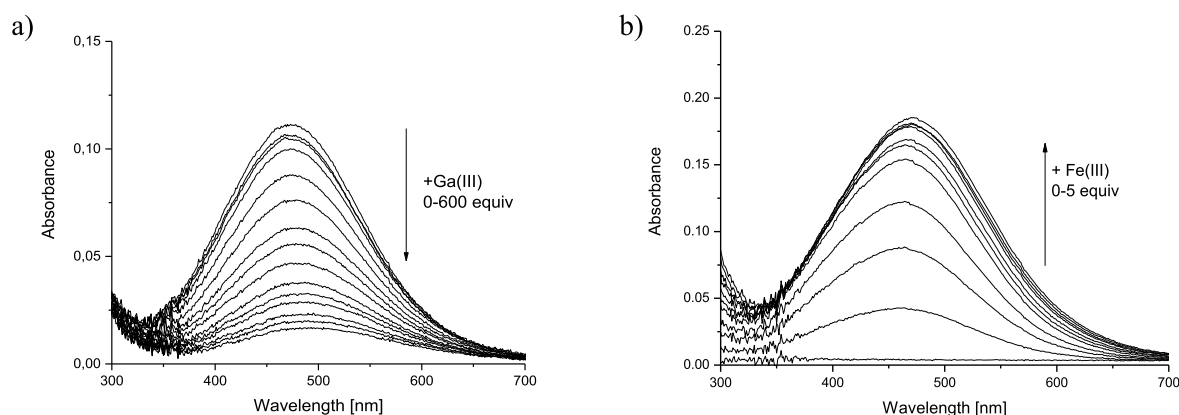


Figure 2. UV-vis metal-metal competition experiment for (a) Fe(III)-H₃L1+Ga(III) ($c_{\text{Fe(III)}} = 0.075$ mM, $c_{\text{L}} = 0.075$ mM) and (b) Ga(III)-H₃L1+Fe(III) ($c_{\text{Fe(III)}} = 0.08$ mM, $c_{\text{L}} = 0.08$ mM) systems at pH 1.5, $I = 0.1$ M (NaClO₄), and $T = 25$ °C.

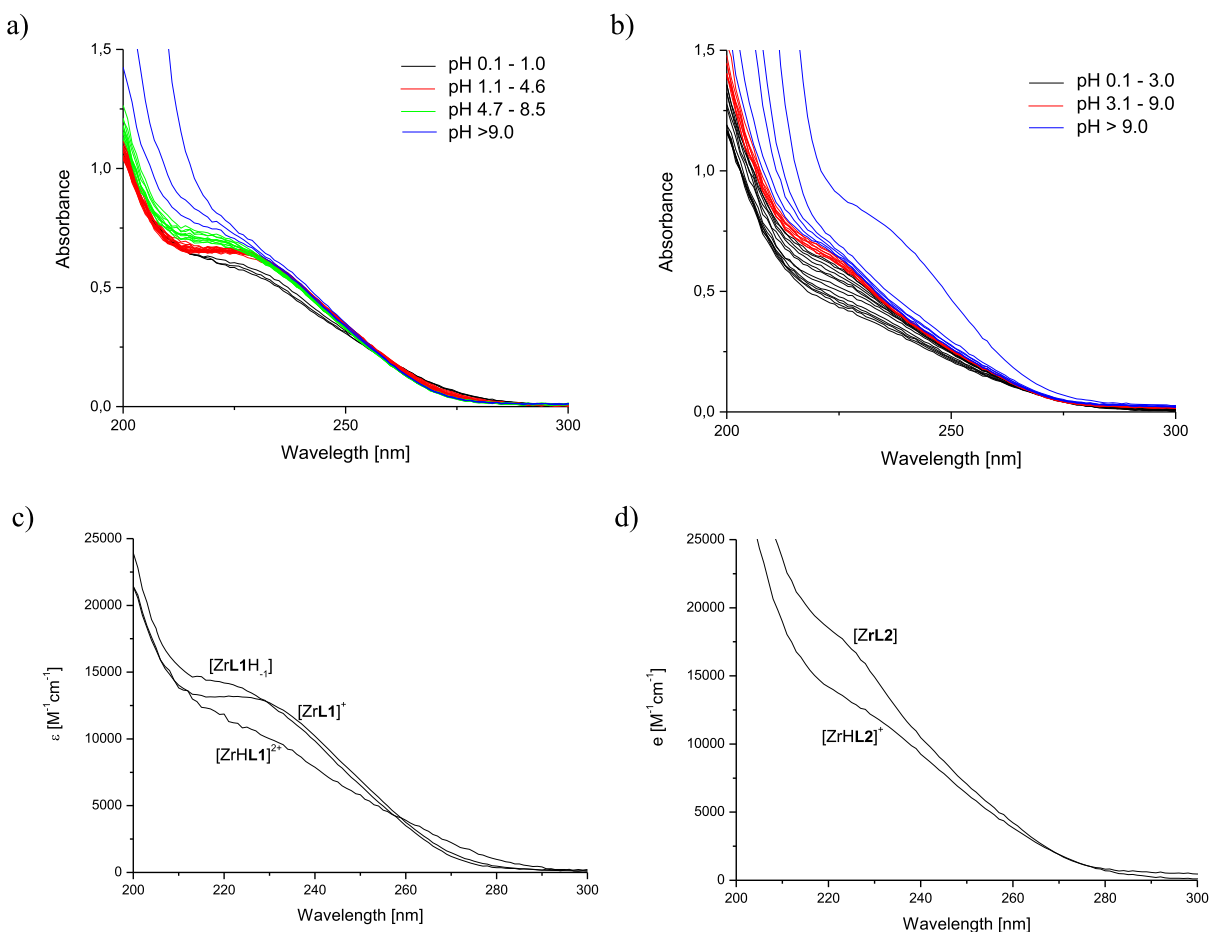


Figure 3. UV spectra of the Zr(IV)-H₃L1 (a, c) and Zr(IV)-H₄L2 (b, d) systems at a metal to ligand molar ratio of 1:1 in the pH range 0.1–11. Conditions: $c_{\text{L1}} = 0.05$ mM, $c_{\text{L2}} = 0.037$ mM, 0.1 M NaClO₄.

[Ga(OH)₄][−]. A similar observation has already been noted for hydroxamate ligands.^{43,52}

Assuming the domination of the [GaHL1]⁺ complex below pH 2, its stability was determined by metal-metal competition titrations, (i) Fe(III)-H₃L1 + Ga(III) and (ii) Ga(III)-H₃L1 + Fe(III), both performed at pH 1.5. This pH was chosen in order to prevent the hydrolysis of the free metal ions and decomposition of the ligand, which is common for hydroxamic acids at a very acidic pH.^{37,51} Upon addition of Ga(III) (up to 600 equiv) to the Fe(III)-H₃L1 solution, the UV-vis band of

[FeHL1]⁺ ($\lambda_{\text{max}} = 470$ nm, Figure 2a) slowly disappeared as a result of the [GaHL1]⁺ complex formation. In the next competition experiment, the appearance of an LMCT transition band ($\lambda_{\text{max}} = 470$ nm) upon addition of Fe(III) to the Ga(III)-H₃L1 solution was observed (Figure 2b). The data refinement using H₃L1 protonation constants (Table 1), Fe(III)-H₃L1 stability constants (Table 1), and stability constants of hydroxocomplexes of both metals (see the Experimental Section) yielded a $\log \beta_{[\text{GaHL1}]^+}$ value of 29.44(7) for Fe(III)-H₃L1 + Ga(III) (Table 1) and of

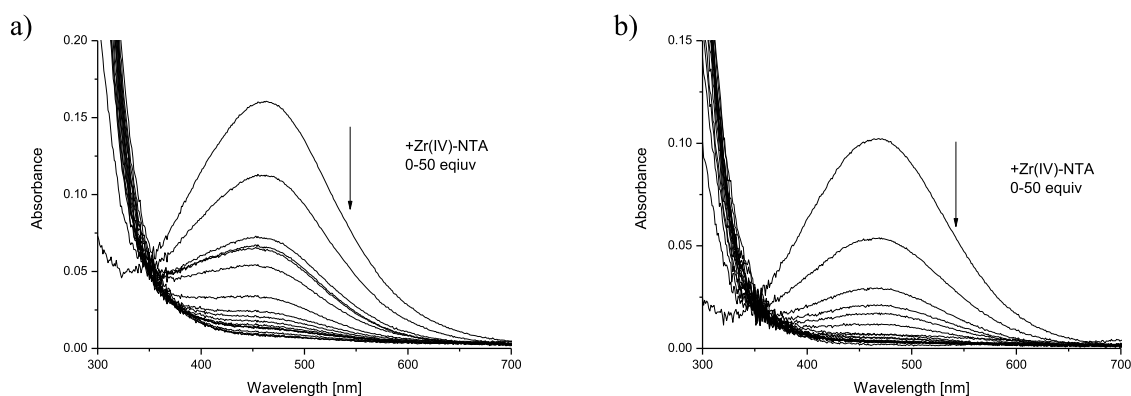


Figure 4. UV-vis spectra of competition titrations of the Fe(III)-H₃L1 + Zr(IV)-NTA ($c_{\text{Fe(III)}} = 0.098$ mM, $c_{\text{L1}} = 0.098$ mM) (a) and Fe(III)-H₄L2 + Zr(IV)-NTA ($c_{\text{Fe(III)}} = 0.055$ mM, $c_{\text{L2}} = 0.055$ mM) (b) systems at pH 1.5 with 0.1 M NaClO₄.

28.3(6) in the case of Ga(III)-H₃L1 + Fe(III) competition titrations.

For the Ga(III)-H₄L2 system, [GaH₂L₂]⁺ was assumed to be the most abundant species at pH 1.5, and its stability was again determined via (i) Fe(III)-H₄L2 + Ga(III) and (ii) Ga(III)-H₄L2 + Fe(III) titrations (Figure S8). The refinement of the titration data yielded $\log \beta_{[\text{GaH}_2\text{L}_2]^+} = 38.11(3)$ and 39.06(4) for Fe(III)-H₄L2 + Ga(III) and Ga(III)-H₄L2 + Fe(III) competition titrations, respectively.

For the Ga(III)-H₃L4 system, [GaHL₄]⁺ was assumed to be the major complex at pH 1.5, and its stability was determined via an Fe(III)-H₃L4 + Ga(III) competition titration (Figure S9). The refinement of the titration data yielded $\log \beta_{[\text{GaHL}_4]^+} = 27.92(3)$.

For the Ga(III)-H₃L1 and Ga(III)-H₄L2 systems, the constants obtained with the two types of titrations are not far from each other but are significantly different. Most likely, the constants measured by means of Ga(III)-H₃L1/H₄L2 + Fe(III) competition experiments are endowed with a greater error, attributable to the overlapping absorption of free iron, present in excess. Therefore, we retained the constants obtained from the Fe(III)-H₃L1/H₄L2 + Ga(III) competition experiments ($\log \beta_{[\text{GaHL}_1]^+} = 29.44(7)$ and $\log \beta_{[\text{GaH}_2\text{L}_2]^+} = 38.11(3)$) as fixed values in the subsequent potentiometric data calculations. The best-fitted speciation model for Ga(III)-H₃L1 and Ga(III)-H₃L4 systems revealed the presence of one additional complex, [GaL], with $\log \beta$ values of 26.79(2) and 25.56(1), respectively ($\text{p}K_{\text{a}1} = 2.65$ for Ga(III)-H₃L1 and 2.36 for Ga(III)-H₃L4; Table 1). For the Ga(III)-H₄L2 system, in addition to [GaH₂L₂]⁺, [GaHL₂] and [GaL₂]⁻ complexes were found with $\log \beta_{[\text{GaHL}_2]} = 35.91(7)$ and $\log \beta_{[\text{GaL}_2]^-} = 27.60(6)$ ($\text{p}K_{\text{a}1} = 2.20$ and $\text{p}K_{\text{a}2} = 8.13$; Table 1). The $\text{p}K_{\text{a}2}$ value of 8.13 is in good agreement with the $\text{p}K_{\text{a}}$ values of the free ligand (Table 1) and could be assigned to the deprotonation of an unbound hydroxamate group.

Zr(IV) Complex Formation Equilibria. The UV-vis titrations of Zr(IV)-H₃L1 equimolar solution over the pH range 0.1–11 (Figure 3a,c) showed a well-defined absorbance band in the 200–300 nm range. The significant changes and the presence of an isosbestic point observed when the pH was increased from 0.1 to 0.9, allowed us to calculate a $\text{p}K_{\text{a}1}$ value of 0.4(2); afterward, the observed 230 nm shoulder remained stable up to pH 4.6. When the pH was increased to 7.0, the development of a 220 nm shoulder with an isosbestic point at 230 nm was observed, characterized by $\text{p}K_{\text{a}2} = 5.34(5)$. From

pH 7.0, the spectra do not reveal any significant changes until pH 9.0, where the hydrolysis probably started. Since information about hydrolysis of the hydroxamate ligands at acidic pH has been widely described in the literature,^{51,52} $\text{p}K_{\text{a}1} = 0.4$ indicates that the three hydroxamate groups are dissociated and therefore most probably bound to the Zr(IV) ion, already at pH <2.0. Assuming the formation of only monomeric complexes, the stability of [ZrL₁]⁺ was determined via UV-vis competition batch experiments, using a Zr(IV)-NTA solution as a competing system for the Fe(III)-H₃L1 complex (Figure 4a). The large LMCT band centered at 470 nm characteristic of dihydroxamate [FeHL₁]⁺ species decreased gradually, and the refinement of the titration data, using the Fe(III)-H₃L1 stability constants (Table 1) together with the Fe(III) and Zr(IV) hydrolysis constants, yielded a $\log \beta_{[\text{ZrL}_1]^+}$ value of 34.8(2).

The UV-vis titration data of the Zr(IV)-H₄L2 system (Figure 3b,d) showed the development of a 230 nm shoulder upon an increase in pH from 0.1 to 3.0 and allowed us to calculate a $\text{p}K_{\text{a}}$ value of 2.2(1). Further, the spectra did not reveal any significant changes over the pH range 3.1–9.0, where the hydrolysis probably started. [ZrHL₂]⁺ was assumed to be the major complex at pH 1.5, and its stability was determined via Fe(III)-H₄L2 + Zr(IV)-NTA (Figure 4b) competition titrations. The refinement of the titration data yielded $\log \beta_{[\text{ZrHL}_2]^+} = 45.9(3)$.

The UV-vis spectra of the Zr(IV)-H₃L3 and Zr(IV)-H₃L4 systems were very similar to the spectra described above (Figure S11); at a pH of around 7 a slight blue shift of the 230 nm band appeared together with the isosbestic point, allowing us to calculate $\text{p}K_{\text{a}} = 7.2(1)$ for Zr(IV)-H₃L3 and 5.5(2) for Zr(IV)-H₃L4 (Table 1). For Zr(IV)-H₃L4, an additional $\text{p}K_{\text{a}} = 2.2(2)$ was calculated, probably corresponding to the formation of the three-hydroxamate complex (Table 1). An evaluation of the competition titration data for Fe(III)-H₃L3 + Zr(IV)-NTA revealed $\log \beta_{[\text{ZrL}_3]^+} = 35.46(5)$, while for Fe(III)-H₃L4 + Zr(IV)-NTA $\log \beta_{[\text{ZrHL}_4]^{2+}} = 36.4(5)$ (Figure S12 and Table 1). Additionally, we have performed the same kind of titration for Fe(III)-DFOE + Zr(IV)-NTA, which gave $\log \beta_{[\text{ZrDFOE}]^+} = 35.54(9)$ (Figure S12 and Table 1). It is worth noting that, during the competition titrations for the H₃L3 ligand and DFOE, a decrease in the 430 nm (typical for three-hydroxamate iron complexes) band was observed, confirming that all three hydroxamate groups are bound already at pH <2.³⁸

Using the constants obtained from competition titrations ($\log \beta_{[\text{ZrL1}]^+} = 34.8(2)$ and $\log \beta_{[\text{ZrHL2}]^+} = 45.9(3)$) as fixed values, the potentiometric data were processed. The best-fitted speciation models revealed the presence of one additional complex for the Zr(IV)-H₃L1 system ($[\text{ZrL1H}_{-1}]$, $\log \beta_{[\text{ZrL1H}_{-1}]} = 29.32(8)$) and one additional complex for the Zr(IV)-H₄L2 system ($[\text{ZrL2}]$, $\log \beta_{[\text{ZrL2}]} = 43.3(1)$) (Table 1).

The $[\text{ZrL1}]^+$ complex dominates the solution from pH 3 up to pH ~ 5 , where its deprotonation to $[\text{ZrL1H}_{-1}]$ occurs, with $\text{p}K_a = 5.48$ (Figure S13a). These results are in line with the spectroscopic data ($\text{p}K_a = 5.34(5)$) and could be ascribed to the dissociation of a water molecule from the coordination sphere of the Zr(IV) ion. For the Zr(IV)-EDTA system, the $\text{p}K_a$ attributed to the deprotonation of a water molecule is 6.2,²⁷ while for Zr(VI)-DFOB it is 6.36.²⁴

The stability constants calculated for the Zr(IV)-H₄L2 system reveal $\text{p}K_a = 2.6$, which is in good agreement with the spectroscopic data ($\text{p}K_a = 2.2(1)$) and could be assigned to the deprotonation of the hydroxamic group and the formation of a fully coordinated tetrahydroxamate complex. The $[\text{ZrL2}]$ complex dominates the solution from pH 3 up to pH ~ 9 , when the dissociation of the complex probably occurs (Figure S13b).

An evaluation of the potentiometric data of Zr(IV)-H₃L3 (with $\log \beta_{[\text{ZrL3}]} = 35.46$ fixed) revealed the presence of an additional complex, $[\text{ZrL3H}_{-1}]$ ($\log \beta_{[\text{ZrL3H}_{-1}]} = 28.35(5)$) (Figure S13c), while for Zr(IV)-H₃L4 (with $\log \beta_{[\text{ZrHL4}]} = 36.4$ fixed), there are two additional complexes, $[\text{ZrL4}]$ ($\log \beta_{[\text{ZrL4}]} = 34.25(5)$) and $[\text{ZrL4H}_{-1}]$ ($\log \beta_{[\text{ZrL4H}_{-1}]} = 28.8(1)$) (Figure S13d). For both systems, the $\text{p}K_a$ values calculated from potentiometric experiments are in excellent agreement with those from the pH-dependent UV-vis titrations (Table 1) and match the data obtained for the whole series of hydroxamate-based ligands. Of importance, the $\text{p}K_a$ value attributed to the deprotonation of a water molecule in the Zr(IV)-H₃L3 system ($\text{p}K_a = 7.2$) is higher than those for Zr(VI)-DFOB ($\text{p}K_a = 6.36$)²⁴ and Zr(IV)-H₃L1 ($\text{p}K_a = 5.48$), suggesting that a longer linker between the binding groups is advantageous. In Zr(IV)-H₃L4, this process is not observed, as all the coordinating positions of the Zr(IV) ion are occupied by four hydroxamate ligands.

Ligand Sequestering Ability. Despite a large variety of PET chelators synthesized and tested in order to provide strong coordination of Ga(III) and Zr(IV) *in vivo*, until now it is has been hard to avoid the release of these metal ions in the body. Here we report two new Ga(III) and Zr(IV) hydroxamate chelators, designed to achieve an efficient sequestering of these metal ions but also to understand how the cyclization and the introduction of an additional hydroxamate group influences the stability of Zr(IV) complexes. Therefore, it is important to evaluate and compare the Ga(III)- and Zr(IV)-sequestering abilities of H₃L1 and H₄L2 with those of other chelators. However, the direct comparison of the stability constants of metal complexes is not straightforward, and other tools taking into account all the physicochemical properties of the ligands, i.e. their denticity, coordination modes, and acid-base properties, should be used.⁵² In order to reliably compare the chelating abilities of H₃L1 and H₄L2 toward Ga(III) and Zr(IV) ions, the pM values were calculated. pFe was originally introduced by Raymond for the comparison of iron-side-rephore systems;⁵³ $\text{pGa} = -\log[\text{Ga(III)}_{\text{free}}]$ and $\text{pZr} =$

$-\log[\text{Zr(IV)}_{\text{free}}]$ were calculated at pH 7.4 with $c_L = 10 \mu\text{M}$ and $c_{\text{Ga(III)}/\text{Zr(IV)}} = 1 \mu\text{M}$ (Table 2).

Table 2. pGa and pZr Values for Various Synthetic and Natural Chelators^a

| ligand | pGa | pZr | chelating groups and ligand geometry |
|--------------------|--------------------|-----------------------|--|
| H ₃ L1 | 22.5 | 32.4 | 3 hydroxamate groups in a cyclic arrangement |
| H ₄ L2 | 22.3 | 37.0 | 4 hydroxamate groups in a linear arrangement |
| H ₃ L3 | 25.4 | 31.5 | 3 hydroxamate groups in a cyclic arrangement |
| H ₃ L4 | 21.9 | 32.6 | 3 hydroxamate groups in a linear arrangement |
| DFOB | 21.6 ⁴³ | 32.2 ²⁴ | 3 hydroxamate groups in a linear arrangement |
| DFOE | 25.2 ³⁸ | 31.0 | 3 hydroxamate groups in a cyclic arrangement |
| DOTA | 20.5 ¹¹ | | 4 macrocyclic amine groups and 4 carboxylate pendant arms |
| NOTA | 27.4 ⁵⁴ | | 3 macrocyclic amine groups and 3 carboxylate pendant arms |
| H ₂ hox | 28.4 ⁵⁵ | | 2 8-hydroxyquinoline groups and 2 amino groups in a linear arrangement |
| PrP9 | 23.1 ⁵⁶ | | 3 macrocyclic amine groups and 3 carboxylate pendant arms |
| HBED | 28.0 ⁵⁷ | | 2 hydroxyaromatic donor groups and 2 carboxylate pendant arms |
| THPN | | 42.7 ² | 4 3-hydroxy-4-pyridinone pendant arms |
| 3,4,3-LI-HOPO | | 44.0 ^{25,58} | 4 1-hydroxy-2-pyridinonates in a linear arrangement |
| DTPA | | 32.3 ^{25,59} | 3 amino groups in linear arrangement and 4 carboxylate pendant arms |

^aValues (re)calculated at pH 7.4 and $c_L = 10 \mu\text{M}$ and $c_{\text{Ga(III)}/\text{Zr(IV)}} = 1 \mu\text{M}$, on the basis of the protonation and stability constants given in original publications. The hydrolysis constants of Ga(III) and Zr(IV) ions were taken from the literature⁶⁰ and are given in the Experimental Section.

The pGa values for Ga(III)-H₃L1 and Ga(III)-H₄L2 systems are on the same order of magnitude as those of the well-known gallium chelators DFOB and PRP9 but are higher than that of the clinically used DOTA (Table 2). On the other hand, H₂hox,⁵⁵ NOTA,⁵⁴ and HBED⁵⁷ present much higher Ga(III) chelating efficacy. The observed effect reflects the differences in the number and type of chelating groups present in the ligands (and therefore the number and type of donor atoms), as well as the ligand dimensions. Of importance, there is only about a 1 order of magnitude increase between pGa values for the linear trihydroxamate ligand DFOB and the cyclic tri- and tetrahydroxamates H₃L1 and H₄L2, respectively. This indicates that the cyclization of the structure only slightly influences the complex stability. It is worth underlining that H₃L1 has shorter spacers between the hydroxamate groups (9 bonds) in comparison to those in DFOB and DFOE (10 bonds). In H₃L3, the spacers have the same length of 10 bonds and the Ga(III) complexes reach the stability of DFOE.³⁸ Additionally, there is almost no difference in complex stability between Ga(III)-H₃L1 and Ga(III)-H₄L2, even though the cavity of H₄L2 is much larger than that of H₃L1, allowing higher flexibility and entropy of the complex structure.

For the Zr(IV) complexes of trihydroxamate H₃L1, H₃L3, and H₃L4 systems, the pZr value is on the same order as those for DFOB²⁴ and DTPA²⁵ chelators. This suggests that ligand cyclization does not provide any increase in complex stability with respect to its linear analogue; for H₃L3 and DFOE one

may even claim a slight decrease in relation to DFOB. A similar conclusion was drawn from the comparison of Zr(IV) complexes of DFOB with fusarinine C (FCS, Scheme 1), where only minor differences in complex stability were observed in *in vivo* studies.⁶¹ An elongation of the chain between hydroxamate binding units from 9 bonds in **H₃L1** to 10 in **H₃L3** is not reflected in the corresponding pZr values. The flexibility of the tripodal **H₃L4** ligand does not produce a higher stability of Zr(IV) complexes. Of importance, the pZr value for the tetrahydroxamate analogue **H₄L2** is >4 units higher than the values calculated for Zr(IV)-DFOB and Zr(IV)-**H₃L1** systems, reflecting the higher affinity of tetrahydroxamate **H₄L2** for Zr(IV), as expected. This feature was already observed in biological studies of other tetrahydroxamate chelators.^{4,39,41} The high thermodynamic stability is certainly the result of the involvement of the fourth hydroxamate coordinating group of the ligand moiety.

The **H₃L1** and **H₄L2** ligands are very good examples to directly compare the stability of Zr(IV) complexes formed with tri- and tetrahydroxamate compounds. For these two ligands, we observe an increase in $\log \beta$ of 8.5 orders of magnitude for the Zr(IV)-**H₄L2** complex with respect to Zr(IV)-**H₃L1** (Table 1). However, this increase is lower than that predicted from the computational calculations performed by Holland³² for tri- and tetrahydroxamate chelators: i.e., DFOB ($\log \beta_{[\text{Zr}(\text{DFOB})]} = 41.20$) versus linear DFO*⁴¹ ($\log \beta_{[\text{Zr}(\text{DFO}^*)]} = 51.56$) and cyclic CTH36³⁹ ($\log \beta_{[\text{Zr}(\text{CTH36})]} = 52.84$). Also, $\log \beta_{[\text{ZrL2}]} = 43.3(1)$ does not reach the values predicted from the above calculations for eight-coordinate Zr(IV)-DFO* and Zr(IV)-CTH36. Of note, the $\log \beta_{[\text{ZrH}(\text{DFOB})\text{OH}]}$ value previously determined by us for DFOB (40.04)²⁴ matches very well the computationally predicted $\log \beta_{[\text{Zr}(\text{DFOB})]}$ value (41.20).³² Still, in the $[\text{ZrH}(\text{DFOB})\text{OH}]^+$ complex, dominating at pH 6.5–10.5, we have suggested the presence of an unbound protonated amino group and a hydrolyzed water molecule bound to Zr(IV). For the cyclic trihydroxamate ligand **H₃L1**, characterized by 9-bond linkers between its hydroxamate units (Scheme 1), $\log \beta_{[\text{ZrL1}]} = 34.8(2)$ is not far from the value estimated for the trihydroxamate cyclic siderophore FSC,^{7,62} $\log \beta_{[\text{Zr}(\text{FSC})]} = 38.92$;³² this difference can be ascribed to the alterations in geometry and dimensions of the ligands. Zirconium is known to form complexes with a complicated geometry of a dodecahedron or square antiprism,^{63,64} and numerous DFT studies have revealed that minor variations in ligand geometry (such as a pendant arm elongation or a modification of the ligand cavity size) could result in significant changes in the stability of Zr(IV) complexes.^{39,65,66} **H₄L2** presented in this work possesses four hydroxamate units and amide units in the linker, but a larger cavity size and a significant asymmetry (coming from one much longer linker and with two amides and amino group) with respect to CTH36 (Scheme 1).^{32,39} These structural alterations are most probably the reason for the lower thermodynamic stability of **H₄L2** complexes, as the coordination sphere might not be uniformly closed around the central ion. Unfortunately, the thermodynamic characterization of Zr(IV)-CTH36 complexes has not yet been reported; thus, the pZr value cannot be quantified. Another cyclic hydroxamic ligand, PPDDFOT₁, that possesses four hydroxamic groups in a symmetrical arrangement and a cavity size even larger than that of **H₄L2** (11 bonds between hydroxamic groups) showed superior stability vs DFOB in EDTA challenging assays.⁶⁷ These results confirm that the Zr(IV) complex stability is strongly

dependent on the ligand geometry and emphasize the demand for a thermodynamic solution study in order to understand this dependence. Other octadentate hydroxy-pyridinone chelators, such as THPN² and 3,4,3-LI-HOPO,²⁵ form the strongest complexes (Table 2). The reason could be not only the type of chelating groups present in the ligands but also the ligand architecture and dimensions.

CONCLUSIONS

In the present work we have developed new chelating agents for the complete saturation of the coordination spheres of Ga(III) and Zr(IV) metals. The trihydroxamic (**H₃L1**) and tetrahydroxamic (**H₄L2**) ligands were successfully synthesized, and the thermodynamic properties of their Ga(III) and Zr(IV) complexes were evaluated. In addition, a series of other synthetic (**H₃L3**, **H₃L4**) and natural (DFOE) compounds was investigated. **H₃L1** proved to be an efficient Ga(III) chelator, but the stability of its Zr(IV) complexes is about 1 order of magnitude lower than that reported for the Zr(IV)-DFOB system. **H₄L2** is the first tetrahydroxamate ligand for which the formation constants and speciation with Zr(IV) were experimentally determined. Of importance, it revealed an enhanced stability of 8.5 orders of magnitude ($\log \beta_{[\text{ZrL2}]} = 43.3$, pZr = 37.0) with respect to Zr(IV)-**H₃L1** ($\log \beta_{[\text{ZrL1}]} = 34.8$, pZr = 32.4), as a consequence of the introduction of a fourth hydroxamate binding unit. However, the stability increase is lower than that predicted by computational calculations for the tetrahydroxamate chelators DFO* ($\log \beta_{[\text{Zr}(\text{DFO}^*)]} = 51.56$) and CTH36 ($\log \beta_{[\text{Zr}(\text{CTH36})]} = 52.84$), and this effect can be ascribed to the structural alterations of the **H₄L2** ligand.

Overall, the determination of the thermodynamic stability of metal complexes coupled with a suitable chelator design will help further developments of optimal chelators for PET imaging applications. However we are aware that there are still a great number of tests to do in order for these ligands to be used as a PET chelators, such as radiolabeling and kinetics studies, biodistribution assays, etc. Current efforts are focused on the design and studies of tetrapodal hydroxamate ligands, to interrogate how their shape and size tune the thermodynamic stability of Zr(IV) complexes.

According to the literature, the ⁸⁹Zr radiolabeling strategies for hydroxamate ligands is usually simple, robust, and relatively rapid. They are performed under mild pH conditions, at room temperature, and take around 1 h for DFOB derivatives on their own without being attached to any targeting vectors and 1–3 h for DFO derivatives attached to targeting vectors, such as trastuzumab.^{14,68,69} Cyclic hydroxamate ligands such as C7⁴⁰ and CTH36³⁹ (both with 8 bonds between hydroxamate groups) have demonstrated excellent complexation abilities at ambient temperature (>99% complexation after 120 min for C7 and >90% of the activity within 5 min reaction time for CTH36, respectively). The slightly smaller ligands C6 and C5, reported by Guerard et al.,⁴⁰ appear to be less suitable for radiolabeling, with higher temperatures being required to obtain high complexation yields. The cyclic ligands reported in this paper possess a larger cavity size, with at least 9 bonds between hydroxamate groups, which allows us to assume that the radiolabeling process will be highly efficient and performed under mild conditions. Preliminary radiolabeling of artificial FOXE siderophores with ⁶⁸Ga, represented here by FOXE 2–5, was achieved after 10 min at room temperature with

moderate yields and high specific activities of ^{68}Ga .³⁸ Further characterization is ongoing.

EXPERIMENTAL SECTION

Synthesis. General Considerations. Unless stated otherwise, all commercially available reagents and solvents were of analytical grade. For the synthesis of **H₃L1** and **H₄L2**, solvents and reagents were purchased from Bachem, BLDpharm, and Fluka. Crude products were purified via flash column chromatography on silica gel (Merck, 230–400 Mesh) or, for compounds **10**, **11**, **16**, and **17**, by semipreparative RP-HPLC using a Waters Prep 600 system equipped with a C18 Jupiter column (250 × 30 mm, 300 Å, 15 μm spherical particle size). Gradients were established each time by considering the analytical HPLC profile of the crude product. The column was perfused at a flow rate of 20 mL/min over 30 min with a binary system of solvent A (H₂O + 0.1% v/v TFA) and solvent B (60% CH₃CN in water + 0.1% v/v TFA). Analytical RP-HPLC analyses were performed on a XBridge C18 column (4.6 × 150 mm, 5 μm particle size) using a flow rate of 0.7 mL/min and a linear gradient of acetonitrile (and 0.1% TFA) in water (and 0.1% TFA) from 0% to 100% over 25 min. The mass spectra were recorded on an ESI-Micromass ZMD 2000 instrument. TLC was performed on precoated plates of silica gel F254 (Merck, Darmstadt, Germany). ¹H NMR analyses were obtained using a Varian spectrometer (400 MHz) and were referenced to residual ¹H signals of the deuterated solvents ($\delta(^1\text{H})$ 7.26 for CDCl₃; $\delta(^1\text{H})$ 2.50 for DMSO). The following abbreviations are used to describe the shape of the peaks: s, singlet; d, doublet; dd, doublet of doublets; t, triplet; m, multiplet.

Synthesis of tert-Butyl(benzyloxy)carbamate (1). To an ice-cooled solution of *O*-benzyloxyethylamine·HCl (4.00 g, 25 mmol) in a 1,4-dioxane/H₂O mixture (60 mL, 1/1 v/v) was added K₂CO₃ (10.37 g, 75 mmol). Boc₂O (8.18 g, 37.5 mmol), previously dissolved in dioxane, was then added dropwise, and the reaction mixture was stirred at room temperature overnight. The solvent was removed under vacuum, and the crude product extracted using ethyl acetate (30 mL) and water (3 × 15 mL). The organic phase was dried over Na₂SO₄, filtered, and evaporated. Compound **1** (4.86 g, 87% yield) was obtained as a colorless oil, which was used without any further purification. NMR data match those reported in the literature (PMID: 11906271). ESI-MS: calcd for C₁₂H₁₈NO₃, 224.28 [M + H]⁺; found, 224.13 [M + H]⁺. T_R = 19.70 min.

Synthesis of Ethyl 4-((Benzyloxy)(tert-butoxycarbonyl)amino)butanoate (2). To a solution of **1** (4.86 g, 21.79 mmol) in DMF (15 mL) was added NaH (60% dispersion in mineral oil, 1.20 g, 23.94 mmol). The mixture was initially stirred at rt for 30 min, and then the reaction mixture was warmed to 60 °C and ethyl 4-bromobutyrate was added dropwise. At the completion of the reaction, the solvent was removed, and the residue was extracted with ethyl acetate and water (3 × 30 mL), dried over Na₂SO₄, and concentrated under vacuum. The product (**2**) was obtained as a yellowish oil (5.36 g, 73% yield). The NMR data correspond to those in the literature (PMID: 28715615). MS (ESI): calcd for C₁₈H₂₈NO₅, 338.20 [M + H]⁺; found, 360.18 [M + Na]⁺, 697.37 [2M + Na]⁺. T_R = 23.51 min.

Synthesis of Ethyl 4-(N-(Benzyloxy)-4-((tert-butoxycarbonyl)amino)butanamido)butanoate (4). Compound **2** (5.36 g, 15.90 mmol) was dissolved in trifluoroacetic acid (TFA, 6 mL), and the mixture was stirred at room temperature for 2 h. The reaction mixture was monitored by MS (ESI) before being concentrated under vacuum. The deprotected amino ester **3** was used without further purification in the next step. To an ice-cold solution of Boc-γ-aminobutyric acid (2.7 g, 13.45 mmol) in DMF (20 mL) were added 1-[[bis(dimethylamino)methylene]-1*H*-1,2,3-triazolo[4,5-*b*]pyridinium 3-oxide hexafluorophosphate (HATU, 5.6 g, 14.75 mmol) and DIPEA (2.6 mL, 14.75 mmol). A portion of **3** (3.5 g, 14.75 mmol) was dissolved in DMF (10 mL), and this solution was added dropwise to the first one. Then the reaction mixture was warmed to room temperature and stirred for 1 h. After removal of the solvent, the residue was dissolved in ethyl acetate and washed with a 5% aqueous solution of citric acid, a 10% aqueous solution of NaHCO₃, and brine.

The crude product was purified by column chromatography using ethyl acetate/petroleum ether (from 1/4 to 1/1 by volume) as an eluent mixture. Compound **4** was obtained as a slightly yellowish oil (4.35 g, 76.6% yield). ESI-MS: calcd for C₂₂H₃₅N₂O₆, 423.53 [M + H]⁺; found, 423.25 [M + H]⁺, 445.23 [M + Na]⁺, 867.47 [2M + Na]⁺. T_R = 21.11 min. ¹H NMR (400 MHz, CDCl₃): δ 7.44–7.32 (m, 5H), 4.80 (s, 2H), 4.11 (qd, *J* = 7.1, 2.9 Hz, 2H), 3.70 (t, *J* = 6.8 Hz, 2H), 3.12 (t, *J* = 6.6 Hz, 2H), 2.42 (t, *J* = 7.2 Hz, 2H), 2.32 (t, *J* = 7.3 Hz, 2H), 1.99–1.91 (m, 2H), 1.80–1.73 (m, 2H), 1.42 (s, 9H), 1.26–1.20 (m, 3H). ¹³C NMR (CDCl₃): δ 172.9, 156.0, 134.3, 129.2, 129.0, 128.7, 79.1, 60.4, 44.6, 40.3, 31.4, 29.6, 28.4, 24.7, 22.3, 14.2.

Synthesis of Ethyl 10,20-Bis(benzyloxy)-2,2-dimethyl-4,9,14,19-tetraoxo-3-oxa-5,10,15,20-tetraazatetracosan-24-oate (7). The Boc-protected derivative **5** (3.40 g, 7.8 mmol) was obtained as previously described for **3**. Compound **6** was synthesized by dissolving the ethyl ester **4** (3.0 g, 7.1 mmol) in a 1,4-dioxane/H₂O mixture in the presence of LiOH (1 M aqueous solution, 12.5 mmol). The mixture was stirred at rt for 20–30 min. Once the reaction was complete, dioxane was evaporated and the crude product was acidified using 1 M HCl to reach pH 6. Then, the aqueous phase was extracted using ethyl acetate. Compound **6** (0.91 g, 2.31 mmol) was used in the next step without further purification. The coupling reaction was conducted as previously described for **4**, and derivative **7** was obtained as a yellowish oil (1.24 g, 77% yield) after column chromatography. ESI-MS: calcd for C₃₇H₅₅N₄O₉, 699.87 [M + H]⁺; found, 699.96 [M + H]⁺. T_R = 21.18 min. ¹H NMR (400 MHz, CDCl₃): δ 7.50–7.31 (m, 10H), 7.04 (bs, 1H), 4.81 (d, *J* = 5.3 Hz, 4H), 4.12 (q, *J* = 7.1 Hz, 2H), 3.71–3.69 (m, 4H), 3.26 (dd, *J* = 11.9, 6.2 Hz, 2H), 3.17–3.09 (m, 2H), 2.53–2.39 (m, 4H), 2.33 (t, *J* = 7.3 Hz, 2H), 2.20 (t, *J* = 6.8 Hz, 2H), 1.98–1.92 (m, 4H), 1.84–1.73 (m, 4H), 1.42 (s, 9H), 1.24 (t, *J* = 7.1 Hz, 3H). ¹³C NMR (CDCl₃): δ 174.7, 173.3, 173.0, 134.2, 129.3, 129.1, 129.1, 128.8, 60.5, 44.7, 44.3, 40.0, 39.6, 33.1, 31.4, 30.0, 29.4, 28.5, 24.8, 23.9, 23.2, 22.3, 14.3.

Synthesis of Ethyl 10,20,30-Tris(benzyloxy)-2,2-dimethyl-4,9,14,19,24,29-hexaaxo-3-oxa-5,10,15,20,25,30-hexaazatetracontan-34-oate (9). Compound **9** was synthesized under the same coupling conditions used for **4** by starting from the acid derivative **6** (0.91 g, 2.31 mmol) and the amino derivative **8** (1.81 g, 2.54 mmol). The desired product was obtained as a colorless oil (1.89 g, 84% yield) after column chromatography. ESI-MS: calcd for C₅₂H₇₅N₆O₁₂, 976.20 [M + H]⁺; found, 975.94 [M + H]⁺. T_R = 17.83 min. ¹H NMR (400 MHz, CDCl₃): δ 7.39–7.35 (m, 15H), 5.05 (bs, 3H), 4.83–4.77 (m, 6H), 4.11 (q, *J* = 7.1 Hz, 2H), 3.72–3.67 (m, 6H), 3.34–3.19 (m, 4H), 3.16–3.11 (m, 2H), 2.49–2.45 (m, 6H), 2.32 (t, *J* = 7.3 Hz, 2H), 2.25–2.12 (m, 4H), 2.02–1.89 (m, 6H), 1.87–1.72 (m, 6H), 1.42 (s, 9H), 1.24 (t, *J* = 7.1 Hz, 3H). ¹³C NMR (CDCl₃): δ 175.3, 174.1, 172.5, 157.3, 135.4, 128.9, 128.3, 128.2, 80.7, 73.8, 61.2, 48.6, 41.2, 40.4, 35.2, 31.8, 31.5, 28.4, 24.1, 21.7, 18.5, 14.7.

Synthesis of 1,11,21-Tris(benzyloxy)-1,6,11,16,21,26-hexaazacyclotriacontane-2,7,12,17,22,27-hexaone (10). Compound **9** was Boc-protected as described for **3**. Then, the ethyl group was hydrolyzed by LiOH as for **6**. To a dilute solution of the fully deprotected trimer (1.92 g, 2.0 mmol) in DMF (100 mL) were added HATU (0.84 g, 2.2 mmol) and DIPEA (0.38 mL, 2.2 mmol) dropwise at 0 °C. The reaction mixture was stirred for 3 h. Then, the solvent was removed, and the residue was extracted with ethyl acetate and an aqueous solution of citric acid (10%), a solution of NaHCO₃ (5%), and brine. The crude product was purified via semipreparative HPLC, giving the desired product as a colorless oil (0.70 g, 42% yield). ESI-MS: calcd for C₄₅H₆₁N₆O₉, 830.02 [M + H]⁺; found, 829.90 [M + H]⁺. T_R = 21.47 min. ¹H NMR (400 MHz, CDCl₃): δ 7.39–7.31 (m, 18H), 4.77 (s, 6H), 3.69–3.67 (m, 6H), 3.25–3.23 (m, 6H), 2.46 (t, *J* = 6.7 Hz, 6H), 2.19 (t, *J* = 7.0 Hz, 6H), 1.97–1.91 (m, 6H), 1.85–1.70 (m, 6H). ¹³C NMR (CDCl₃): δ 174.9, 173.8, 133.8, 129.3, 128.8, 44.2, 39.4, 33.0, 29.5, 23.7, 23.2.

Synthesis of 1,11,21-Trihydroxy-1,6,11,16,21,26-hexaazacyclotriacontane-2,7,12,17,22,27-hexaone (11, H₃L1). To a solution of the benzyl-protected derivative **10** (0.70 g, 0.84 mmol) in MeOH (30 mL) was added glacial acetic acid (1 mL). The mixture was treated with a catalytic amount (0.084 mmol) of palladium on activated

charcoal (10% Pd basis) under a hydrogen atmosphere. After 24 h, the reaction mixture was filtered through Celite, concentrated under reduced pressure, diluted with water, and alkalized with saturated sodium bicarbonate. The aqueous phase was extracted with ethyl acetate (4 × 10 mL), and the combined organic layers were dried over Na₂SO₄, filtered, and concentrated under vacuum. Compound **11** was obtained as a light yellow oil after preparative HPLC purification (0.43 g, 91% yield). ESI-MS: calcd for C₂₄H₄₃N₆O₉, 559.64 [M + H]⁺; found, 559.79 [M + H]⁺. T_R = 15.79 min. ¹H NMR (400 MHz, CDCl₃): δ 9.59 (bs, 3H), 7.83–7.81 (m, 3H), 3.53–3.38 (m, 6H), 3.15–2.93 (m, 6H), 2.40–2.24 (m, 5H), 2.20–2.16 (m, 2H), 2.02 (t, J = 7.1 Hz, 5H), 1.76–1.70 (m, 6H), 1.64–1.50 (m, 6H). ¹³C NMR (CDCl₃): δ 174.5, 173.0, 172.2, 169.5, 158.9, 158.5, 129.9, 50.5, 47.3, 47.1, 38.7, 38.6, 33.0, 31.5, 31.2, 30.3, 29.7, 24.9, 24.8, 23.0, 22.3, 19.7. HR-ESI-MS *m/z* 559.30894; calcd for C₂₄H₄₃N₆O₉ ([M + H]⁺) 559.30860. Anal. Calcd for C₂₄H₄₂N₆O₉: C, 51.6; H, 7.6; N, 15.0. Found: C, 51.4; H, 7.5; N, 14.9.

Synthesis of Ethyl 17-(Benzyloxy)-10-(((benzyloxy)carbonyl)amino)-2,2-dimethyl-4,11,16-trioxo-3-oxa-5,12,17-triazahenicosan-21-oate (13). Compound **13** was synthesized under the same coupling conditions used for compounds **4** and **9** by starting from Z-Lys(Boc)-OH (1.05 g, 2.75 mmol) and the amino derivative **5** (1.09 g, 2.50 mmol). The desired product was obtained as a yellowish oil (1.40 g, 80% yield) after column chromatography. ESI-MS: calcd for C₃₆H₅₃N₄O₉, 685.84 [M + H]⁺; found, 685.73 [M + H]⁺. T_R = 26.16 min. ¹H NMR (400 MHz, CDCl₃): δ 7.37–7.35 (m, 5H), 7.31–7.26 (m, 5H), 6.99 (s, 1H), 5.85 (d, J = 7.7 Hz, 1H), 5.13–4.96 (m, 2H), 4.76 (s, 2H), 4.07 (q, J = 7.1 Hz, 2H), 3.73–3.59 (m, 2H), 3.28–3.13 (m, 2H), 3.03–2.97 (m, 2H), 2.47–2.40 (m, 2H), 2.28 (t, J = 7.3 Hz, 2H), 1.94–1.88 (m, 2H), 1.84–1.69 (m, 3H), 1.62–1.58 (m, 1H), 1.39 (s, 11H), 1.31 (dd, J = 19.1, 12.2 Hz, 2H), 1.19 (t, J = 7.1 Hz, 3H). ¹³C NMR (CDCl₃): δ 13C NMR (101 MHz, cdcl3) δ 174.3, 172.9, 172.0, 156.2, 136.2, 134.1, 129.2, 128.9, 128.6, 128.4, 128.0, 127.9, 78.9, 76.2, 66.8, 60.4, 54.8, 44.4, 39.9, 39.2, 38.5, 32.2, 31.2, 29.7, 29.4, 28.3, 23.7, 22.4, 22.1, 14.1.

Synthesis of Ethyl 17,27,37,47-Tetrakis(benzyloxy)-10-(((benzyloxy)carbonyl)amino)-2,2-dimethyl-4,11,16,21,26,31,36,41,46-nonaoxo-3-oxa-5,12,17,22,27,32,37,42,47-nonaazahenpentacontan-51-oate (15). The tetramer **15** was synthesized under the same coupling conditions used for compounds **4**, **9**, and **13** by starting from the acid derivative **14** (0.61 g, 0.93 mmol) and the amino derivative **12** (0.90 g, 1.02 mmol). The desired product was obtained as a colorless oil (0.97 g, 69% yield) after column chromatography. ESI-MS: calcd for C₈₁H₁₁₃N₁₀O₁₈, 1514.85 [M + H]⁺; found, 1514.16 [M + H]⁺, 775.95 [M + 2H]²⁺. T_R = 25.88 min. ¹H NMR (400 MHz, CDCl₃): δ 7.47–7.28 (m, 25H), 5.13–4.96 (m, 2H), 4.86–4.65 (m, 8H), 4.09 (dd, J = 13.8, 6.8 Hz, 3H), 3.80–3.53 (m, 8H), 3.32–3.10 (m, 7H), 3.04–3.00 (m, 2H), 2.54–2.34 (m, 7H), 2.29 (t, J = 7.0 Hz, 2H), 2.23–2.05 (m, 6H), 1.96–1.90 (m, 8H), 1.82–1.68 (m, 8H), 1.66–1.53 (m, 2H), 1.40 (s, 12H), 1.25–1.19 (m, 4H), 0.94–0.85 (m, 2H). ¹³C NMR (CDCl₃): δ 174.5, 172.9, 136.2, 134.0, 129.2, 129.1, 128.8, 128.5, 128.2, 128.0, 67.0, 60.5, 55.0, 44.1, 39.4, 39.0, 32.9, 32.0, 31.3, 29.8, 29.7, 29.5, 28.4, 23.9, 23.0, 22.5, 22.2, 14.2.

Synthesis of Benzyl (6,16,26,36-Tetrakis(benzyloxy)-2,7,12,17,22,27,32,37,42-nonaoxo-1,6,11,16,21,26,31,36,41-nonaazacycloheptatetracontan-43-yl)carbamate (16). Compound **15** was Boc-protected as described for **3**. Then, the ethyl group was hydrolyzed by LiOH as for **6**. To a dilute solution of the fully deprotected tetramer (0.55 g, 0.367 mmol) in DMF (40 mL) were added HATU (0.154 g, 0.40 mmol) and DIPEA (0.07 mL, 0.40 mmol) dropwise at 0 °C. The reaction mixture was stirred for 3 h. Then, the solvent was removed, and the residue was extracted with ethyl acetate and an aqueous solution of citric acid (10%), a solution of NaHCO₃ (5%), and brine. The crude product was purified via semipreparative HPLC, giving the desired product as a colorless oil (0.29 g, 57% yield). ESI-MS: calcd for C₇₄H₉₉N₁₀O₁₅, 1368.66 [M + H]⁺; found, 1368.36 [M + H]⁺, 684.74 [M+2H]²⁺. T_R = 25.73 min. ¹H NMR (400 MHz, DMSO-*d*₆): δ 7.96–7.63 (m, 6H), 7.56–7.13 (m, 25H), 5.06–4.89 (m, 2H), 4.85–4.71 (m, 8H), 4.00–3.76 (m,

3H), 3.09–2.88 (m, 10H), 2.84–2.61 (m, 1H), 2.43–2.25 (m, 8H), 2.09–1.93 (m, 9H), 1.75–1.71 (m, 9H), 1.59–1.53 (m, 8H), 1.39–1.08 (m, 7H), 1.07–0.76 (m, 2H). ¹³C NMR (DMSO-*d*₆): δ 171.8, 156.4, 135.2, 129.8, 129.1, 128.9, 128.7, 128.2, 128.1, 75.7, 65.8, 55.1, 44.5, 38.5, 33.0, 29.5, 24.7, 23.2.

Synthesis of (S)-43-Amino-6,16,26,36-tetrahydroxy-1,6,11,16,21,26,31,36,41-nonaazacycloheptatetracontane-2,7,12,17,22,27,32,37,42-nonaone (17, H₄L2). Compound **17** was synthesized as previously described for **11** by starting from derivative **16** (0.13 g, 71% yield). ESI-MS: calcd for C₃₈H₆₉N₁₀O₁₃, 874.03 [M + H]⁺; found, 873.75 [M + H]⁺. T_R = 21.28 min. ¹H NMR (400 MHz, DMSO-*d*₆): δ 9.64–9.59 (m, 3H), 8.07–8.04 (m, 2H), 7.86–7.71 (m, 4H), 3.47–3.44 (m, 9H), 3.04–2.99 (m, 10H), 2.42–2.22 (m, 8H), 2.04–2.00 (m, 8H), 1.76–1.49 (m, 20H), 1.47–1.16 (m, 5H). ¹³C NMR (DMSO-*d*₆): δ 172.9, 172.2, 168.7, 158., 56.5, 52.7, 47.2, 38.7, 33.0, 31.2, 29.7, 29.1, 24.8, 24.5, 23.0, 22.1. HR-ESI-MS *m/z* 873.50488; calcd for C₃₈H₆₉N₁₀O₁₃ ([M + H]⁺) 873.50401. Anal. Calcd for C₃₈H₆₈N₁₀O₁₃: C, 52.3; H, 7.9; N, 16.0. Found: C, 52.3; H, 7.8; N, 16.1.

Thermodynamic Solution Studies. General Considerations.

Unless otherwise stated, all commercially available reagents and solvents were of analytical grade, were purchased from commercial suppliers (Sigma-Aldrich, Titripur, Merck, Fisher Scientific, Fluka), and were used as received without further purification. All solutions were prepared in doubly distilled water. A stock solution of Fe(III) was prepared immediately before use from Fe(ClO₄)₃·xH₂O in 0.01 M HClO₄ and standardized by an inductively coupled plasma–optical emission spectrometer (ICP-OES; iCAP 7400 Duo ICP-OES) along with spectrophotometric determination, on the basis of the molar extinction coefficient ε = 4160 M⁻¹ cm⁻¹ at 240 nm.^{70,71} Stock solutions of Ga(III) and Zr(IV) were prepared immediately before use from Ga(ClO₄)₃·xH₂O and anhydrous ZrCl₄, respectively, in 0.1 M HClO₄ to prevent hydrolysis and standardized by ICP-OES (iCAP 7400 Duo ICP-OES) along with direct titration with ethylenediaminetetraacetic acid (EDTA).^{72,73} The HClO₄ solutions were titrated with standardized NaOH (0.1 N). The carbonate-free NaOH solution was standardized by titration with potassium hydrogen phthalate (KHP). All stock solutions were prepared using a R200D Sartorius analytical balance (with 0.01 mg precision).

All measurements were performed at 0.1 M NaClO₄ ionic strength, which was chosen instead of 1.0 M NaClO₄ ionic strength in order to increase the solubility of the investigated ligands and their complexes. We are aware that some measurements were performed at a very acidic pH (<1), where the ionic strength 0.1 M is not enough to keep the ionic activity stable, but due to the decomposition of hydroxamate ligands in strong acids,^{43,51} all measurements performed below pH 1 were assumed to be endowed with a large error and (i) were not taken into account during data evaluation or (ii) precluded from the discussion.

Electrospray Ionization Mass Spectrometry (ESI-MS). ESI-MS data were recorded on a Bruker Q-FTMS spectrometer. The instrumental parameters were as follows: scan range, *m/z* 200–1600; dry gas, nitrogen; temperature, 170 °C; capillary voltage, 4500 V; ion energy, 5 eV. The capillary voltage was optimized to the highest signal to noise ratio. The spectra were recorded in the positive mode. Compounds were dissolved in a MeOH/H₂O solution (80/20 by weight); the same solvent mixture was used to dilute the matrix solutions to the concentration range of 0.01 mM. The Fe(III), Ga(III), and Zr(IV) and stock solutions were prepared as described previously and added to the ligand solutions in 1/1, 2/1 and 1/3 mixtures for Fe(III) and Ga(III) and 1/1 and 1/3 mixtures for Zr(IV), all at pH 3 (the pH was adjusted by using acetic acid). The free hydrogen ion concentration was measured with a Mettler-Toledo InLab Semi-Micro combined glass electrode filled with NaCl in MeOH/H₂O (80/20 by weight). Potential differences were measured with a Beckman φ72 pH meter, standardized according to the classical methods with buffers prepared according to reported procedures in MeOH/H₂O solvent.^{74,75}

Potentiometric Titrations. The potentiometric titrations of ligands and their complexes were carried out using a Titrand 905

(Metrohm) automatic titrator system, equipped with a combined glass electrode (Mettler Toledo, InLab Semi-Micro, with XEROLYT EXTRA Polymer filling) and a 800 Dosino dosing system, equipped with a 2 mL micro buret. The ionic strength was fixed at $I = 0.1$ M with NaClO_4 . The electrode was calibrated daily in terms of hydrogen ion concentration using HClO_4 (0.1 M) with CO_2 -free NaOH solutions (0.1 M).⁷⁶ A stream of high-purity argon, presaturated with water vapor, was passed over the surface of the solution cell, the cell was filled with 50 mL of the studied solution, and the system was thermostated at 25.0 ± 0.2 °C. At least three titrations were performed for each system, with a starting concentration of the ligand of 1 mM and a 1:1 metal to ligand molar ratio with a 10% excess of the ligand in the pH range 2–11. The purity and exact concentration of the ligand solutions were determined using the Gran method.⁷⁷ Special care was taken to ensure that complete equilibration was attained. The titration curves were carefully checked and did not display any pH fluctuations that often accompany the precipitation of metal hydroxides. The potentiometric data were refined with the SUPERQUAD⁷⁸ and HYPERQUAD⁷⁹ programs, which use nonlinear least-squares methods. The successive protonation constants of the ligand were calculated from the cumulative constants determined with the program and defined by eqs 1 and 2 (charges are omitted for clarity).



$$K_n^{\text{H}} = \frac{[\text{H}_n\text{L}]}{[\text{H}_{n-1}\text{L}][\text{H}]} \quad (2)$$

The stability constants calculated for metal complexes are defined by eqs 3 and 4:



$$\beta_{\text{M}_p\text{H}_q\text{L}_r} = \frac{[\text{M}_p\text{H}_q\text{L}_r]}{[\text{M}]^p[\text{H}]^q[\text{L}]^r} \quad (4)$$

The uncertainties in the log K values correspond to the added standard deviations in the cumulative constants.

pH-Dependent UV–Vis Titrations. The pH-dependent UV–vis spectrophotometric experiments for the Fe(III)- $\text{H}_3\text{L1}$, Fe(III)- $\text{H}_4\text{L2}$, Ga(III)- $\text{H}_3\text{L1}$, Ga(III)- $\text{H}_4\text{L2}$, Ga(III)- $\text{H}_3\text{L4}$, Zr(IV)- $\text{H}_3\text{L1}$, Zr(IV)- $\text{H}_4\text{L2}$, Zr(IV)- $\text{H}_3\text{L3}$, and Zr(IV)- $\text{H}_3\text{L4}$ systems were carried out as a function of concentration with a Varian Cary 300 Bio spectrophotometer in the 300–700 nm range for iron complexes and 200–300 nm range for gallium and zirconium solutions using Hellma quartz optical cells with a 1 cm path length. To calculate the stability constants for investigated systems, two sets of pH-dependent UV–vis titrations were carried out: in the pH ranges (i) 0.1–2 and (ii) 2–11. In the (i) series, the experiments were performed by making 20 samples, differing by 0.1 pH unit, with a constant total volume of 0.7 mL and concentration of metal ion of ~ 0.05 –0.1 mM and metal to ligand molar ratio of 1:1; for all samples, the ionic strength was adjusted to 0.1 M by the addition of NaClO_4 and the pH (range 0.1–2.0) was controlled by the concentration of HClO_4 . After preparation, each solution was allowed to equilibrate for about 1 h, and then its UV–vis spectrum was recorded. This was necessary to minimize the effects of hydroxamate ligand hydrolysis, which occurs in strong acid.^{43,51} In the (ii) set of experiments, 3 mL of a solution containing a 1:1 Fe(III):ligand molar ratio, where the ferric concentration was around 0.20 mM, was introduced into a cell and the pH was adjusted by adding the proper microvolume of HClO_4 ; the solutions were allowed to equilibrate (up to 30 min) and checked with a Mettler Toledo Super Easy pH meter with an accuracy of ± 0.01 , and then the spectra were recorded.

Metal Competition Batch UV–Vis Titrations. In order to calculate the stability constants of the investigated complexes, several competitive titrations were performed. All of them were carried out as a function of concentration with a Varian Cary 300 Bio spectrophotometer in the 300–650 nm range (with 1 nm precision) using Hellma quartz optical cells with a 1 cm path length.

Spectrophotometric titrations were performed on samples with a concentration of the ligand of ~ 0.05 –0.1 mM and $I = 0.1$ M (completed by adding NaClO_4), at 25.0 ± 0.1 °C; pH 1.5 or 2.0 was adjusted by adding the proper volume of HClO_4 . In general, the stock solution of the starting complex (Fe(III)-L, Ga(III)-L, or Zr(IV)-L, respectively) was divided into several aliquots to which an excess of titrant (Ga(III), Fe(III), or Zr(IV)-NTA, respectively) was added. After preparation, each solution was allowed to equilibrate for about 1 h, and then its UV–vis spectrum was recorded. The vials were kept in the dark, and the absorbance was measured again after 24, 48, and 120 h. Changes were observed only between the spectra collected after 1 and 24 h, indicating that an equilibrium was attained.

In order to determine the log β values of $[\text{GaHL}]^+$ for $\text{H}_3\text{L1}$ and $\text{H}_3\text{L4}$ and of $[\text{GaH}_2\text{L}_2]^+$, competition experiments at pH 1.5 of (i) Fe(III)- H_3L + Ga(III) and (ii) Ga(III)-L + Fe(III) were performed. For (i) 15 samples with a constant concentration of Fe(III) ions and L (1:1) were titrated by up to 600 equiv of Ga(III) ions; for Ga(III)-L + Fe(III), 18 samples with a constant concentration of Ga(III) ions and H_3L (1:1) were titrated by up to 4 equiv of Fe(III) ions.

In order to determine the log β values of $[\text{ZrL}]^+$ for $\text{H}_3\text{L1}$, $\text{H}_4\text{L2}$, $\text{H}_3\text{L3}$, $\text{H}_3\text{L4}$, and DFOE and for $[\text{ZrHL}_2]^+$, competition experiments at pH 1.5 or 2, Fe(III)-L + Zr(IV)-NTA, were performed. In each experiment 18 samples with a constant concentration of Fe(III) ions and ligands were titrated by up to 50 equiv of a Zr(IV)-NTA solution with a metal to ligand molar ratio of 1:3, starting from 0 equiv.

Data Treatment. In the calculations of complex stability constants, the protonation constants of free ligands (Table 1) and the constants were related to hydrolytic species being taken into account: Ga(III),⁶⁰ $\text{Ga}(\text{OH})^{2+}$ log $\beta_{\text{GaH}_-1} = -3.11$, $\text{Ga}(\text{OH})_2^+$ log $\beta_{\text{GaH}_-2} = -7.66$, $\text{Ga}(\text{OH})_3$ log $\beta_{\text{GaH}_-3} = -11.94$, $\text{Ga}(\text{OH})_4^-$ log $\beta_{\text{GaH}_-4} = -15.66$, to Fe(III),⁸⁰ $\text{Fe}(\text{OH})^{2+}$ log $\beta_{\text{FeH}_-1} = -2.56$, $\text{Fe}(\text{OH})_2^+$ log $\beta_{\text{FeH}_-2} = -6.2$, $\text{Fe}(\text{OH})_3$ log $\beta_{\text{FeH}_-3} = -11.44$, $\text{Fe}(\text{OH})_4^-$ log $\beta_{\text{FeH}_-4} = -21.88$, $\text{Fe}(\text{OH})_5^{2-}$ log $\beta_{\text{FeH}_-5} = -2.74$, $\text{Fe}(\text{OH})_6^{3-}$, and Zr(IV),⁶⁰ $\text{Zr}(\text{OH})^{3+}$ log $\beta_{\text{ZrH}_-1} = -0.56$, $\text{Zr}(\text{OH})_2^{2+}$ log $\beta_{\text{ZrH}_-2} = -1.44$, $\text{Zr}(\text{OH})_4$ log $\beta_{\text{ZrH}_-4} = -8.85$, $\text{Zr}(\text{OH})_6^{2-}$ log $\beta_{\text{ZrH}_-6} = -30.6$, $\text{Zr}_3(\text{OH})_4^{8+}$ log $\beta_{\text{Zr}_3\text{H}_-4} = -6.96$, $\text{Zr}_4(\text{OH})_8^{8+}$ log $\beta_{\text{Zr}_4\text{H}_-8} = 6.52$, $\text{Zr}_3(\text{OH})_9^{3+}$ log $\beta_{\text{Zr}_3\text{H}_-9} = 12.19$. The Zr(IV) hydrolysis constants for the species $\text{Zr}(\text{OH})^{3+}$, $\text{Zr}(\text{OH})_2^{2+}$, $\text{Zr}(\text{OH})_4$, and $\text{Zr}_3(\text{OH})_4^{8+}$ were recalculated for 0.1 M NaClO_4 ionic strength according to literature parameters.⁶⁰ The $\text{p}K_w$ value used in the calculation at the 0.1 M NaClO_4 ionic strength was -13.77 .⁸¹

The UV–vis data were refined to obtain the overall binding constant using SPECFIT/32 software^{82–84} that adjusts the absorptivity and the stability constants of the species formed at equilibrium. Specfit uses factor analysis to reduce the absorbance matrix and to extract the eigenvalues prior to the multiwavelength fit of the reduced data set according to the Marquardt algorithm.^{82–84} Uncertainties in log β were calculated from the standard deviation.

The competition data were refined to obtain the overall binding constant using SPECFIT/32 software.^{82–84} The protonation constants of ligands and formation constants for iron complexes (Table 1 and the literature²⁴) were used as fixed parameters during data analysis. The concentration of iron complexes was calculated from the absorbance spectra (collected in the 300–700 nm range). Hydrolytic forms of the ferric ion in the studied pH range are characterized by an absorption band with a λ_{max} value of below 300 nm, and therefore they are beyond the experimental wavelength window. However, the spectrum of Fe(III) in solution at the pH of the experiment was fixed in the calculations.^{85,86} The stability constants of the Fe(III)-NTA⁴⁸ and Zr(IV)-NTA²⁷ complexes were taken from the literature and were used as fixed constants during the evaluation of the stability constants of the zirconium complexes.

The competition equilibrium is described by eqs 5 and 6:



$$K = \frac{[\text{FeL}][\text{ZrNTA}]}{[\text{FeNTA}][\text{ZrL}]} \quad (6)$$

The molecular charges are omitted for clarity. The data were processed using Origin 7.0. The species distribution diagrams were computed with the HYSS program.⁷⁹

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.inorgchem.1c01622>.

ESI-MS spectra and UV–vis spectroscopic data from solution studies, detail of the Fe(III) complex solution study, and ¹H/¹³C NMR spectra and chromatograms of the synthesized compounds (PDF)

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Notes

The authors declare no competing financial interest.

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