



Cite this: *Dalton Trans.*, 2024, **53**, 14634Chelation of [¹¹¹In]In³⁺ with the dual-size-selective macrocycles py-macrodipa and py₂-macrodipa†Kevin K. Lee,  ‡^a Mou Chakraborty, ^{b,c} Aohan Hu, ^a Thines Kanagasundaram, ^a Daniel L. J. Thorek ^{b,c,d} and Justin J. Wilson  *^{‡a}

Indium-111 (¹¹¹In) is a diagnostic radiometal that is important in nuclear medicine for single-photon emission computed tomography (SPECT). In order to apply this radiometal, it needs to be stably chelated and conjugated to a targeting vector that delivers it to diseased tissue. Identifying effective chelators that are capable of binding and retaining [¹¹¹In]In³⁺ *in vivo* is an important research area. In this study, two 18-membered macrocyclic chelators, py-macrodipa and py₂-macrodipa, were investigated for their ability to form stable coordination complexes with In³⁺ and to be effectively radiolabeled with [¹¹¹In]In³⁺. The In³⁺ complexes of these two chelators were characterized by NMR spectroscopy, X-ray crystallography, and density functional theory calculations. These studies show that both py-macrodipa and py₂-macrodipa form 8-coordinate In³⁺ complexes and attain an asymmetric conformation, consistent with prior studies on this ligand class with small rare earth metal ions. Spectrophotometric titrations were carried out to determine the thermodynamic stability constants (log *K*_{ML}) of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺, which were found to be 18.96(6) and 19.53(5), respectively, where the values in parentheses are the errors of the last significant figures obtained from the standard deviation from three independent replicates. Radiolabeling studies showed that py-macrodipa and py₂-macrodipa can quantitatively be radiolabeled with [¹¹¹In]In³⁺ at 25 °C within 5 min, even at ligand concentrations as low as 1 μM. The *in vitro* stability of the radiolabeled complexes was investigated in human serum at 37 °C, revealing that ~90% of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺ remained intact after 7 days. The bio-distribution of these radiolabeled complexes in mice was investigated, showing lower uptake in the kidneys, liver, and blood at the 24 h mark compared to [¹¹¹In]InCl₃. These results demonstrate the potential of py-macrodipa and py₂-macrodipa as chelators for [¹¹¹In]In³⁺, suggesting their value for SPECT radiopharmaceuticals.

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Introduction

In recent years, substantial efforts in the pharmaceutical industry have been directed toward the development of new

therapeutic radiopharmaceutical agents for the treatment of disease.^{1–6} Notably, these efforts have resulted in FDA approval of Xofigo® (radium-223 dichloride citrate) and Pluvicto® (lutetium-177 vipivotide tetraxetan, also recognized as ¹⁷⁷Lu-PSMA-617) for treating adult patients with metastatic castration-resistant prostate cancer,^{7–9} and Lutathera® ([¹⁷⁷Lu]Lu-DOTA-TATE) for the treatment of gastroenteropancreatic neuroendocrine tumors.¹⁰ These radiotherapeutic agents use a radionuclide with a biologically compatible half-life that emits particles with sufficiently high linear energy transfer to damage biomolecules, and are usually coupled to a targeting moiety designed to selectively deliver them to cancer cells. Once properly localized, these compounds cause DNA damage, leading to cell death.¹¹ When implemented correctly, this targeted approach provides precise and effective treatments with fewer adverse effects compared to traditional cytotoxic drug candidates.

As reflected by this type of FDA-approved drugs, therapeutic radiopharmaceuticals have primarily relied on beta particle

^aDepartment of Chemistry and Chemical Biology, Cornell University, Ithaca, New York, 14853, USA. E-mail: justinwilson@cornell.edu

^bDepartment of Radiology, Washington University School of Medicine, St. Louis, Missouri, 63110, USA

^cProgram in Quantitative Molecular Therapeutics, Washington University School of Medicine, St. Louis, Missouri, 63110, USA

^dDepartment of Biomedical Engineering, Washington University, St. Louis, Missouri, 63110, USA

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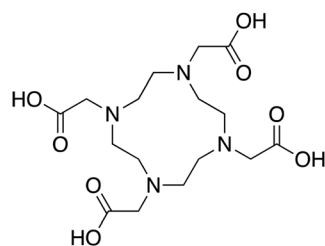
‡ Present Address: Department of Chemistry and Biochemistry, University of California Santa Barbara, Santa Barbara, California, 93117, USA.

(β)-emitting radionuclides, such as ^{131}I , ^{177}Lu , and ^{90}Y .^{12–18} The investigation of alpha particle (α)-emitters for therapeutic applications, enabled in part by advancements in isotope production and purification, has arisen as a promising alternative to β -emitters.¹⁹ For example, α -particles have a substantially higher linear energy transfer, meaning they deposit more of their energy over a shorter path length, giving rise to more damaging biological effects and greater cytotoxicity per decay. Additionally, the short range of α -particles allows for precise absorbed dose delivery to tumors while sparing healthy tissues, a property that may enhance the efficacy of targeted internal radiotherapy.^{20–23} One of the major challenges in harnessing α -emitting radionuclides for nuclear medicine has been developing chemical strategies to conjugate them to biological targeting vectors. Generally, a bifunctional chelator, which can stably bind radiometal ions and form a covalent linkage to targeting moieties, is needed. The macrocyclic ligand DOTA (1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid), shown in Chart 1, is an established chelator that works well for many different radiometals.^{24–27} A key limitation of DOTA is that it is significantly less effective for large metal ions, as reflected by its poorer thermodynamic affinity for larger compared to smaller lanthanide ions.²⁸ This limitation presents challenges associated with its use for promising α -emitting radionuclides, like ^{223}Ra , ^{225}Ac , and ^{227}Th , which are characterized by their large ionic radii.²⁹

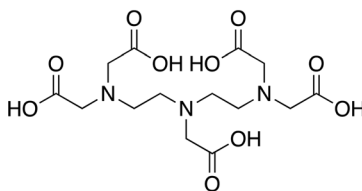
To address this problem, our group has developed chelating agents that possess high affinity and selectivity for large metal ions. These efforts have built upon the success of *N,N'*-bis[[6-carboxy-2-pyridil)methyl]-4,13-diaza-18-crown-6, or macropa

(Chart 1), a ligand that binds preferentially to large over small ions.³⁰ Given its high affinity for large metal ions, macropa has been shown to be effective for chelating the large α -emitting radiometal ions ^{225}Ac , ^{223}Ra , and ^{213}Bi , in addition to the large diagnostic radiometals ^{203}Pb , ^{132}La , ^{131}Ba , and ^{134}Ce .^{31–43} A limitation of macropa, however, is that it cannot effectively bind to smaller radiometal ions that are widely available and commonly employed in diagnostic applications, including ^{68}Ga , ^{111}In , and ^{44}Sc .⁴⁴

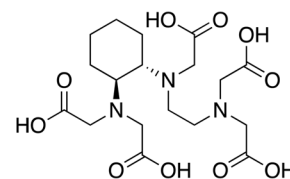
Continuing our investigations on expanded 18-membered macrocyclic chelators, we discovered the “dipa” class of ligands, which includes macrodipa, py-macrodipa, and py₂-macrodipa (Chart 1).^{45–47} A unique feature of this ligand class is its metal ion selectivity pattern that enables it to bind strongly to both large and small metal ions, but relatively poorly to intermediately sized ones. This property arises from the flexibility of this ligand, allowing it to undergo a conformational toggle when binding to small *versus* large metal ions. Within the realm of nuclear medicine, this ligand class can be applied simultaneously for large therapeutic radiometal ions and small diagnostic radiometal ions. This principle was recently demonstrated using the large radiometal ^{135}La , a therapeutic Auger electron-emitter, and the small radiometal ^{44}Sc , a diagnostic positron-emitter, in conjunction with py-macrodipa. Despite the significant ionic radii differences between La^{3+} and Sc^{3+} , py-macrodipa could be efficiently radio-labeled with both radionuclides and formed complexes that were stable to human serum over several half-lives.⁴⁶ Accordingly, a bifunctional version of py-macrodipa was synthesized and conjugated to a PSMA-targeting vector to deliver



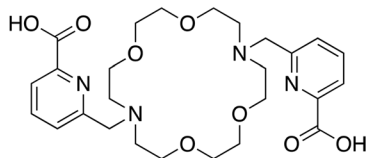
DOTA



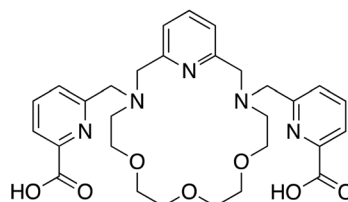
DTPA



CHX-A''-DTPA



macropa



py-macrodipa

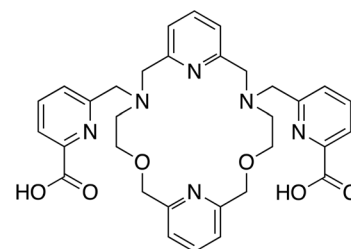
py₂-macrodipa

Chart 1 Structures of ligands discussed in this work.

^{135}La and ^{44}Sc to tumors *in vivo*.⁴⁸ Extending these efforts towards α -emitters, we demonstrated that py-macrodipa was also effective for $^{213}\text{BiBi}^{3+}$ chelation, and py₂-macrodipa could be applied for $^{225}\text{AcAc}^{3+}$.⁴⁷ These studies show the versatility of py-macrodipa and py₂-macrodipa for theranostic applications because of their abilities to chelate both large α -emitters and smaller diagnostic radiometals.

Among other conventional and highly promising diagnostic radiometals, ^{111}In has been extensively used within the clinic for SPECT imaging and has therapeutic potential as a robust Auger electron emitter.^{49–51} This cyclotron-produced isotope undergoes decay *via* electron capture with a half-life of 2.8 d, releasing γ rays (245 and 172 keV) that are well-suited for diagnostic applications.⁵² Although $^{111}\text{InIn}^{3+}$ can be effectively chelated with the common bifunctional chelators DOTA and CHX-A'-DTPA, its small 6-coordinate ionic radii of 0.80 Å (ref. 53) makes it unamenable for chelation by macropa. Thus, using ^{111}In as a partner diagnostic radionuclide for macropa-based bioconjugates for α therapy is not possible. In this work, we investigate the potential of the dual-size-selective chelators py-macrodipa and py₂-macrodipa for ^{111}In . Our results demonstrate that these chelators can effectively be used with this diagnostic radionuclide, thereby opening the possibility of pairing it with the larger α therapy isotopes ^{225}Ac and ^{213}Bi .

Results and discussion

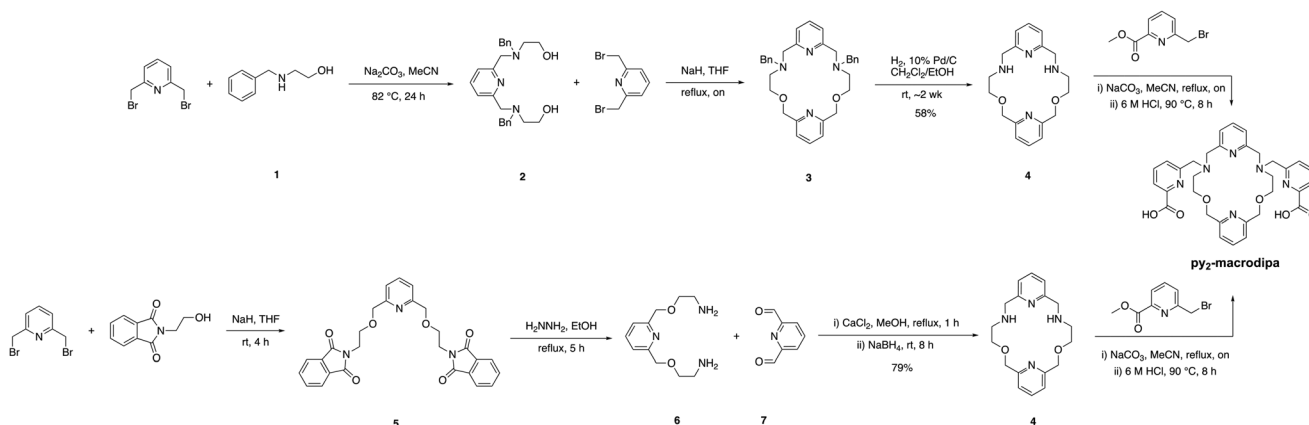
Ligand synthesis

The syntheses of py-macrodipa and py₂-macrodipa have previously been reported.^{46,47} Py-macrodipa can be accessed through a 3-step procedure, with the key reaction being the construction of the pyridine-containing macrocycle *via* a Ca^{2+} -templated reductive amination. Following the macrocycle formation, installation of the picolinate pendent arms through alkylation and acidic deprotection affords py-macrodipa. By contrast, the prior synthesis of py₂-macrodipa required a more extensive route to form the key dipyridine-containing macro-

cycle (4). To access 4, a procedure involving the construction of a benzyl-protected macrocycle was previously reported. This method requires deprotection of the benzyl group through reduction with H_2 over Pd/C catalysts, followed by the incorporation of picolinate pendent arms *via* alkylation (Scheme 1, top). Despite the success of this initial approach, the deprotection step is particularly time-consuming and low-yielding, thus prompting the pursuit of more efficient synthetic strategies. As an alternative approach, we explored the metal ion template-driven synthesis that was used successfully for py-macrodipa (Scheme 1, bottom). The key macrocycle-forming step in these new reactions involves the condensation of diamine 6 with the dialdehyde 7 in the presence of the templating Ca^{2+} ion, followed by the direct *in situ* reduction with NaBH_4 . The subsequent addition of picolinate arms to 4 was accomplished as previously reported.⁴⁷ The identity and purity of both intermediate and final products were confirmed *via* NMR spectroscopy and mass spectrometry (Fig. S1–S8, ESI†). This new synthetic strategy is advantageous over the previously reported one with a significantly shorter reaction time of 8 h compared to 2 weeks, which is required for benzyl deprotection. Additionally, the template-driven approach substantially increases the yield, achieving an overall yield of 79%.

Indium complex characterization

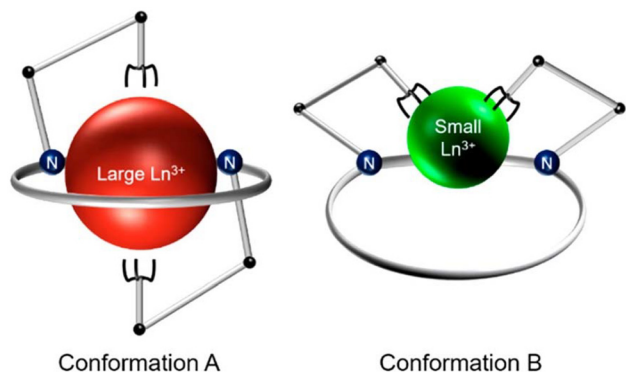
To assess the suitability of py-macrodipa and py₂-macrodipa for ^{111}In SPECT imaging applications, we characterized the In^{3+} complexes of these ligands. Because In^{3+} is diamagnetic, its complexes can be readily investigated *via* NMR spectroscopy. The ^1H NMR spectra of $[\text{In}(\text{py-macrodipa})]^+$ and $[\text{In}(\text{py}_2\text{-macrodipa})]^+$, obtained in D_2O , show sharp well-resolved resonances within the aromatic region and display diastereotopically split signals for the aliphatic protons. Notably, for both complexes, there are two distinct sets of signals for the picolinate arms, indicating that the complexes lack symmetry. In the context of prior studies with these ligands, it is well known that they form complexes of distinct symmetries depending on the size of the metal ion. We have previously described these



Scheme 1 Previously reported synthetic approach of dipyridine macrocyclic backbone (top) and improved synthetic approach of dipyridine macrocyclic backbone by using metal ion template synthesis (bottom).

different conformations as Conformation A and Conformation B (Scheme 2).^{46,47} The symmetric Conformation A is attained by large ions, like La^{3+} , whereas the asymmetric Conformation B occurs in complexes of small ions, like Sc^{3+} . Fig. 1 compares the ^1H NMR spectra of the In^{3+} complexes with those of La^{3+} and Sc^{3+} . The number of distinct resonances, as well as their chemical shift values, within the spectra of the In^{3+} complexes closely mirror those of the Sc^{3+} . Thus, like Sc^{3+} , In^{3+} attains the asymmetric Conformation B with both py-macrodipa and py_2 -macrodipa. Because the six-coordinate ionic radius of In^{3+} (0.80 Å) is comparable to that of Sc^{3+} (0.745 Å), this result is expected.⁵³

Further characterization of these In^{3+} complexes was achieved through single-crystal X-ray diffraction. The crystal structures of these complexes are depicted in Fig. 2. Consistent with the NMR spectroscopy results described above, these structures show both In^{3+} complexes to adopt the asymmetric Conformation B. For $[\text{In}(\text{py-macrodipa})]^+$, the



Scheme 2 Depiction of the conformational toggle present in Ln^{3+} -py-macrodipa and Ln^{3+} - py_2 -macrodipa complexes. Reproduced with permission from A. Hu, S. N. MacMillan, J. J. Wilson, Macrocylic Ligands with an Unprecedented Size-Selectivity Pattern for the Lanthanide Ions, *J. Am. Chem. Soc.* 2020, **142**, 13500–13506. Copyright 2020 American Chemical Society.

immediate coordination sphere of the ion is provided by the pyridyl group, two tertiary amines of the macrocycle, and the two picolinate arms. An eighth donor is provided by a macrocyclic ether oxygen atom. The In–O distance within this interaction is 2.64 Å, notably longer than In–O distances found in related In-crown ether complexes.^{54,55} Within $[\text{In}(\text{py}_2\text{-macrodipa})(\text{OH}_2)]^+$, a similar coordination sphere is observed. However, in place of the macrocycle ether oxygen atom, an inner-sphere water molecule is present. The In–O distance afforded by this interaction is 2.21 Å, longer than the In–OH₂ distances observed in other crystal structures.^{56–58} The presence of this inner-sphere water molecule leads to a more expanded conformation of the macrocycle that is stabilized by intramolecular hydrogen-bonding interactions. These two structures highlight some of the variability that is possible within Conformation B. The differential presence and absence of inner-sphere water within related Conformation B structures of $[\text{Lu}(\text{macrodipa})]^+$, $[\text{Sc}(\text{py-macrodipa})(\text{OH}_2)]^+$, $[\text{Lu}(\text{py-macrodipa})]^+$, and $[\text{Sc}(\text{py}_2\text{-macrodipa})(\text{OH}_2)]^+$ has also been previously observed.^{45–47} These results suggest a fluxionality associated with the exchange of the inner-sphere solvent molecules for the ether oxygen of the macrocycle. Slightly different crystallization conditions may lead to the isolation of these different forms.

DFT calculations

To further investigate the conformational preferences of the In^{3+} complexes of py-macrodipa and py_2 -macrodipa, DFT calculations were undertaken. The geometries of the In^{3+} complexes of these two ligands in both Conformations A and B were optimized using the ωB97XD functional, the LANL2DZ effective core potential and basis set for In, and the 6-31+G(d, p) basis set for other atoms. This method has been previously employed for Ln^{3+} -py-macrodipa and Ln^{3+} - py_2 -macrodipa complexes.^{46,47} Subsequently, their standard free energies (G°) were calculated. Given our observation of Conformation B with and without inner-sphere water molecules, both forms of this conformation were investigated in these calculations. The

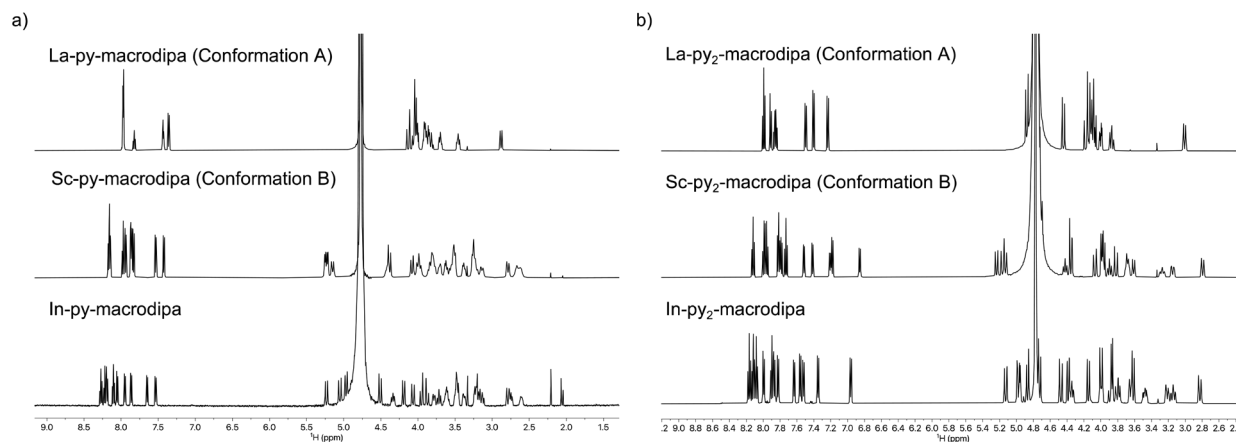


Fig. 1 (a) ^1H NMR spectra of $[\text{La}(\text{py-macrodipa})]^+$, $[\text{Sc}(\text{py-macrodipa})]^+$ and $[\text{In}(\text{py-macrodipa})]^+$ (500 MHz, D_2O , pD 5, 25 °C). (b) ^1H NMR spectra of $[\text{La}(\text{py}_2\text{-macrodipa})]^+$, $[\text{Sc}(\text{py}_2\text{-macrodipa})]^+$, and $[\text{In}(\text{py}_2\text{-macrodipa})]^+$ (500 MHz, D_2O , pD 5, 25 °C).

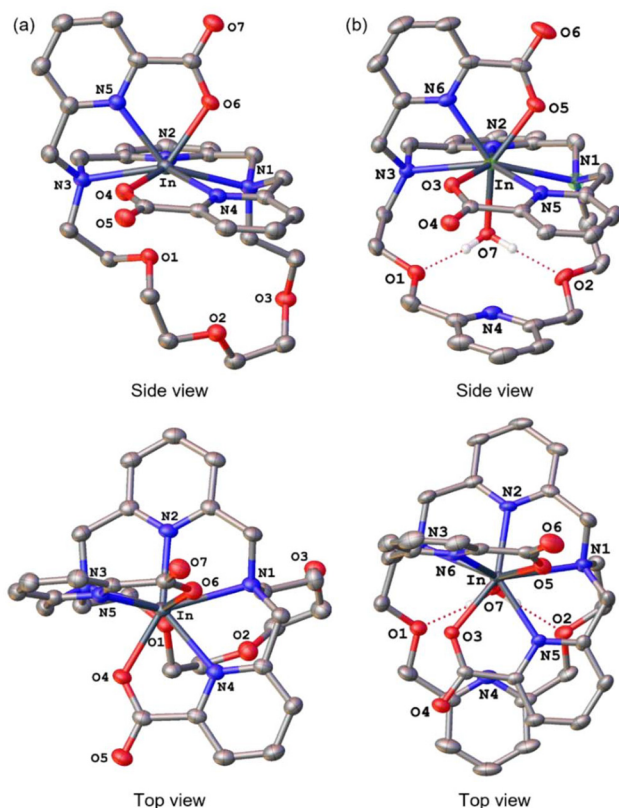
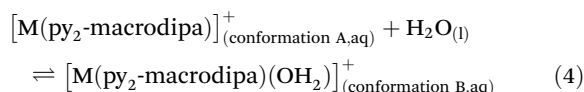
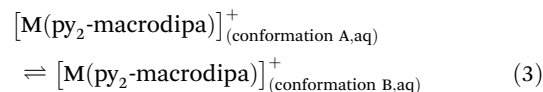
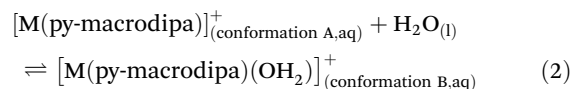
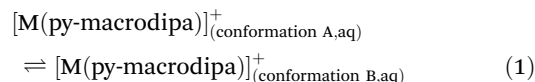


Fig. 2 Crystal structures of (a) $[\text{In}(\text{py-macrodipa})]^+$ and (b) $[\text{In}(\text{py}_2\text{-macrodipa})(\text{OH}_2)]^+$. Thermal ellipsoids are drawn at the 50% probability level. Solvent, counterions, and non-acidic hydrogen atoms are omitted for clarity.

equilibria between conformations A and B in an aqueous solution can be represented by balanced chemical reactions (eqn (1)–(4)), with the standard free energy changes denoted as ΔG° and the metal ions denoted as M. Our previous studies with Ln^{3+} -py-macrodipa and Ln^{3+} -py₂-macrodipa complexes have shown the ΔG° values are positive for the large La^{3+} ion, but negative for the small Sc^{3+} ion, indicating a thermodynamically driven transition from conformation A to B.^{46,47} For the equilibria that do not invoke an inner-sphere water molecule (eqn (1) and (3)), the ΔG° for $[\text{In}(\text{py-macrodipa})]^+$ and $[\text{In}(\text{py}_2\text{-macrodipa})]^+$ complexes are -17.1 and -20.5 kJ mol^{-1} , respectively, as detailed in Table 1. When an inner-

sphere water is included in Conformation B (eqn (2) and (4)), these ΔG° values become substantially more negative, attaining -74.6 and -84.6 kJ mol^{-1} for $[\text{In}(\text{py-macrodipa})]^+$ and $[\text{In}(\text{py}_2\text{-macrodipa})]^+$, respectively. The negative values, which indicate a thermodynamic preference for Conformation B, are expected due to the relatively small size of the In^{3+} ion, which is comparable to that of Sc^{3+} .⁵³ Thus, both In^{3+} complexes favor Conformation B, regardless of the presence or absence of an inner-sphere water molecule.



In^{3+} complex thermodynamic stability

To determine the thermodynamic stability of the In^{3+} complexes of py-macrodipa and py₂-macrodipa, UV-vis spectrophotometric titrations were carried out. The values for the protonation constants (K_i) of py-macrodipa and py₂-macrodipa were obtained from previous pH potentiometric titrations of these ligands.^{46,47} Although pH potentiometric titrations of $[\text{In}(\text{py-macrodipa})]^+$ and $[\text{In}(\text{py}_2\text{-macrodipa})]^+$ were initially attempted, their stability constants (K_{ML}) were too large to be determined over the pH range accessible to this method. Consequently, we conducted spectrophotometric titrations over an acidic pH (0.2–2.4) to determine these K_{ML} values. The UV-vis spectra of these titrations are shown in Fig. 3, and the data fitting analysis is shown in Fig. S9 and S10, ESI.† As shown in Table 2, the $\log K_{\text{InL}}$ values for both py-macrodipa and py₂-macrodipa (18.96 and 19.53, respectively) are several orders of magnitude larger than those of other metal ions. Notably, these values are even larger than those of the Sc^{3+} complexes of these ligands. Compared to the conventional chelators DOTA and DTPA, however, the K_{InL} values are still over 10 orders of magnitude smaller.^{59,60} Although the In^{3+} complexes of py-macrodipa and py₂-macrodipa do not attain the same level of thermodynamic stability as those of DOTA and DTPA, this property does not preclude their use for radiopharmaceutical applications. For example, the stability constant of macropa with La^{3+} is only 14.99, but this ligand has proven to be highly effective for *in vivo* applications using ²²⁵Ac, ^{132/135}La, and ¹³⁴Ce. Thus, in addition to thermodynamic stability, the inertness of relevant metal ion complexes needs to be considered under biologically relevant conditions.

Table 1 Changes of standard free energy in aqueous solution of DFT-optimized py-macrodipa and py₂-macrodipa complexes at 298 K

	$\Delta G^\circ(\text{A, B})$ (kJ mol^{-1})		
	La^{3+}	Sc^{3+}	In^{3+}
(1)	+55.4 ^a	-56.6 ^a	-17.1
(2)	+24.6 ^a	-80.5 ^a	-74.6
(3)	+61.2	-49.2	-20.5
(4)	+27.0 ^b	-87.4 ^b	-84.6

^a Ref. 46. ^b Ref. 47.

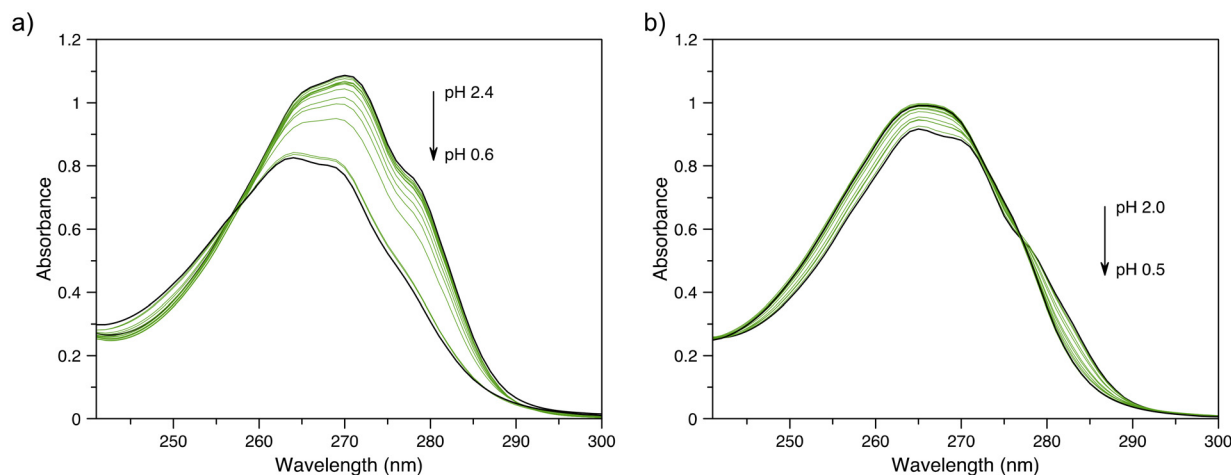


Fig. 3 (a) Representative UV-vis spectra of solutions containing py-macrodipa and InCl_3 in spectrophotometric titrations over the pH range of 0.6–2.4. (b) Representative UV-vis spectra of solutions containing py₂-macrodipa and InCl_3 in spectrophotometric titrations over the pH range 0.5–2.0.

Table 2 Stability constants of complexes formed with py-macrodipa, py₂-macrodipa, DOTA, and DTPA

	py-macrodipa	py ₂ -macrodipa	DOTA	DTPA
$\log K_{\text{InL}}$	18.96 ⁽⁶⁾	19.53 ⁽⁵⁾	23.9 ^a	29.0 ^d
$\log K_{\text{LaL}}$	14.31 ^b	16.68 ^c	22.9 ^e	18.02 ^f
$\log K_{\text{CeL}}$	14.65 ^b	17.13 ^c	23.4 ^e	19.06 ^f
$\log K_{\text{PrL}}$	14.81 ^b	17.28 ^c	23.0 ^e	19.64 ^f
$\log K_{\text{NdL}}$	14.51 ^b	17.11 ^c	23.0 ^e	20.23 ^f
$\log K_{\text{SmL}}$	13.66 ^b	16.56 ^c	23.0 ^e	20.79 ^f
$\log K_{\text{EuL}}$	13.29 ^b	15.93 ^c	23.5 ^e	20.74 ^f
$\log K_{\text{GdL}}$	12.63 ^b	15.25 ^c	24.7 ^e	21.15 ^f
$\log K_{\text{TbL}}$	11.95 ^b	14.76 ^c	24.2 ^e	21.15 ^f
$\log K_{\text{DyL}}$	11.47 ^b	14.04 ^c	24.8 ^e	21.23 ^f
$\log K_{\text{HoL}}$	10.69 ^b	12.68 ^c	24.5 ^e	21.43 ^f
$\log K_{\text{ErL}}$	10.60 ^b	12.17 ^c	24.4 ^e	21.41 ^f
$\log K_{\text{TmL}}$	10.92 ^b	11.98 ^c	24.4 ^e	21.10 ^f
$\log K_{\text{YbL}}$	11.31 ^b	11.82 ^c	25.0 ^e	20.96 ^f
$\log K_{\text{LuL}}$	11.54 ^b	11.90 ^c	25.4 ^e	21.14 ^f
$\log K_{\text{ScL}}$	15.83 ^b	16.28 ^c	30.79 ^g	27.43 ^g
pIn	19.3 ^h	19.7 ^h	18.8 ^h	25.7 ^h

^a Ref. 59 (0.1 M KCl, 25 °C). ^b Ref. 46 (0.1 M KCl, 25 °C). ^c Ref. 47 (0.1 M KCl, 25 °C). ^d Ref. 60 (0.1 M KNO₃, 25 °C). ^e Ref. 61 (0.1 M KCl, 25 °C). ^f Ref. 62 (0.1 M KNO₃, 25 °C). ^g Ref. 63 (0.1 M NMe₄Cl, 25 °C). ^h $p\text{In} = -\log [\text{In}^{3+}]_{\text{free}}$. Calculated for 10 μM total ligand and 1 μM total metal at pH 7.4 and 25 °C.

Radiolabeling experiments

Based on the promising In^{3+} coordination chemistry of these ligands, we next evaluated their ability to be labeled with radioactive $^{111}\text{In}^{3+}$. Concentration-dependent radiolabeling studies were conducted to ascertain the minimum concentration necessary for achieving high radiochemical conversions (RCCs). Varied concentrations of py-macrodipa and py₂-macrodipa, ranging from millimolar to nanomolar, were incubated with ~668 kBq of $^{111}\text{In}^{3+}$ in a pH 5.5 ammonium acetate buffer at 25 °C for 5 min. The RCCs of the radiolabeled complexes were quantified using radio-thin-layer chromatography (radio-TLC). Remarkably, the RCCs of both py-macrodipa and

py₂-macrodipa ^{111}In complexes were greater than 95% at ligand concentrations as low as 10⁻⁶ M (Fig. 4). By contrast, macropra was unable to incorporate $^{111}\text{In}^{3+}$ under the same conditions, even with ligand concentrations as high as 1 mM (Fig. S13, ESI[†]). To achieve quantitative RCCs of macropra with $^{111}\text{In}^{3+}$, 4 mM of macropra was needed to incubate with $^{111}\text{In}^{3+}$ at 25 °C for 1 h. (Fig. S14, ESI[†]). This result highlights how the preference of macropra for large ions limits its use for the smaller diagnostic $^{111}\text{In}^{3+}$ radiometal ion. In addition, the abilities of py-macrodipa and py₂-macrodipa to radiolabel at 25 °C marks an advantage over DOTA, for which the $^{111}\text{In}^{3+}$ complexes are most effectively formed by heating at 100 °C for 30 min.⁶⁴ The effective and mild ^{111}In -radiolabeling capabilities of py-macrodipa and py₂-macrodipa are particularly important in the context of the fact that these chelators can also similarly radiolabel the large therapeutic

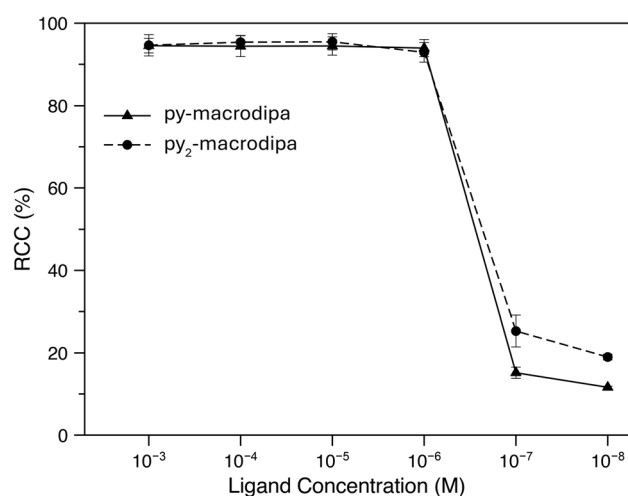


Fig. 4 Radiochemical conversions of $^{111}\text{In}^{3+}$ with py-macrodipa and py₂-macrodipa at different ligand concentrations (25 °C; pH 5.5; 5 min).

$[^{225}\text{Ac}]\text{Ac}^{3+}$, $[^{135}\text{La}]\text{La}^{3+}$, and $[^{213}\text{Bi}]\text{Bi}^{3+}$ radiometal ions. Lastly, the identities of the radiolabeled $[^{111}\text{In}]\text{In}^{3+}$ complexes were confirmed with radio-high-performance liquid chromatography (radio-HPLC). In these studies, HPLC retention times of the non-radioactive In^{3+} complexes of py-macrodipa and py_2 -macrodipa were used as reference values. As shown in Fig. 5, the UV-vis-HPLC chromatograms of the cold In^{3+} complexes display peaks with nearly identical retention times as the radio-HPLC chromatogram of the ^{111}In -labeled complexes. Therefore, the $[^{111}\text{In}]\text{In}^{3+}$ complexes are most likely identical in structure and conformation to their well-characterized non-radioactive analogs.

Human serum stability

After the successful formation of the $[^{111}\text{In}]\text{In}^{3+}$ complexes of py-macrodipa and py_2 -macrodipa, their stabilities under biologically relevant conditions were assessed. The complexes were formed using 8.77 MBq of $[^{111}\text{In}]\text{InCl}_3$ and a 1 μM ligand solution, with radiochemical purity confirmed by radio-TLC and radio-HPLC, and subsequently incubated in human serum at 37 $^\circ\text{C}$. As shown in Fig. 6, both $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ remained $98.4 \pm 1.8\%$ and $94.8 \pm 4.7\%$ intact, respectively, after 1 week based on radio-TLC. Radio-HPLC was also conducted at each time point to further confirm the identity of the activity (Fig. S15 and S16, ESI †). Despite the gradual decrease in the signal-to-noise ratio of the chromatograms due to the physical decay of the radionuclides, only a single peak corresponding to the $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ complexes were observed throughout the experiment. These results

indicate that both $[^{111}\text{In}]\text{In}^{3+}$ complexes are inert under these biologically relevant conditions, making them promising for radiopharmaceutical applications. In comparison, $73.2 \pm 10.2\%$ of $[^{111}\text{In}][\text{In}(\text{macropa})]^+$, formed with 4 mM of macropa and 8.77 MBq $[^{111}\text{In}]\text{In}^{3+}$, remained intact in human serum after 12 h at 37 $^\circ\text{C}$ (Fig. S17, ESI †). Furthermore, the similarly high stability of $[^{225}\text{Ac}][\text{Ac}(\text{py}_2\text{-macrodipa})]^+$ in human serum underscores the potential for using ^{225}Ac and ^{111}In as a therapeutic pair with py_2 -macrodipa.

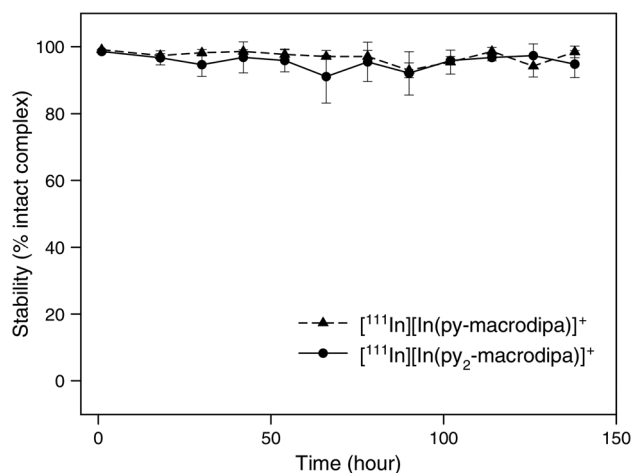


Fig. 6 Stability of $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ in human serum at 37 $^\circ\text{C}$.

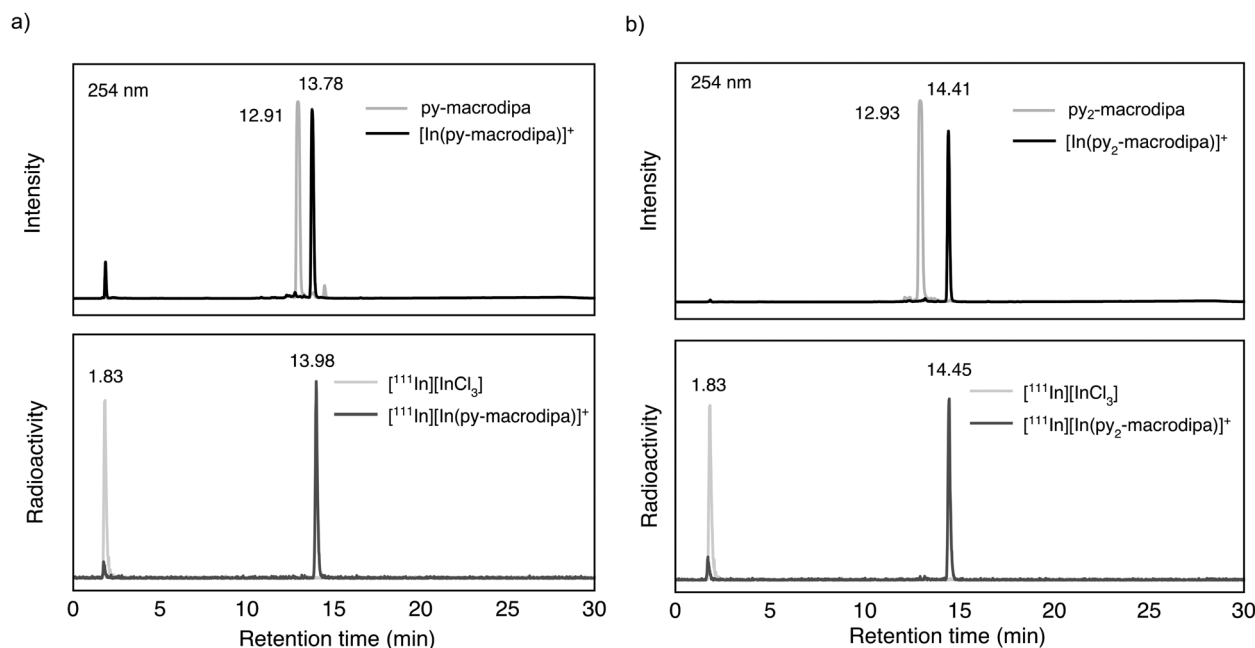


Fig. 5 (a) UV-vis-HPLC chromatogram (254 nm) of nonradioactive $[\text{In}(\text{py-macrodipa})]^+$ and py-macrodipa (top) compared with radio-HPLC chromatogram of $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}]\text{InCl}_3$ (bottom). (b) UV-vis-HPLC chromatogram (254 nm) of nonradioactive $[\text{In}(\text{py}_2\text{-macrodipa})]^+$ and py_2 -macrodipa (top) compared with the radio-HPLC chromatogram of $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ and $[^{111}\text{In}]\text{InCl}_3$ (bottom).

Biodistribution studies

To evaluate the *in vivo* stabilities of $[^{111}\text{In}][\text{In}(\text{macropa})]^+$, $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$, and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$, their biodistribution in mice was investigated in comparison to $[^{111}\text{In}]\text{InCl}_3$. The complexes were formed using 17 mM of each chelator and 11.8 MBq of $[^{111}\text{In}]\text{In}^{3+}$ after incubation for 5 min at 25 °C in pH 5.5 NH_4OAc buffer. The RCCs of all $[^{111}\text{In}]\text{In}^{3+}$ complexes were confirmed with radio-TLC. Swiss Webster mice were retro-orbitally (RO) administered with 0.703 MBq of complexes or $[^{111}\text{In}]\text{InCl}_3$ and sacrificed at time points of 0.5, 2, and 24 h for biodistribution analysis. These results are shown in Fig. 7. For $[^{111}\text{In}]\text{InCl}_3$, most of the injected activity is excreted through the kidneys with a smaller amount clearing through the liver. The activity persisted in the blood, heart, lungs, liver, kidneys, tibia, and calvaria. After 24 h post-injection, relatively high activity levels were observed in the kidneys, and moderate uptake was found in the liver and bone (Fig. 7a). These results are consistent with prior $[^{111}\text{In}]\text{InCl}_3$ biodistribution studies.⁶⁵

$[^{111}\text{In}][\text{In}(\text{macropa})]^+$ exhibited a similar biodistribution and clearance profile as $[^{111}\text{In}]\text{InCl}_3$ with high uptake in the kidney and slow blood clearance (Fig. 7b). The organs with the greatest amount of activity were the kidneys and liver. Low activities were observed in the heart, lungs, stomach, small intestine (SI), large intestine (LI), skin, fat, muscle, and brain, and moderate levels were found in blood, spleen, and bone. By contrast, the biodistribution profiles of $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ differ significantly from $[^{111}\text{In}]\text{InCl}_3$ (Fig. 7c and d). Low accumulation in the kidney, liver, and SI was observed, and a very low accumulation of activity in blood was found for both complexes 2 h post-injection, indicating rapid clearance. During this study, the complexes exhibited negligible uptake by other organs ($<1\%$ IA g^{-1} at 30 min post-injection), and activity did not accumulate over time, indicating no release of $[^{111}\text{In}]\text{In}^{3+}$ *in vivo*. These biodistribution studies reveal that $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$ and $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ are highly stable *in vivo*, a property that is crucial for their use as radiopharmaceutical agents.

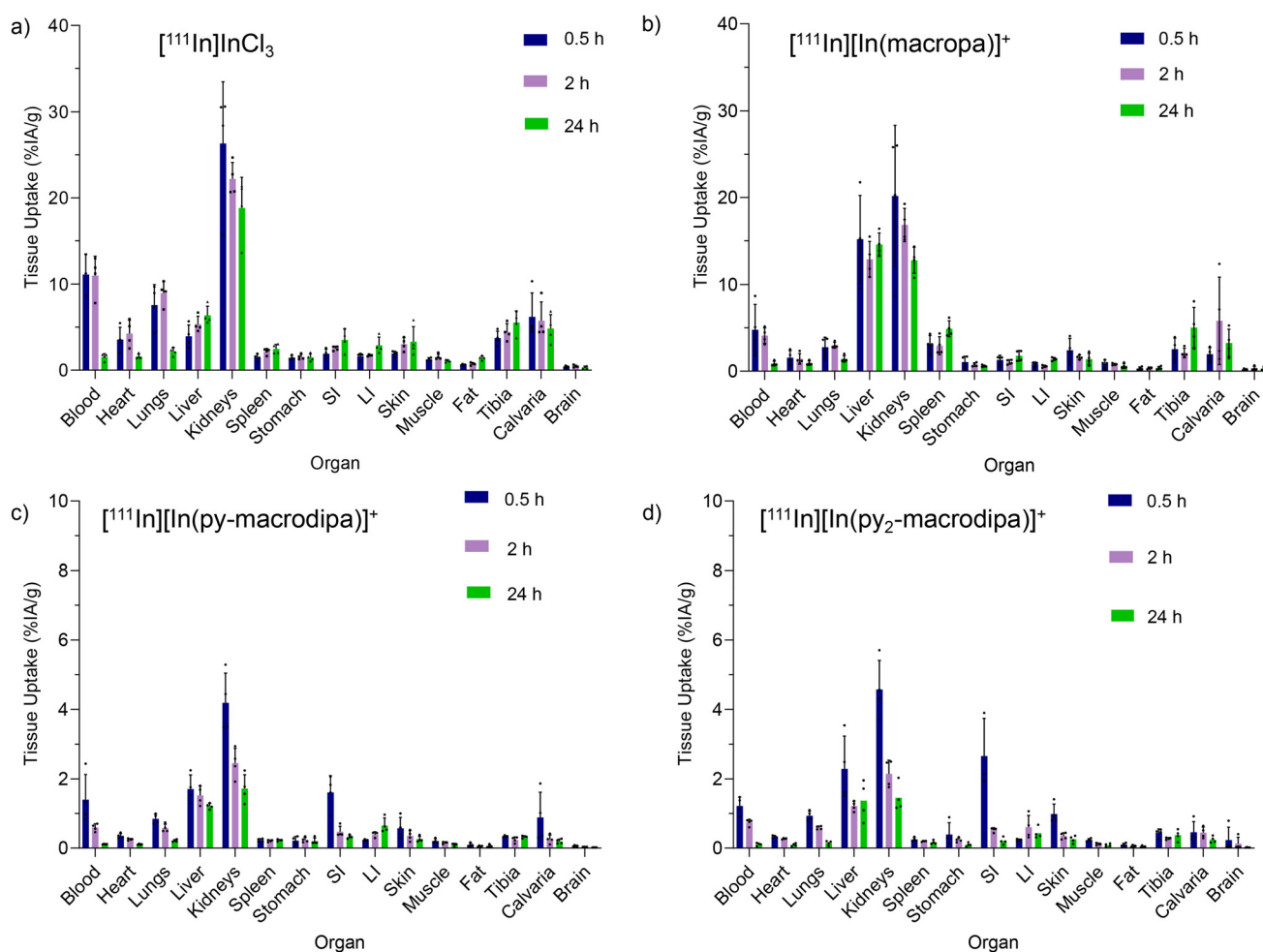


Fig. 7 *In vivo* biodistribution of (a) $[^{111}\text{In}]\text{InCl}_3$, (b) $[^{111}\text{In}][\text{In}(\text{macropa})]^+$, (c) $[^{111}\text{In}][\text{In}(\text{py-macrodipa})]^+$, and (d) $[^{111}\text{In}][\text{In}(\text{py}_2\text{-macrodipa})]^+$ following retro-orbital injection in mice. Four Naive Swiss Webster females of 4–6 weeks old were sacrificed after 0.5, 2, and 24 h. Values for each time point are given as percent injected activity per gram tissue ($\%IA\text{ g}^{-1}$).

Conclusions

In this study, we assessed the potential of macropa, py-macrodipa, and py₂-macrodipa as chelators for the widely used SPECT radioisotope ¹¹¹In. Through computational and experimental analyses, we characterized their coordination chemistry with In³⁺ and observed that both py-macrodipa and py₂-macrodipa ligands form stable complexes with this ion, adopting asymmetric Conformation B upon complexation. Furthermore, radiolabeling experiments demonstrated effective complexation with [¹¹¹In]In³⁺ under mild conditions, and the resulting radiolabeled complexes were stable in human serum. This excellent stability was also verified in mice, which showed that [¹¹¹In][In(py-macrodipa)]⁺ and [¹¹¹In][In(py₂-macrodipa)]⁺ did not accumulate in the kidneys, liver, and blood at the 0.5 h or 24 h mark, unlike [¹¹¹In]InCl₃ and [¹¹¹In][In(macropa)]⁺. In the context of earlier studies on the suitability of these chelators for large therapeutic radiometals like ¹³⁵La, ²¹³Bi, and ²²⁵Ac, their ability to be used with the small diagnostic ¹¹¹In radionuclide portend well for use in theranostic applications. The conformational flexibility provided by this ligand class presents an interesting opportunity to chelate useful radiometal ions of different ionic radii. Thus, py-macrodipa and py₂-macrodipa join several other recently reported chelators that have mutually high affinity for large α-emitting and small diagnostic radionuclides.^{66–73} Ongoing efforts are focused on synthesizing a bifunctional analog of py₂-macrodipa to enable its use for SPECT imaging with ¹¹¹In and α therapy with ²²⁵Ac and ²¹³Bi.

Experimental

General

All chemicals were purchased from commercial suppliers and used without further purification. py-Macrodipa, macropa, pyridine-2,6-dicarbaldehyde, and 6-bromomethylpyridine-2-carboxylic methyl ester were synthesized as previously described.⁴⁶ The compounds 2,6-bis(bromomethyl)pyridine and *N*-(2-hydroxyethyl) phthalimide were purchased from VWR (Radnor PA). Anhydrous solvents were obtained following storage over 3 Å molecular sieves, which were activated under heat. 1D-NMR (¹H and ¹³C {¹H}) spectra were acquired at 25 °C on a 500 MHz Bruker AV 3HD spectrometer equipped with a broadband Prodigy cryoprobe (Bruker, Billerica, MA). Water (18 MΩ cm) was purified using an ELGA PURELAB flex 2 (High Wycombe, U.K.). High-resolution mass spectra (MS) were recorded on an Exactive Orbitrap mass spectrometer (Thermo Fisher) in positive ionization mode with sample analysis in real-time (DART) mode. Elemental analyses (C, H, N) were carried out by Atlantic Microlab Inc. (Norcross, GA). High-performance liquid chromatography (HPLC) consisted of a CBM-20A communications bus module, an LC-20AT pump, and an SPD-20AV UV-Vis detector monitoring at 254 nm (Shimadzu, Japan). Analytical chromatography was carried out using a Luna C18 column, 100 Å, 5 μm, 150 mm × 3 mm

(Phenomenex, Torrance, CA) at a flow rate of 1.0 mL min⁻¹. [¹¹¹In]InCl₃ (in 0.01M HCl with radionuclide purity >99.9%) was purchased from BWXT Medical Ltd (Vancouver, Canada). Activity amounts of [¹¹¹In]InCl₃ solution were measured with a dose calibrator (CAPINTEC, CRC-15R) and 2480 automated γ-counter (WIZARD2 PerkinElmer). Glass microfiber chromatography paper impregnated with silica gel (iTLC-SG) was purchased from Agilent Technologies (Folsom CA). Female Swiss Webster mice (4–6 weeks, Charles River Laboratories) were purchased for *in vivo* biodistribution studies. All radioactive material handling and animal experiments were conducted in compliance with institutional regulations and approved by Environmental Health and Safety Radioactive Materials at Washington University in St. Louis under protocol D16-00245 and Institutional Animal Care and Use Committee protocol #22-0023.

Chelator synthesis

2,2'-(((Pyridine-2,6-diylbis(methylene))bis(oxy))bis(ethane-2,1-diyl))bis(isoindoline-1,3-dione) (5). To a solution of *N*-(2-hydroxyethyl)phthalimide (4.00 g, 20.9 mmol) in anhydrous tetrahydrofuran (30 mL) was added NaH (60% dispersion in mineral oil) (1.68 g, 41.0 mmol) at 0 °C over 1 h. Commercially available 2,6-bis(bromomethyl)pyridine (2.63 g, 10.5 mmol) was added to the mixture portion-wise over 30 min and allowed to stir at room temperature for 4 h. The mixture was then filtered, and the filtrate was concentrated under reduced pressure to give a yellow solid. The crude solid was then washed with 100 mL of methanol on filter paper to yield **5** (2.51 g, 52%) as an off-white solid. ¹H NMR (500 MHz, CDCl₃) δ = 7.84 (dq, *J* = 6.0, 3.2 Hz, 4H), 7.71 (dq, *J* = 5.1, 3.0 Hz, 4H), 7.55 (t, *J* = 7.7 Hz, 1H), 7.21 (d, *J* = 7.7 Hz, 2H), 4.58 (s, 4H), 3.97 (t, *J* = 5.7 Hz, 4H), 3.79 (t, *J* = 5.7 Hz, 4H). ¹³C {¹H} NMR (126 MHz, CDCl₃) δ = 168.39, 157.61, 134.08, 132.26, 123.42, 119.89, 68.12, 67.83, 37.71. DART-MS *m/z*: 486.17; calcd for [M + H]⁺: 486.17.

2,2'-((Pyridine-2,6-diylbis(methylene))bis(oxy))bis(ethan-1-amine) (6). To a solution of **5** (2.00 g, 4.12 mmol) in absolute ethanol (30 mL) was added 80% hydrazine monohydrate (0.60 mL, 10.1 mmol). The reaction mixture was heated at 80 °C for 4 h. A quantity of 10 mL of 6 M HCl was added, and the mixture was stirred at 80 °C for an additional 1 h. After cooling to room temperature, the mixture was filtered, and the collected solid was washed with water (10 mL). The combined filtrate was concentrated under reduced pressure to give a yellow oil. Ethanol was added to this crude oil and the precipitates were collected by filtration to afford **6** (0.94 g, 68%) as a white solid. ¹H NMR (500 MHz, D₂O, pD ≈ 5) δ = 8.58 (t, *J* = 8.0 Hz, 1H), 8.02 (d, *J* = 8.0 Hz, 2H), 5.04 (s, 4H), 3.94 (t, *J* = 5.0 Hz, 4H), 3.34 (t, *J* = 4.9 Hz, 4H). ¹³C {¹H} NMR (126 MHz, DMSO-*d*₆) δ = 158.26, 137.82, 120.24, 79.63, 73.43, 73.31, 41.77. DART-MS *m/z*: 226.15; calcd for [M + H]⁺: 226.15.

3,13-Dioxa-6,10-diaza-1,8(2,6)-dipyridinacyclotetradecaphane (4). Anhydrous methanol (10 mL) was added to a vial containing pyridine-2,6-dicarbaldehyde (0.20 g, 1.49 mmol) and CaCl₂·2H₂O (0.24 g, 1.57 mmol). The mixture was stirred at

room temperature until pyridine-2,6-dicarbaldehyde was fully dissolved. A solution of **6** (0.50 g, 1.49 mmol) in anhydrous methanol (10 mL) was then added dropwise. The mixture was stirred at room temperature for 30 min and subsequently heated at 65 °C for 1 h. After cooling on ice to 0 °C, solid NaBH₄ (0.28 g, 7.47 mmol) was added in small portions over 15 min. The reaction mixture was allowed to slowly return to RT and stirred for 8 h. The reaction was quenched with H₂O (10 mL), and methanol was removed under reduced pressure. The mixture was then extracted with 3 × 10 mL of CH₂Cl₂. The combined organic layers were dried over Na₂SO₄ and removed under reduced pressure. The crude product was purified by column chromatography (25 g silica gel, eluted with a gradient of 100% DCM to 75% DCM/MeOH) to afford **4** (0.39 g, 79%) as a clear oil. ¹H NMR (500 MHz, MeOD-*d*₄) δ = 7.80 (t, 1H, *J* = 7.7 Hz), 7.69 (t, 1H, *J* = 7.7 Hz), 7.35 (d, 2H, *J* = 7.7 Hz), 7.22 (d, 2H, *J* = 7.7 Hz), 4.56 (s, 4H), 3.80 (s, 4H), 3.73 (t, 4H, *J* = 4.8 Hz), 2.86 (t, 4H, *J* = 4.8 Hz). DART-MS *m/z*: 329.20; calcd for [M + H]⁺: 329.20.

X-ray crystallography

Low-temperature X-ray diffraction data of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺ were collected on a Rigaku XtaLAB Synergy diffractometer coupled to a Rigaku Hypix detector with Cu Kα radiation (λ = 1.54184 Å), from a PhotonJet micro-focus X-ray source at 100 K. The diffraction images were processed and scaled using the CrysAlisPro software (Oxford Diffraction LTD, TX). The structures were solved through intrinsic phasing using SHELXT⁷⁴ and refined against F² on all data by full-matrix least squares with SHELXL⁷⁵ following established refinement strategies.⁷⁶ All non-hydrogen atoms were refined anisotropically. All hydrogen atoms bound to carbon were included in the model at geometrically calculated positions and refined using a riding model. Hydrogen atoms bound to oxygen were located in the difference Fourier synthesis and subsequently refined semi-freely with the help of distance restraints. The isotropic displacement parameters of all hydrogen atoms were fixed to 1.2 times the *U*_{eq} value of the atoms they are linked to (1.5 times for methyl groups). [In(py₂-macrodipa)]⁺ contains disordered solvent molecules of Et₂O and MeOH but could not be satisfactorily modeled. Therefore, those solvents were treated as diffuse contributions to the overall scattering without specific atom positions using the solvent mask routine in Olex2 with a volume of 327.6 Å³ corresponding to 81.2 electrons.⁷⁷ The X-ray crystallographic data is deposited in the Cambridge Structural Database (CCDC) as crystallographic information files (cif) under deposition numbers 2373568 and 2373569.†

Stability constant measurements

The stability constants of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺ complexes were measured by UV-vis spectrophotometric titrations. Ligand stock solutions (17.83 mM py-macrodipa and 19.57 mM py₂-macrodipa) were made by dissolving the solid ligand in 18 MΩ·cm H₂O. An In³⁺ stock solution (21.18 mM) was made by dissolving InCl₃·4H₂O (99.9% purity)

in standardized HCl (0.1 M). The exact concentrations were determined by complexometric titrations with a standardized Na₂H₂EDTA solution (Alfa Aesar). The complexometric titrations were performed at pH 5.4 maintained by a sodium acetate buffer, and the endpoint was indicated by xylenol orange.

UV-vis spectra were recorded on a Shimadzu UV-1900 UV-vis spectrometer with a 1 cm quartz cuvette. The stability constants were determined with UV-vis spectrophotometric titrations using a batch method. Specifically, a set of solutions (around 15 samples) containing In³⁺ and py-macrodipa, and py₂-macrodipa (1 : 1 metal to ligand molar ratio, 45–80 μM) at different pH values were prepared. All samples were maintained at low pH (1.1 ≤ pH ≤ 2.4), which was adjusted with standardized HCl (0.1 M). Under these conditions, pH cannot be accurately measured with the glass electrode; pH values were calculated directly from the concentration of HCl used in each sample. The ionic strength was fixed at 0.1 M with KCl when possible. The solutions were incubated at 25 °C overnight after which all samples had reached equilibrium. The pH-dependent spectrophotometric data were analyzed with HypeSpec 2014 software.⁷⁸ The absorption band from 240–300 nm was used for analysis and the spectra for free ligand species were included in the fitting. The stability constants were calculated from the average of three independent titrations.

Radiolabeling

Py-macrodipa and py₂-macrodipa were dissolved in ammonium acetate buffer solution (pH 5.5) to give 10 mM stock solutions from which serial dilutions were used to prepare ligand solutions of concentrations ranging from 10^{−3} to 10^{−7} M. For [¹¹¹In]In³⁺ labeling studies, a 5 μL aliquot of the respective ligand solution was further diluted with 44 μL of 0.5 M NH₄OAc buffer solution (pH = 5.5), and 1 μL of [¹¹¹In]InCl₃ solution in 0.01 M HCl (0.668 MBq) was added. RCCs were determined with radio-iTLC using a mobile phase of 50 mM EDTA (pH 5.5). Under these conditions, chelated [¹¹¹In]In³⁺ remained at the baseline (*R*_f = 0), and uncomplexed [¹¹¹In]In³⁺ migrated with the solvent front (*R*_f ~1).⁷⁹ At 5 min, ~5 μL aliquots were spotted on the plates and developed. Once dried, RCCs were measured using a Dual Scan-Ram Scanner (Lablogic, UK) equipped with a NaI detector and quantified with Laura software. Radio-HPLC was further used to confirm the identity of the [¹¹¹In]In³⁺ complex with elution conditions: A: H₂O + 0.1% TFA, B: MeOH; 0 to 90% B linear gradient 20 min. [¹¹¹In][In(py-macrodipa)]⁺ (*t*_R = 13.98 min), [¹¹¹In][In(py₂-macrodipa)]⁺ (*t*_R = 14.45 min), and [¹¹¹In]InCl₃ (*t*_R = 1.83 min).

Human serum stability

The compounds [¹¹¹In][In(py-macrodipa)]⁺ and [¹¹¹In][In(py₂-macrodipa)]⁺ were prepared with the radiolabeling protocol as described above. A quantity of 1.5 mL of 0.33 mM ligand solutions of py-macrodipa and py₂-macrodipa were incubated with 9 μL of [¹¹¹In]InCl₃ (8.77 MBq) solution in 114.5 μL of 0.5 M

NH₄OAc buffer (pH = 5.5) for 5 min at 25 °C. A 375 µL portion of human serum, purchased from Sigma (Sera human Lot# SLCF9666), was added, and the mixture was incubated at 37 °C. At each time point, 10 µL of each solution was analyzed by radio-TLC to determine the % intact complex, and the values were tabulated as shown in Table S2, ESI.† At each time point, 50 µL of the mixture was added to 250 µL of MeOH to precipitate the proteins. After centrifugation, 10 µL of the supernatant liquid was used for radio-HPLC to identify the activity.⁸⁰

Biodistribution study

For preparation of [¹¹¹In][In(macropa)]⁺, 20 µL of [¹¹¹In]InCl₃ (11.8 MBq) in 0.01 M HCl (Sigma-Aldrich, trace-metal grade, 99.999%) was added to ammonium acetate buffer (250 µL, 0.5 M, pH 5.5) to adjust the pH to 5.5, and then 100 µL of 17 mM of macropa chelator was added. The mixture was vortexed and incubated at 25 °C for 5 min. Radiochemical purity was evaluated using iTLC-SG with a 50 mM EDTA pH 5.5 mobile phase. Radio-TLC was developed using a gas proportional scanner (Bioscan AR-2000) with a 1 min acquisition time. For subsequent *in vivo* evaluation, the labeled material was then diluted with 1.5 mL of normal saline (0.9% NaCl in deionized H₂O; Hospira), and activities were prepared in 30G syringes for administration. [¹¹¹In][In(py-macrodipa)]⁺ and [¹¹¹In][In(py₂-macrodipa)]⁺ were also prepared following the same labeling conditions. Healthy Swiss webster female mice from Charlies River Laboratories of 4–6 weeks old, were injected *via* the RO sinus with 0.703 MBq/100 µL [¹¹¹In]InCl₃, [¹¹¹In][In(macropa)]⁺, [¹¹¹In][In(py-macrodipa)]⁺, or [¹¹¹In][In(py₂-macrodipa)]⁺ complexes for biodistribution studies. Four animals for each complex at each time point 0.5, 2, and 24 h post-injection were euthanized by CO₂ asphyxiation before tissue dissection. A blood sample was acquired, and the organs of interest were collected (heart, lungs, liver, kidneys, spleen, stomach, small intestine, large intestine, skin, muscle, fat, tibia, calvaria, and brain), rinsed of excess blood, weighed, and measured by γ counting. Data were corrected for background and decay, and the activity of each tissue was normalized to an injection standard. The organ uptake values were calculated as percent of injected activity per gram of tissue (% IA g⁻¹).

DFT calculations

DFT calculations were carried out with the Gaussian 16 software package.⁸¹ The molecular geometries of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺ (Conformation A and B) were optimized employing the ωB97XD functional,^{82,83} and the basis set 6-31+G(d,p) was used for the lighter elements (C, N, H, and O).^{84,85} For the heavier indium atom, the LANL2DZ effective core potential, as well as its associated basis set, was applied.⁸⁶ In addition, frequency calculations were carried out on optimized geometries to confirm, *via* the absence of imaginary frequencies, that these structures are local minima on the potential energy surface. Zero-point energies and thermochemical correction factors were also afforded by the frequency

calculations. The effect of the aqueous environment was considered using the polarizable continuum model (PCM) by a single-point calculation on the optimized structures.⁸⁷ The initial Conformation B geometries of [In(py-macrodipa)]⁺ and [In(py₂-macrodipa)]⁺ were taken from the crystal structures. The initial Conformation A geometries were taken from the previously obtained crystal structure of La³⁺ complexes with the metal center replaced with In³⁺.^{46,47} For all aqueous species (everything except H₂O), a correction factor of 7.9 kJ mol⁻¹ was applied to adjust the free energy values from the software state default (298 K, 1 atm) to standard state in aqueous solution (298 K, 1 M). For H₂O(l), this correction factor is 17.9 kJ mol⁻¹, accounting for its larger standard state concentration in the liquid phase of 55.5 M.

Data availability

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

Conflicts of interest

There are no conflicts to declare.

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References

- 1 J. Czernin, I. Sonni, A. Razmaria and J. Calais, *J. Nucl. Med.*, 2019, **60**, 3S–12S.
- 2 D. A. Mulford, D. A. Scheinberg and J. G. Jurcic, *J. Nucl. Med.*, 2005, **46**, 199S–204S.
- 3 D. Delbeke and G. M. Segall, *J. Nucl. Med.*, 2011, **52**, 24S–28S.
- 4 M. Dondi, R. Kashyap, D. Paez, T. Pascual, J. Zaknun, F. M. Bastos and Y. Pynda, *J. Nucl. Med.*, 2011, **52**, 16S–23S.
- 5 G. Sgouros, L. Bodei, M. R. McDevitt and J. R. Nedrow, *Nat. Rev. Drug Discovery*, 2020, **19**, 589–608.
- 6 G. Kramer-Marek and J. Capala, *Tumor Biol.*, 2012, **33**, 629–640.
- 7 J. Fallah, S. Agrawal, H. Gittleman, M. H. Fiero, S. Subramaniam, C. John, W. Chen, T. K. Ricks, G. Niu,

- A. Fotenos, M. Wang, K. Chiang, W. F. Pierce, D. L. Suzman, S. Tang, R. Pazdur, L. Amiri-Kordestani, A. Ibrahim and P. G. Kluetz, *Clin. Cancer Res.*, 2023, **29**, 1651–1657.
- 8 M. Benešová, M. Schäfer, U. Bauder-Wüst, A. Afshar-Oromieh, C. Kratochwil, W. Mier, U. Haberkorn, K. Kopka and M. Eder, *J. Nucl. Med.*, 2015, **56**, 914–920.
- 9 P. G. Kluetz, W. Pierce, V. E. Maher, H. Zhang, S. Tang, P. Song, Q. Liu, M. T. Haber, E. E. Leutzinger, A. Al-Hakim, W. Chen, T. Palmby, E. Alebachew, R. Sridhara, A. Ibrahim, R. Justice and R. Pazdur, *Clin. Cancer Res.*, 2014, **20**, 9–14.
- 10 U. Hennrich and K. Kopka, *Pharmaceuticals*, 2019, **12**, 114.
- 11 J. Nonnekens, K. L. S. Chatalic, J. D. M. Molkenboer-Kuenen, C. E. M. T. Beerens, F. Bruchertseifer, A. Morgenstern, J. Veldhoven-Zweistra, M. Schottelius, H.-J. Wester, D. C. van Gent, W. M. van Weerden, O. C. Boerman, M. de Jong and S. Heskamp, *Cancer Biother. Radiopharm.*, 2017, **32**, 67–73.
- 12 M. S. Kaminski, K. R. Zasadny, I. R. Francis, A. W. Milik, C. W. Ross, S. D. Moon, S. M. Crawford, J. M. Burgess, N. A. Petry, G. M. Butchko, S. D. Glenn and R. L. Wahl, *N. Engl. J. Med.*, 1993, **329**, 459–465.
- 13 C. T. Sawin and D. V. Becker, *Thyroid*, 1997, **7**, 163–176.
- 14 T. E. Witzig, L. I. Gordon, F. Cabanillas, M. S. Czuczman, C. Emmanouilides, R. Joyce, B. L. Pohlman, N. L. Bartlett, G. A. Wiseman, N. Padre, A. J. Grillo-López, P. Multani and C. A. White, *J. Clin. Oncol.*, 2002, **20**, 2453–2463.
- 15 A. Imhof, P. Brunner, N. Marincek, M. Briel, C. Schindler, H. Rasch, H. R. Mäcke, C. Rochlitz, J. Müller-Brand and M. A. Walter, *J. Clin. Oncol.*, 2011, **29**, 2416–2423.
- 16 D. J. Kwekkeboom, J. J. Teunissen, W. H. Bakker, P. P. Kooij, W. W. de Herder, R. A. Felders, C. H. van Eijck, J.-P. Esser, B. L. Kam and E. P. Krenning, *J. Clin. Oncol.*, 2005, **23**, 2754–2762.
- 17 J. R. Strosberg, M. E. Caplin, P. L. Kunz, P. B. Ruzsniowski, L. Bodei, A. Hendifar, E. Mittra, E. M. Wolin, J. C. Yao, M. E. Pavel, E. Grande, E. Van Cutsem, E. Seregni, H. Duarte, G. Gericke, A. Bartalotta, M. F. Mariani, A. Demange, S. Mutevelic and E. P. Krenning, *Lancet Oncol.*, 2021, **22**, 1752–1763.
- 18 O. Sartor, J. de Bono, K. N. Chi, K. Fizazi, K. Herrmann, K. Rahbar, S. T. Tagawa, L. T. Nordquist, N. Vaishampayan, G. El-Haddad, C. H. Park, T. M. Beer, A. Armour, W. J. Pérez-Contreras, M. DeSilvio, E. Kpamegan, G. Gericke, R. A. Messmann, M. J. Morris and B. J. Krause, *N. Engl. J. Med.*, 2021, **385**, 1091–1103.
- 19 V. Radchenko, A. Morgenstern, A. R. Jalilian, C. F. Ramogida, C. Cutler, C. Duchemin, C. Hoehr, F. Haddad, F. Bruchertseifer, H. Gausemel, H. Yang, J. A. Osso, K. Washiyama, K. Czerwinski, K. Leufgen, M. Pruszyński, O. Valzdorf, P. Causey, P. Schaffer, R. Perron, S. Maxim, D. S. Wilbur, T. Stora and Y. Li, *J. Nucl. Med.*, 2021, **62**, 1495–1503.
- 20 Y.-S. Kim and M. W. Brechbiel, *Tumor Biol.*, 2012, **33**, 573–590.
- 21 F. D. C. Guerra Liberal, J. M. O'Sullivan, S. J. McMahon and K. M. Prise, *Cancer Biother. Radiopharm.*, 2020, **35**, 404–417.
- 22 C. F. Ramogida and C. Orvig, *Chem. Commun.*, 2013, **49**, 4720–4739.
- 23 M. W. Brechbiel, *Dalton Trans.*, 2007, **43**, 4918–4928.
- 24 S. N. M. Chilla, C. Henoumont, L. V. Elst, R. N. Muller and S. Laurent, *Isr. J. Chem.*, 2017, **57**, 800–808.
- 25 G. J. Stasiuk and N. J. Long, *Chem. Commun.*, 2013, **49**, 2732–2746.
- 26 R. E. Mewis and S. J. Archibald, *Coord. Chem. Rev.*, 2010, **254**, 1686–1712.
- 27 Z. Baranyai, G. Tirsó and F. Rösch, *Eur. J. Inorg. Chem.*, 2020, **2020**, 36–56.
- 28 S. L. Wu and W. D. Horrocks Jr, *J. Chem. Soc., Dalton Trans.*, 1997, **9**, 1497–1502.
- 29 H. Yang, J. J. Wilson, C. Orvig, Y. Li, D. S. Wilbur, C. F. Ramogida, V. Radchenko and P. Schaffer, *J. Nucl. Med.*, 2022, **63**, 5–13.
- 30 A. Roca-Sabio, M. Mato-Iglesias, D. Esteban-Gómez, É. Tóth, A. de Blas, C. Platas-Iglesias and T. Rodríguez-Blas, *J. Am. Chem. Soc.*, 2009, **131**, 3331–3341.
- 31 E. Aluicio-Sarduy, N. A. Thiele, K. E. Martin, B. A. Vaughn, J. Devaraj, A. P. Olson, T. E. Barnhart, J. J. Wilson, E. Boros and J. W. Engle, *Chem. – Eur. J.*, 2020, **26**, 1238–1242.
- 32 N. A. Thiele, V. Brown, J. M. Kelly, A. Amor-Coarasa, U. Jermilova, S. N. MacMillan, A. Nikolopoulou, S. Ponnala, C. F. Ramogida, A. K. H. Robertson, C. Rodríguez-Rodríguez, P. Schaffer, C. Williams Jr, J. W. Babich, V. Radchenko and J. J. Wilson, *Angew. Chem., Int. Ed.*, 2017, **56**, 14712–14717.
- 33 A. P. King, N. T. Gutsche, N. Raju, S. Fayn, K. E. Baidoo, M. M. Bell, C. S. Olkowski, R. E. Swenson, F. I. Lin, S. M. Sadowski, S. S. Adler, N. A. Thiele, J. J. Wilson, P. L. Choyke and F. E. Escorcía, *J. Nucl. Med.*, 2023, **64**, 549–554.
- 34 J. M. Kelly, A. Amor-Coarasa, S. Ponnala, A. Nikolopoulou, C. Williams, N. A. Thiele, D. Schlyer, J. J. Wilson, S. G. DiMaggio and J. W. Babich, *J. Nucl. Med.*, 2019, **60**, 649–655.
- 35 K. N. Bobba, A. P. Bidkar, N. Meher, C. Fong, A. Wadhwa, S. Dhrona, A. Sorlin, S. Bidlingmaier, B. Shuere, J. He, D. M. Wilson, B. Liu, Y. Seo, H. F. VanBrocklin and R. R. Flavell, *J. Nucl. Med.*, 2023, **64**, 1076–1082.
- 36 K. N. Bobba, A. P. Bidkar, A. Wadhwa, N. Meher, S. Drona, A. M. Sorlin, S. Bidlingmaier, L. Zhang, D. M. Wilson, E. Chan, N. Y. Greenland, R. Aggarwal, H. F. VanBrocklin, J. He, J. Chou, Y. Seo, B. Liu and R. R. Flavell, *Theranostics*, 2024, **14**, 1344–1360.
- 37 M. K. Blei, L. Waurick, F. Reissig, K. Kopka, T. Stumpf, B. Drobot, J. Kretschmar and C. Mamat, *Inorg. Chem.*, 2023, **62**, 20699–20709.
- 38 B. J. B. Nelson, S. Ferguson, M. Wuest, J. Wilson, M. J. M. Duke, S. Richter, H. Soenke-Jans, J. D. Andersson, F. Juengling and F. Wuest, *J. Nucl. Med.*, 2022, **63**, 584–590.
- 39 D. S. Abou, N. A. Thiele, N. T. Gutsche, A. Villmer, H. Zhang, J. J. Woods, K. E. Baidoo, F. E. Escorcía, J. J. Wilson and D. L. J. Thorek, *Chem. Sci.*, 2021, **12**, 3733–3742.

- 40 B. J. B. Nelson, J. D. Andersson and F. Wuest, *Nucl. Med. Biol.*, 2022, **110–111**, 59–66.
- 41 F. Reissig, D. Bauer, M. Ullrich, M. Kreller, J. Pietzsch, C. Mamat, K. Kopka, H.-J. Pietzsch and M. Walther, *Pharmaceuticals*, 2020, **13**, 272.
- 42 D. Abou, N. Thiele, A. Villmer, N. Gustche, F. Escorcía, J. Wilson and D. Thorek, *J. Nucl. Med.*, 2020, **61**, 587–587.
- 43 D. J. Fiszbein, V. Brown, N. A. Thiele, J. J. Woods, L. Wharton, S. N. MacMillan, V. Radchenko, C. F. Ramogida and J. J. Wilson, *Inorg. Chem.*, 2021, **60**, 9199–9211.
- 44 A. Hu and J. J. Wilson, *Acc. Chem. Res.*, 2022, **55**, 904–915.
- 45 A. Hu, S. N. MacMillan and J. J. Wilson, *J. Am. Chem. Soc.*, 2020, **142**, 13500–13506.
- 46 A. Hu, E. Aluicio-Sarduy, V. Brown, S. N. MacMillan, K. V. Becker, T. E. Barnhart, V. Radchenko, C. F. Ramogida, J. W. Engle and J. J. Wilson, *J. Am. Chem. Soc.*, 2021, **143**, 10429–10440.
- 47 A. Hu, M. E. Simms, V. Kertesz, J. J. Wilson and N. A. Thiele, *Inorg. Chem.*, 2022, **61**, 12847–12855.
- 48 A. Hu, K. E. Martin, D. Śmiłowicz, E. Aluicio-Sarduy, S. J. Cingoranelli, S. E. Lapi, J. W. Engle, E. Boros and J. J. Wilson, *Eur. J. Inorg. Chem.*, 2023, **26**, e202300457.
- 49 J. Sörensen, D. Sandberg, M. Sandström, A. Wennborg, J. Feldwisch, V. Tolmachev, G. Åström, M. Lubberink, U. Garske-Román, J. Carlsson and H. Lindman, *J. Nucl. Med.*, 2014, **55**, 730–735.
- 50 D. Rosik, A. Orlova, J. Malmberg, M. Altai, Z. Varasteh, M. Sandström, A. E. Karlström and V. Tolmachev, *Eur. J. Nucl. Med. Mol. Imaging*, 2012, **39**, 693–702.
- 51 D. Wild, M. Fani, M. Behe, I. Brink, J. E. F. Rivier, J. C. Reubi, H. R. Maecke and W. A. Weber, *J. Nucl. Med.*, 2011, **52**, 1412–1417.
- 52 T. J. Wadas, E. H. Wong, G. R. Weisman and C. J. Anderson, *Chem. Rev.*, 2010, **110**, 2858–2902.
- 53 R. D. Shannon, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffr., Theor. Gen. Crystallogr.*, 1976, **32**, 751–767.
- 54 L. A. Kloo and M. J. Taylor, *J. Chem. Soc., Dalton Trans.*, 1997, **15**, 2693–2696.
- 55 B. F. T. Cooper, C. G. Andrews and C. L. B. Macdonald, *J. Organomet. Chem.*, 2007, **692**, 2843–2848.
- 56 D. G. Samsonenko, M. N. Sokolov, A. V. Virovets, N. V. Pervukhina and V. P. Fedin, *Eur. J. Inorg. Chem.*, 2001, **2001**, 167–172.
- 57 L. Zhang, F. Wang, X. Zhang, F. Liang, Z. Hu and Y. Wu, *Cryst. Growth Des.*, 2024, **24**, 627–631.
- 58 R. L. Davidovich, A. V. Gerasimenko, E. I. Voit and V. B. Logvinova, *J. Fluorine Chem.*, 2019, **226**, 109343.
- 59 E. T. Clarke and A. E. Martell, *Inorg. Chim. Acta*, 1991, **190**, 37–46.
- 60 C. F. G. C. Geraldes, R. Delgado, A. M. Urbano, J. Costa, F. Jasanada and F. Nepveu, *J. Chem. Soc., Dalton Trans.*, 1995, **13**, 327–335.
- 61 W. P. Cacheris, S. K. Nickle and A. D. Sherry, *Inorg. Chem.*, 1987, **26**, 958–960.
- 62 T. S. Grimes and K. L. Nash, *J. Solution Chem.*, 2014, **43**, 298–313.
- 63 M. Pniok, V. Kubíček, J. Havlíčková, J. Kotek, A. Sabatie-Gogová, J. Plutnar, S. Huclier-Markai and P. Hermann, *Chem. – Eur. J.*, 2014, **20**, 7944–7955.
- 64 S. Liu, J. Pietryka, C. E. Ellars and D. S. Edwards, *Bioconjugate Chem.*, 2002, **13**, 902–913.
- 65 R. A. Fawwaz, T. S. T. Wang, A. Estabrook, J. M. Rosen, M. A. Hardy, P. O. Alderson, S. C. Srivastava, P. Richards and S. Ferrone, *J. Nucl. Med.*, 1985, **26**, 488–492.
- 66 P. Cieslik, M. Kubeil, K. Zarschler, M. Ullrich, F. Brandt, K. Anger, H. Wadepohl, K. Kopka, M. Bachmann, J. Pietzsch, H. Stephan and P. Comba, *J. Am. Chem. Soc.*, 2022, **144**, 21555–21567.
- 67 I. Kopp, P. Cieslik, K. Anger, T. Josephy, L. Neupert, G. Velmurugan, M. Gast, H. Wadepohl, S. A. Brühlmann, M. Walther, K. Kopka, M. Bachmann, H. Stephan, M. Kubeil and P. Comba, *Inorg. Chem.*, 2023, **62**, 20754–20768.
- 68 L. Wharton, M. D. G. Jaraquemada-Peláez, C. Zhang, J. Zeisler, C. Rodríguez-Rodríguez, M. Osooly, V. Radchenko, H. Yang, K.-S. Lin, F. Bénard, P. Schaffer and C. Orvig, *Bioconjugate Chem.*, 2022, **33**, 1900–1921.
- 69 L. Wharton, C. Zhang, J. Zeisler, C. Rodríguez-Rodríguez, M. Osooly, V. Radchenko, H. Yang, K.-S. Lin, F. Bénard, P. Schaffer and C. Orvig, *Bioconjugate Chem.*, 2022, **33**, 2381–2397.
- 70 L. Wharton, H. Yang, M. D. G. Jaraquemada-Peláez, H. Merckens, G. Engudar, A. Ingham, H. Koniar, V. Radchenko, P. Kunz, P. Schaffer, F. Bénard and C. Orvig, *J. Med. Chem.*, 2023, **66**, 13705–13730.
- 71 L. Wharton, C. Zhang, H. Yang, J. Zeisler, V. Radchenko, C. Rodríguez-Rodríguez, M. Osooly, B. O. Patrick, K.-S. Lin, F. Bénard, P. Schaffer and C. Orvig, *Bioconjugate Chem.*, 2022, **33**, 505–522.
- 72 D. Fiaccabrino, T. Masvikeni, M. D. G. Jaraquemada-Peláez, C. Orvig and P. Schaffer, *Inorg. Chem.*, 2024, **63**, 13911–13923.
- 73 M. E. Simms, Z. Li, M. M. Sibley, A. S. Ivanov, C. M. Lara, T. C. Johnstone, V. Kertesz, A. Fears, F. D. White, D. L. J. Thorek and N. A. Thiele, *Chem. Sci.*, 2024, **15**, 11279–11286.
- 74 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 3–8.
- 75 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 76 P. Müller, *Crystallogr. Rev.*, 2009, **15**, 57–83.
- 77 O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- 78 P. Gans, A. Sabatini and A. Vacca, *Ann. Chim.*, 1999, **89**, 45–49.
- 79 B. L. McNeil, K. J. Kadassery, A. W. McDonagh, W. Zhou, P. Schaffer, J. J. Wilson and C. F. Ramogida, *Inorg. Chem.*, 2022, **61**, 9638–9649.
- 80 C. Müller, H. Struthers, C. Winiger, K. Zhernosekov and R. Schibli, *J. Nucl. Med.*, 2013, **54**, 124–131.

- 81 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr, J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16 Rev. C.01* 2016.
- 82 J.-D. Chai and M. Head-Gordon, *J. Chem. Phys.*, 2008, **128**, 084106.
- 83 J.-D. Chai and M. Head-Gordon, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6615–6620.
- 84 P. C. Hariharan and J. A. Pople, *Theor. Chim. Acta*, 1973, **28**, 213–222.
- 85 W. J. Hehre, R. Ditchfield and J. A. Pople, *J. Chem. Phys.*, 1972, **56**, 2257–2261.
- 86 W. R. Wadt and P. J. Hay, *J. Chem. Phys.*, 1985, **82**, 284–298.
- 87 A. V. Marenich, C. J. Cramer and D. G. Truhlar, *J. Phys. Chem. B*, 2009, **113**, 6378–6396.