








Cite this: *Chem. Sci.*, 2024, 15, 13550 All publication charges for this article have been paid for by the Royal Society of Chemistry

A versatile fluorinated azamacrocyclic chelator enabling ^{18}F PET or ^{19}F MRI: a first step towards new multimodal and smart contrast agents†

Charline Sire, ^a Vincent Meneyrol,^b Nathalie Saffon-Merceron, ^c
Enzo Terreno, ^d Francesca Garello, ^d Lorenzo Tei, ^e Emmanuelle Jestin,^b
Raphaël Tripier ^a and Thibault Troadec ^{*a}

Macrocyclic chelators play a central role in medical imaging and nuclear medicine owing to their unparalleled metal cation coordination abilities. Their functionalization by fluorinated groups is an attractive design, to combine their properties with those of ^{18}F for Positron Emission Tomography (PET) or natural ^{19}F for Magnetic Resonance Imaging (MRI), and access potential theranostic or smart medical imaging probes. For the first time, a compact fluorinated macrocyclic architecture has been synthesized, based on a cyclen chelator bearing additional pyridine coordinating units and simple methyltrifluoroborate prosthetic groups. This ligand and its corresponding model Zn(II) complex were investigated to evaluate the ^{18}F -PET or ^{19}F MRI abilities provided by this novel molecular structure. The chelator and the complex were obtained via a simple and high-yielding synthetic route, present excellent solvolytic stability of the trifluoroborate groups at various pH, and provide facile late-stage ^{18}F -radiolabeling (up to 68% radiochemical yield with high activity) as well as a satisfying detection limit for ^{19}F MRI imaging (low mM range).

Received 30th April 2024
Accepted 17th July 2024

DOI: 10.1039/d4sc02871f

rsc.li/chemical-science

Introduction

Fluorine is a ubiquitous element in medical imaging with its two main isotopes, natural ^{19}F and radioactive ^{18}F , that allow Magnetic Resonance Imaging (MRI) and Positron Emission Tomography (PET), respectively. On the one hand, ^{18}F is the most used and readily available radionuclide for PET, with a wide range of radiolabeling strategies currently available,^{1,2} as it has the advantages of a biologically relevant radioactive half-life ($t_{1/2} = 110$ min), low radiotoxicity and low positron (β^+) emission energy resulting in high resolution images. On the other hand, ^{19}F -MRI presents the intrinsic advantages of the

natural abundance of ^{19}F (100%) and its good sensitivity compared to other nuclei (second best behind ^1H used in MRI). Moreover, the lack of *in vivo* background owing to the absence of fluorine in the body soft tissues allows direct quantitative detection of the tracer, in opposition to conventional ^1H MRI that relies on the water content of the body.^{3,4} However this also leads to a low overall sensitivity of the ^{19}F MRI technique that is currently tackled by two distinct strategies when it comes to molecular probes: high fluorine content of the tracer (ex: perfluorinated alkanes)⁴ or combination with paramagnetic metals or lanthanides that lower the relaxation time of the sensors.⁵

Concurrently, in the field of molecular probes for medical imaging and therapy, saturated polyazamacrocycles, such as cyclen (1,4,7,10-tetraazacyclododecane), cyclam (1,4,8,11-tetraazacyclotetradecane) and tacn (1,4,7-triazacyclononane) are cornerstone for the complexation of metallic and lanthanide cations for MRI,⁶ PET,^{7,8} Single-Photon Emission Computed Tomography (SPECT)⁹ or internal Radiotherapy,^{7,8} owing to their exceptional coordination properties and *in vivo* inertness of their complexes. In that context, fluorinated azamacrocycles are highly attractive as they can combine, within a single molecular architecture, the properties of metals or radiometals coordinated within their cavity with the ones of ^{19}F or ^{18}F . In particular, two major applications have been targeted recently: (i) complexes combining radiometals (α or β^- emitters) for radiotherapy and prosthetic ^{18}F -radiolabeled units for diagnosis, leading to so-called theranostic tracers that merge

^aUniv. Brest, UMR CNRS 6521 CEMCA, 6 Avenue Victor Le Gorgeu, 29200 Brest, France. E-mail: Thibault.troadec@univ-brest.fr

^bCyclotron Réunion Océan Indien CYROI, 2 rue Maxime Rivière 97490 Sainte-Clotilde, France

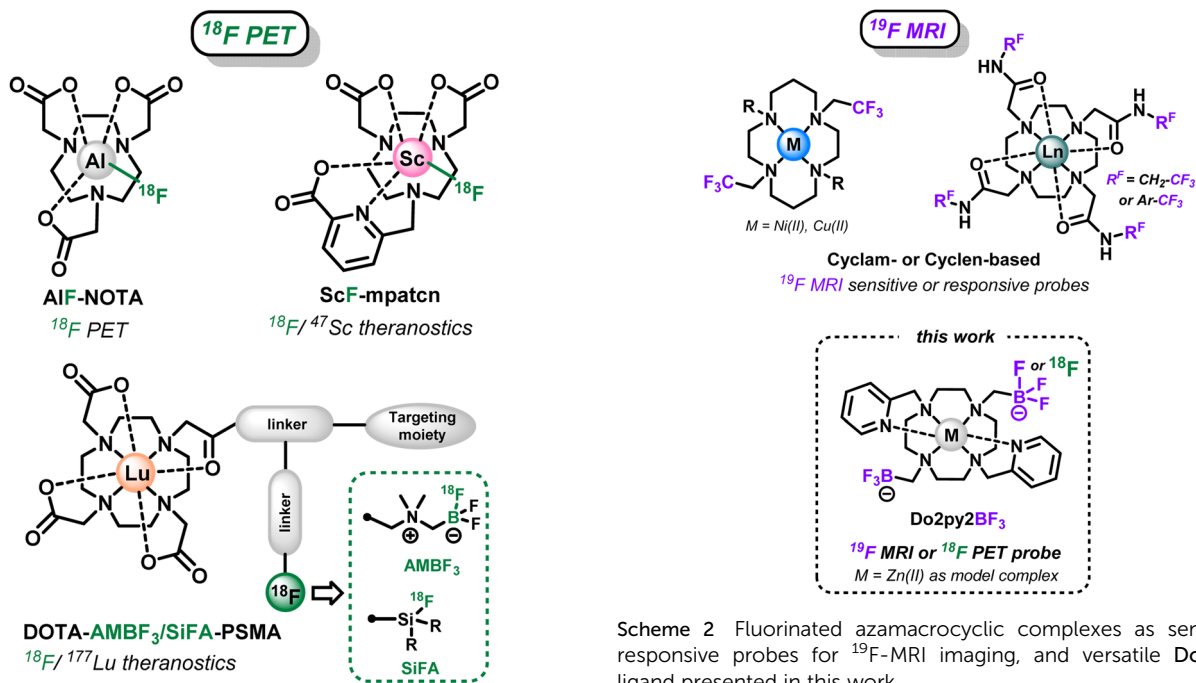
^cInstitut de Chimie de Toulouse (UAR 2599), 118 route de Narbonne, 31062 Toulouse Cedex 9, France

^dDepartment of Molecular Biotechnology and Health Sciences, University of Turin, Piazza Nizza 44/bis, 10126 Turin, Italy

^eDipartimento di Scienze ed Innovazione Tecnologica, Università del Piemonte Orientale "Amedeo Avogadro", Viale T. Michel 11, 15121 Alessandria, Italy

† Electronic supplementary information (ESI) available: Experimental procedures, spectroscopic data for characterization and stability studies of the new compounds, and experimental details for ^{18}F labeling and ^{19}F magnetic resonance experiments. CCDC 2314190–2314192. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4sc02871f>



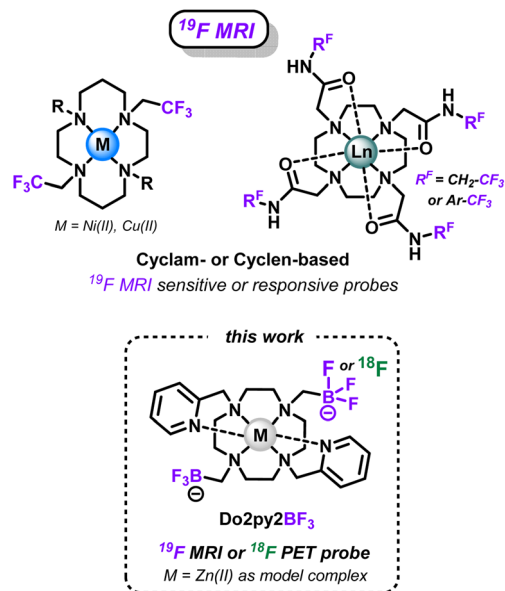


Scheme 1 Fluorinated azamacrocyclic complexes for ^{18}F -PET imaging and theranostics.

therapeutic and diagnostic modalities in a single molecule and provide identical biodistribution *in vivo*; and (ii) complexes of paramagnetic cations with natural ^{19}F pendants as sensitive or responsive ^{19}F MRI probes.

Several fluorinated polyazacycloalane scaffolds have thus been described for these two distinct objectives in recent years. First, azamacrocycles were used in ^{18}F -PET as an alternative to conventional ^{18}F -labeling (*via* C–F bonds formation)^{10,11} for the tagging of biomolecules. In this case, the macrocyclic cavity of tacn-based chelators was used to accommodate metal- ^{18}F synthons: with Al(III) over a decade ago (AIF-NOTA, Scheme 1),^{12–14} and lately with Ga(III) and Fe(III).^{15–18} Linear ligands, such as ResCa and analogues, have also been reported since to allow such radiolabeling at low temperature (<37 °C).^{19–23} However in those cases, the chelating cavity is occupied by innocent metal cations bearing no additional property. Only very recently Boros *et al.* went a step further and developed the corresponding theranostic version, using the Sc(III)–F synthon that can allow the ^{47}Sc (β^- radiotherapy)/ ^{18}F (PET imaging) theranostic couple.²⁴ Nevertheless, the scope of this strategy is limited in terms of possible metal cations and corresponding radiometals as the strength of the M–F interaction is not sufficient with most relevant metallic cations, as already observed for instance with lanthanides.^{25,26}

In a second strategy, the well-known DOTA chelator ((1,4,7,10-tetraazacyclododecane-1,4,7,10-tetrayl)tetraacetic acid), that can coordinate a wide range of metal cations and ^{177}Lu (β^- emitter) in particular, has been functionalized by branched side chains allowing grafting on a targeting moiety and installation of fluoroborate or fluorosilane prosthetic groups.^{27,28} These are particularly attractive in this context, as they allow easy late-stage ^{18}F -labeling *via* $^{19}\text{F}/^{18}\text{F}$ isotopic



Scheme 2 Fluorinated azamacrocyclic complexes as sensitive or responsive probes for ^{19}F -MRI imaging, and versatile Do2py2BF₃ ligand presented in this work.

exchange processes (Scheme 1),^{29–31} as an alternative to sometimes tedious C– ^{18}F bond forming reactions.^{10,11} However, the organic frameworks attached to the chelator in this case require challenging multistep syntheses, and result in large branched structures that can be detrimental to the recognition properties of the targeting unit and biodistribution of the radiopharmaceuticals. Therefore, more compact architecture are highly desirable.

Concerning the ^{19}F MRI modality, small trifluoromethyl units have been efficiently used in sensitive or responsive probes based on azamacrocyclic complexes. First, trifluoromethylated cyclen-based chelators (Scheme 2) have proven excellent to coordinate paramagnetic lanthanide cations (Tb^{3+} , Dy^{3+} , Ho^{3+} , Er^{3+} in particular) that decrease ^{19}F relaxation times and drastically enhance the probes sensitivity.^{5,32–34} Similarly, CF_3 -appended cyclam chelators (Scheme 2) have been used with nickel(II) cations as sensitive probes,^{35,36} or as on-off responsive sensors with redox-active cations (Co^{2+} , Cu^{2+}) that can modulate the ^{19}F MRI signal upon modification of their oxidation state by external stimuli.^{37,38} However, the lipophilic nature of such trifluoromethylated groups can be detrimental to applications in aqueous media and *in vivo* in particular.³²

Therefore, we report in this study a novel compact cyclen-based architecture, bearing methyltrifluoroborate units directly supported by the macrocyclic amines (Do2py2BF₃, Scheme 2), to target two different applications with the same molecular topology: (i) allow easy ^{18}F -radiolabeling *via* $^{19}\text{F}/^{18}\text{F}$ isotopic exchange to allow future theranostic combinations with relevant radiometals; and (ii) present suitable ^{19}F MRI signal, which has never been explored in the case of trifluoroborates, that could be used in responsive or sensitive imaging agents upon coordination with relevant non-radioactive paramagnetic cations. Noteworthy in the field (see AIF-NOTA or ScF-mptacn, Scheme 1), the



chemistry of such chelators is first developed without the introduction of grafting functions and targeting molecules that will be necessary for future *in vivo* targeted applications. Therefore, the bispyridyl-cyclen scaffold has been selected in this proof-of-concept study for its ease of preparation and ubiquitous coordination properties.^{39–41} Herein, the synthesis and characterization of this novel ligand architecture is presented, as well as the corresponding zinc(II) complex, a convenient diamagnetic model complex that allowed easy NMR characterizations and simple stability studies. The ability of the scaffold to undergo ¹⁸F-labeling, as known for other types of trifluoroborates, was investigated on this new structure. Then the potential of BF₃ units as ¹⁹F MRI reporters was also investigated for the first time on the model Zn(II) complex. Finally, a particular focus was given to the study of the solvolytic stability of the new trifluoroborated scaffolds in water at different pH, which is a crucial parameter to envisage future applications for this new type of chelators.

Results & discussion

Synthesis and characterization

Ligand H₂Do2py2BF₃ was prepared from previously described Do2py,⁴⁰ through a 2-step synthetic procedure adapted from Perrin *et al.* for the installation of methyltrifluoroborate functional groups *via* fluorination of a pinacolborane intermediate with potassium bifluoride (Scheme 3).⁴²

The new macrocycle was fully characterized by means of High-Resolution Mass Spectrometry (HRMS) and multinuclear NMR. In 1 : 1 D₂O/CD₃CN (pH 4), the ligand exhibits broad singlets in ¹⁹F ($\delta = -135.6$ ppm) and ¹¹B ($\delta = 2.44$ ppm) NMR and a diagnostic quadruplet for the α -methylene group in ¹H NMR ($\delta = 1.80$ ppm, ³J_{H-F} = 5.2 Hz) (see ESI† for spectral data). In ¹³C NMR, this methylenic carbon could not be directly observed owing to multiplicity and broadness induced by neighboring boron and fluorine nuclei, but was assigned thanks to 2D ¹H–¹³C HSQC NMR (47.8 ppm). Single crystals were grown from a water/acetonitrile mixture (1 : 1, pH 4) revealing the neutral

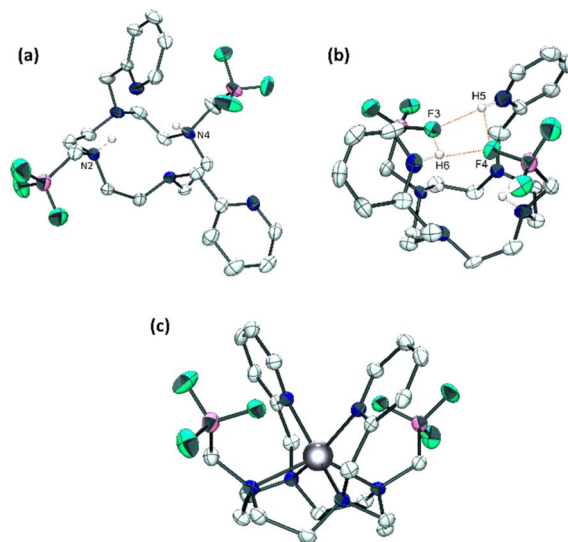
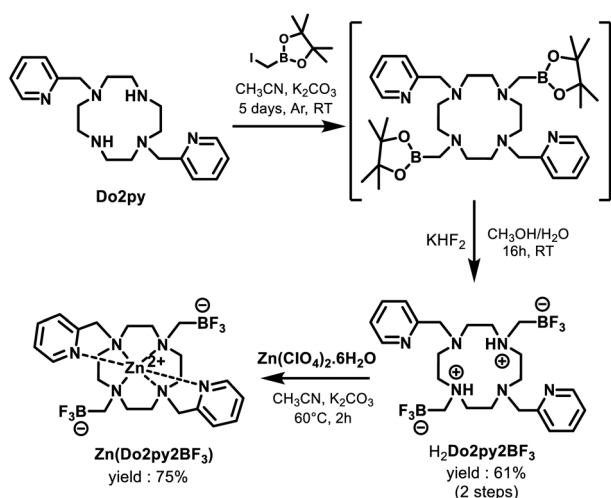


Fig. 1 X-Ray diffraction structures of (a) H₂Do2py2BF₃ ligand at pH 4, (b) H₄Do2py2BF₃ at pH 1 and (c) Zn(Do2py2BF₃) complex (c). Ellipsoids are drawn with 50% probability. H atoms omitted for clarity, except on ammonium groups of H₂Do2py2BF₃ and H₄Do2py2BF₃.

H₂Do2py2BF₃ ligand with both nitrogen atoms holding the BF₃ pendants (N2 and N4, Fig. 1a) that are protonated.⁴³ This feature is classical for cyclen-based ligands at this pH, as two amines in *trans* N1–N3 positions generally have pK_as above 9.⁴⁴

Single crystals were also grown from an acidic aqueous solution (pH 1), revealing a different structure with the ligand under the form H₄Do2py2BF₃²⁺ displaying additional protonation of the two pyridine units (H5 and H6, Fig. 1b) creating a H-bonding network with fluorine atoms (F3 and F4) from both trifluoroborate pendants (*d*_{F4–H5} and *d*_{F3–H6} of 1.89 and 1.91 Å respectively). This constrained geometry is also present in solution at acidic pH, as evidenced by a set of 3 broad signals on the ¹⁹F spectrum from –136 to –140 ppm (D₂O, pH 1), close to the sole chemical shift observed at pH 4 (–135.6 ppm). Similarly, ¹H NMR spectrum reveals the different symmetry of the molecule with a complex multiplet centered at 4.16 ppm accounting for the four methylenic pyridyl protons (*vs.* sharp singlet at 4.49 ppm at pH 4), different multiplicity of the aromatic protons and a set of three broad signals between 1.90 and 2.30 ppm for the methylenic protons linked to the trifluoroborate units.

The corresponding zinc(II) complex was then prepared, from an equimolar mixture of H₂Do2py2BF₃ and zinc(II) perchlorate in acetonitrile at 60 °C, with potassium carbonate as a base, and purified by crystallization in a water/acetonitrile mixture (3 : 7). Owing to its limited solubility in water at high concentrations, NMR analysis was carried out in a 1 : 1 D₂O/CD₃CN mixture (pH 6). At RT, the ¹H spectrum presents very broad signals for the macrocyclic protons, that could however be resolved at 333 K (Fig. S11, ESI†), and ¹⁹F NMR reveals a broad singlet at –137.4 ppm, very close to that of free ligand, as a first indication of the innocent nature of the trifluoroborate pendants upon coordination of the metal. More insight was gained by the structure revealed by X-ray diffraction analysis on single crystals



Scheme 3 Synthesis of H₂Do2py2BF₃ ligand and Zn(Do2py2BF₃) complex.



of the complex grown in CH₃CN/H₂O (7 : 3) solution (Fig. 1c). Zinc(II) cation lies slightly above the cyclen plane (distance with N1–N2–N3–N4 centroid = 0.999 Å, Fig. S13, ESI†) and adopts a distorted trigonal prismatic geometry similar to the Zn(Do2py)²⁺ complex reported in the literature,⁴⁰ highlighting again the innocent behaviour of the trifluoroborated groups in the coordination sphere of the metal. Only the Zn–N bonds with cyclen nitrogen atoms are somewhat longer in this complex (by up to 0.07 Å, Fig. S13, ESI†), probably owing to the different steric bulk on two nitrogen atoms from the ring (N–CH₂–BF₃ vs. NH in Do2py).

Solvolytic stability

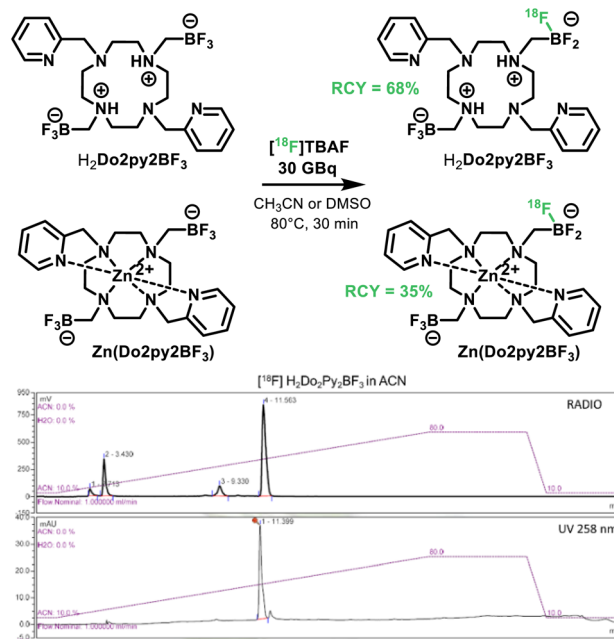
The main limitation of trifluoroborate prosthetic groups for practical use in [¹⁸F]-PET radiotracers can be their lack of stability towards solvolysis, *via* the release of free fluoride and formation of boronic acid derivatives, which has been well-documented by Perrin *et al.*^{45,46}

The solvolytic stability of the new azamacrocyclic motifs was thus investigated for the free ligand and the Zn complex by ¹⁹F NMR, at a concentration of 3 × 10^{−2} M in buffered aqueous solutions at pH 2.0, 7.3 and 9.7 (Fig. S15–S20, ESI†). In these conditions, Do2py2BF₃ demonstrated exceptional robustness with no degradation observed in any of the three pH conditions, after up to 3 days at 25 °C (Table 1). In particular, this observation is in contrast with the closest reported ammonium-based structure (Et₂NH⁺CH₂BF₃) that displays a half-life of 66 hours at physiological pH.⁴⁶ This enhanced stability in Do2py2BF₃ is probably due to the higher basicity of the cyclen tertiary amines providing a poorly labile ammonium proton stabilizing the zwitterionic structure with the trifluoroborate group. On the zinc(II) complex, the stability is slightly different, owing to the loss of the ammonium protons to accommodate the metallic cation. First, at pH 2.0, a fast decoordination of the cation was evidenced by ¹⁹F NMR (*t*_{1/2(Zn)} < 5 min, Fig. S18, ESI†), which is common with such highly basic polyamines at acidic pH. However, no degradation of the BF₃ units was observed as only the ¹⁹F signals of the free ligand were recovered. At neutral and basic pH, no release of the metal occurred, demonstrating the strong coordination ability of the ligand. However, very slow solvolysis of the BF₃ groups appeared (Fig. S19–S20,† ESI). As previously discussed in the literature by Perrin and others,^{45,47} the kinetics of solvolysis follow a pseudo-first order kinetic rate,

with a rate-determining step corresponding to the loss of the first fluoride ion, and the other intermediates towards fully hydrolyzed boronic acid being fast-lived. Very long half-lives (985 and 400 hours at pH 7.3 and 9.7 respectively) could be calculated here for the new BF₃ pendants, that are perfectly suited to consider further use in the context of ¹⁸F PET imaging when compared to the radioactive half-life of ¹⁸F (110 min) or the biodistribution and clearance of radiotracers (from minutes to a few days). Indeed, the stability of these pendants even surpasses the one of the most stable similar trifluoroborate functions described so far in the literature in these pH ranges (366 h for AMBF₃ groups at pH 7.5).

¹⁸F-radiolabeling

Furthermore the ability of the new molecule for easy ¹⁸F-radiolabeling was evaluated on both H₂Do2py2BF₃ ligand and Zn(Do2py2BF₃) (Scheme 4). Radiosyntheses were carried out on a TRACERlab FXFN module (GE) (Fig. S21, ESI†). At the end of bombardment (EOB), the [¹⁸F] fluoride produced by the cyclotron was delivered to the automate and trapped on a pre-conditioned QMA Sep-Pak cartridge to remove [¹⁸O]-water. The elution of the activity (30 GBq at EOB) into the reactor was performed with a solution of TBABr (16 mg in 0.5 ml H₂O) to form [¹⁸F]tetrabutylammonium fluoride ([¹⁸F]TBAF). Acetonitrile was added for the azeotropic evaporation under vacuum and helium flow. Solutions containing 1 mg of H₂Do2py2BF₃ in acetonitrile or Zn(Do2py2BF₃) in DMSO were added into the reactor for a 30 min incubation at 80 °C. It is worth noting that these solvents are not desired for further application *in vivo*, but were used herein as the ligand and complex are not fully soluble in water. However, solubility in aqueous media should be



Scheme 4 ¹⁸F-radiolabeling of H₂Do2py2BF₃ and Zn(Do2py2BF₃) in acetonitrile, and corresponding HPLC (UV) and radioHPLC traces for H₂Do2py2BF₃.

Table 1 Solvolysis data of Do2py2BF₃ ligand and Zn(Do2py2BF₃) complex

Compound	pH	<i>t</i>	% intact BF ₃	<i>t</i> _{1/2}
Do2py2BF ₃	2.0	72 h	100%	—
	7.3	72 h	100%	—
	9.7	72 h	100%	—
Zn(Do2py2BF ₃) ²⁺	2.0	8 days	100% (90% Zn release)	—
	7.3	22 h	89%	985 h
		13 days	77%	
	9.7	18 h	90%	400 h
		11 days	63%	



achieved for future applications upon grafting on water soluble biomolecules, and aqueous ^{18}F -labelling of trifluoroborates has been well documented.^{48,49}

RadioChemical Yields (RCY) were subsequently measured *via* radioHPLC analyses, with UV traces of the “cold” ligand or zinc complex as references (Fig. S22–S23, ESI†). In these conditions, highly satisfactory RCY of up to 68% was obtained for the ligand $\text{H}_2\text{Do2py2BF}_3$. RCY for the $\text{Zn}(\text{Do2py2BF}_3)$ complex is somewhat lower, probably owing to its lower solubility in the reaction medium, but is acceptable to envisage future applications in the field. Indeed, these RCY are excellent when compared to analogous isotopic exchange with trifluoroborates in similar conditions (15–30%)^{42,50} or labeling *via* metal-fluoride synthons (30–45% with Al,⁵¹ 65% with Sc (ref. 24)). In addition to these conversions, labeled compounds could also be easily purified on semi-preparative HPLC with a Gemini C_{18} Column and formulated as injectable aqueous solutions after solubilization in NaCl solutions on C_{18} Sep-Pak cartridges within the automated system, which is one of the main advantages of such facile isotopic exchange on BF_3 units. For the $\text{H}_2\text{Do2py2BF}_3$ radiolabeling, 30.6 $\text{GBq } \mu\text{mol}^{-1}$ at the injection time were associated to the desired product whereas 7.8 $\text{GBq } \mu\text{mol}^{-1}$ were obtained for the radiolabeled $\text{Zn}(\text{Do2py2BF}_3)$. These results on the molar activities are good enough for further *in vivo* studies and consistent with the reference publications (around 40–110 $\text{GBq } \mu\text{mol}^{-1}$ for high specific activity).⁴⁸

^{19}F -magnetic resonance

To investigate the potential of trifluoroborate units as ^{19}F MRI probes, relaxivity studies were carried out at a 7.1 T field, on

a $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (1 : 1) solution of $\text{Zn}(\text{Do2py2BF}_3)$ at a concentration of 0.032 M in complex (*i.e.* ^{19}F concentration of 0.192 M owing to the 6 fluorine atoms per molecule). A single peak in ^{19}F modality was detectable at -137 ppm. Longitudinal (T_1) relaxivity was measured with an inversion recovery sequence, while the transversal (T_2) relaxivity was measured with a CPMG sequence, leading to values of $T_1 = 383.7 \pm 0.01$ ms and $T_2 = 268 \pm 0.02$ ms. These are reasonably short relaxation times compared to corresponding ligands bearing trifluoromethyl groups (*i.e.* two to three times lower T_1 than cyclam methylene- CF_3 analogue with diamagnetic $\text{Cu}(\text{i})$ cation),^{5,38} that allowed a large number of averages to be accumulated in a short time. Then a phantom was prepared with variable ^{19}F concentrations ranging from 1 mM to 150 mM (Fig. 2). Samples with ^{19}F concentration of 50, 100 and 150 mM were clearly detectable already after 10 min of acquisition (corresponding $\text{Zn}(\text{Do2py2BF}_3)$ concentration of 8.3, 16.7 and 25 mM, respectively). After 45 min of acquisition, samples with 15 and 25 mM ^{19}F were barely visible so a different phantom (25–150 mM) was used for detection limit measurement with 10, 45 and 60 minutes acquisition (Fig. 2). From these data, a detection limit between 50 and 75 mM in ^{19}F was observed at $t = 10$ minutes, and of 40 mM in ^{19}F (6.7 mM in complex) could be determined for acquisition times of 45 and 60 minutes. This value is clearly in the range of perfluorocarbons used in preclinical and clinical studies.⁵²

Conclusions

In conclusion, we have successfully prepared the first example of a small trifluoroborated macrocyclic chelator, through a practical synthetic procedure and easy purifications. This ligand presents suitable stability of the BF_3 units, which is the usual limitation of such functions for *in vivo* applications, and a spectator character of these groups upon coordination to $\text{Zn}(\text{II})$ in the corresponding model complex. This architecture subsequently allowed easy late-stage ^{18}F radiolabeling through simple and fast isotopic exchange with great RCY, and trifluoroborates have also been used for the first time as ^{19}F MRI reporters. These first results suggest that such BF_3 -appended azamacrocycles are a new very promising class of chelators to envisage both theranostic applications with the ^{18}F PET modality and suitable radiometal coordination design, and their use as sensitive or responsive contrast agents when using the ^{19}F MRI modality. Following this first step, ligand optimization will now be tackled for the various metals and radiometals relevant for these two distinct applications.

Data availability

CCDC 2314190 ($\text{H}_2\text{Do2py2BF}_3$ ligand), 2314191 ($\text{H}_4\text{Do2py2BF}_3^{2+}$ ligand) and 2314192 ($\text{Zn}(\text{Do2py2BF}_3)$) contain the ESI† for this paper. The datasets supporting this article have been uploaded as part of the ESI†. Additional data can be provided upon reasonable request to Thibault.troade@univ-brest.fr.

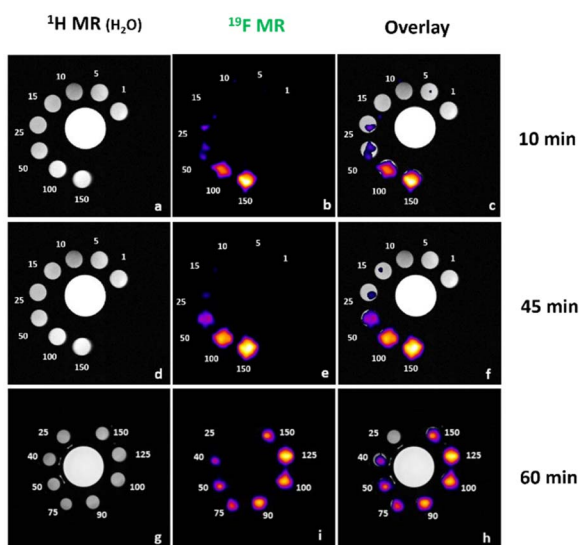


Fig. 2 ^{19}F -Magnetic resonance of $\text{Zn}(\text{Do2py2BF}_3)$ in acetonitrile/water (1 : 1) at 3.2×10^{-2} M. Measurements were conducted at $t = 10$ min (panels a–c), 45 min (panels d–f) and 60 min (panels g–i). For each timepoint, control ^1H MR (H_2O signal) is displayed in left column, ^{19}F signal in the center column, and overlay in the right column. ^{19}F concentration (mM) is indicated next to the corresponding sample.



Author contributions

Conceptualization and funding acquisition by TT; investigation by CS, TT, VM, NSM and FG; supervision by TT, RT, LT, ET; original draft by TT; review and editing by all authors.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work has been supported by CNRS, University of Brest, the "Fondation ARC pour la recherche sur le cancer" (program Macrofluor, TT) and ANR (JCJC program Smartfluor, TT). The authors would like to thank Cyril Colas from the "Fédération de Recherche" ICOA/CBM (FR2708) for HRMS analysis and Muriel Escadeillas from CRMPO (Rennes) for elemental analysis.

Notes and references

- 1 D. van der Born, A. Pees, A. J. Poot, R. V. A. Orru, A. D. Windhorst and D. J. Vugts, Fluorine-18 labelled building blocks for PET tracer synthesis, *Chem. Soc. Rev.*, 2017, **46**, 4709–4773.
- 2 L. C. Erin, N. S. Megan, L. Ryan, H. Raphael and J. H. S. Peter, Radiosyntheses using Fluorine-18: The Art and Science of Late Stage Fluorination, *Curr. Top. Med. Chem.*, 2014, **14**, 875–900.
- 3 T. Guden-Silber, S. Temme, C. Jacoby and U. Flogel, in *Preclinical MRI*, Springer, 2018, pp. 235–257.
- 4 J. Ruiz-Cabello, B. P. Barnett, P. A. Bottomley and J. W. M. Bulte, Fluorine (19F) MRS and MRI in biomedicine, *NMR Biomed.*, 2011, **24**, 114–129.
- 5 K. L. Peterson, K. Srivastava and V. C. Pierre, Fluorinated Paramagnetic Complexes: Sensitive and Responsive Probes for Magnetic Resonance Spectroscopy and Imaging, *Front. Chem.*, 2018, **6**, 160.
- 6 Y.-D. Xiao, R. Paudel, J. Liu, C. Ma, Z.-S. Zhang and S.-K. Zhou, MRI contrast agents: Classification and application (Review), *Int. J. Mol. Med.*, 2016, **38**, 1319–1326.
- 7 E. W. Price and C. Orvig, Matching chelators to radiometals for radiopharmaceuticals, *Chem. Soc. Rev.*, 2013, **43**, 260–290.
- 8 T. I. Kostelnik and C. Orvig, Radioactive Main Group and Rare Earth Metals for Imaging and Therapy, *Chem. Rev.*, 2019, **119**, 902–956.
- 9 D. Sneddon and B. Cornelissen, Emerging chelators for nuclear imaging, *Curr. Opin. Chem. Biol.*, 2021, **63**, 152–162.
- 10 O. Jacobson, D. O. Kiesewetter and X. Chen, Fluorine-18 Radiochemistry, Labeling Strategies and Synthetic Routes, *Bioconjugate Chem.*, 2015, **26**, 1–18.
- 11 V. Rosecker, C. Denk, M. Maurer, M. Wilkovich, S. Mairinger, T. Wanek and H. Mikula, Cross-Isotopic Bioorthogonal Tools as Molecular Twins for Radiotheranostic Applications, *ChemBioChem*, 2019, **20**, 1530–1535.
- 12 W. J. McBride, R. M. Sharkey, H. Karacay, C. A. D'Souza, E. A. Rossi, P. Laverman, C.-H. Chang, O. C. Boerman and D. M. Goldenberg, A Novel Method of 18F Radiolabeling for PET, *J. Nucl. Med.*, 2009, **50**, 991–998.
- 13 P. Laverman, W. J. McBride, R. M. Sharkey, A. Eek, L. Joosten, W. J. G. Oyen, D. M. Goldenberg and O. C. Boerman, A Novel Facile Method of Labeling Octreotide with 18F-Fluorine, *J. Nucl. Med.*, 2010, **51**, 454–461.
- 14 W. J. McBride, C. A. D'Souza, R. M. Sharkey, H. Karacay, E. A. Rossi, C.-H. Chang and D. M. Goldenberg, Improved 18F Labeling of Peptides with a Fluoride-Aluminum-Chelate Complex, *Bioconjugate Chem.*, 2010, **21**, 1331–1340.
- 15 R. Bhalla, C. Darby, W. Levason, S. K. Luthra, G. McRobbie, G. Reid, G. Sanderson and W. Zhang, Triaza-macrocyclic complexes of aluminium, gallium and indium halides: fast 18F and 19F incorporation *via* halide exchange under mild conditions in aqueous solution, *Chem. Sci.*, 2013, **5**, 381–391.
- 16 R. Bhalla, W. Levason, S. K. Luthra, G. McRobbie, G. Sanderson and G. Reid, Radiofluorination of a Pre-formed Gallium(III) Aza-macrocyclic Complex: Towards Next-Generation Positron Emission Tomography (PET) Imaging Agents, *Chem.–Eur. J.*, 2015, **21**, 4688–4694.
- 17 F. M. Monzittu, I. Khan, W. Levason, S. K. Luthra, G. McRobbie and G. Reid, Rapid Aqueous Late-Stage Radiolabelling of [GaF3(BnMe2-tacn)] by ¹⁸F/¹⁹F Isotopic Exchange: Towards New PET Imaging Probes, *Angew. Chem., Int. Ed.*, 2018, **57**, 6658–6661.
- 18 D. E. Runacres, V. K. Greenacre, J. M. Dyke, J. Grigg, G. Herbert, W. Levason, G. McRobbie and G. Reid, Synthesis, Characterization, and Computational Studies on Gallium(III) and Iron(III) Complexes with a Pentadentate Macrocyclic bis-Phosphinate Chelator and Their Investigation As Molecular Scaffolds for 18F Binding, *Inorg. Chem.*, 2023, **62**(50), 20844–20857.
- 19 A. M. S. Musthakahmed, E. Billaud, G. Bormans, F. Cleeren, J. Lecina and A. Verbruggen, *WO2016065435*, 2016.
- 20 F. Cleeren, J. Lecina, E. M. F. Billaud, M. Ahamed, A. Verbruggen and G. M. Bormans, New Chelators for Low Temperature Al18F-Labeling of Biomolecules, *Bioconjugate Chem.*, 2016, **27**, 790–798.
- 21 F. Cleeren, J. Lecina, M. Ahamed, G. Raes, N. Devoogdt, V. Caveliers, P. McQuade, D. J. Rubins, W. Li, A. Verbruggen, C. Xavier and G. Bormans, Al18F-Labeling Of Heat-Sensitive Biomolecules for Positron Emission Tomography Imaging, *Theranostics*, 2017, **7**, 2924–2939.
- 22 S. Schmitt and E. Moreau, Radiochemistry with {Al¹⁸F}²⁺: Current status and optimization perspectives for efficient radiofluorination by complexation, *Coord. Chem. Rev.*, 2023, **480**, 215028.
- 23 L. Russelli, J. Martinelli, F. De Rose, S. Reder, M. Herz, M. Schwaiger, W. Weber, L. Tei and C. D'Alessandria, Room Temperature Al18F Labeling of 2-Aminomethylpiperidine-Based Chelators for PET Imaging, *ChemMedChem*, 2020, **15**, 284–292.
- 24 J. N. Whetter, B. A. Vaughn, A. J. Koller and E. Boros, An Unusual Pair: Facile Formation and *In Vivo* Validation of



- Robust Sc–18F Ternary Complexes for Molecular Imaging, *Angew. Chem., Int. Ed.*, 2022, **61**, e202114203.
- 25 L. M. P. Lima, A. Lecointre, J.-F. Morfin, A. de Blas, D. Visvikis, L. J. Charbonnière, C. Platas-Iglesias and R. Tripier, Positively Charged Lanthanide Complexes with Cyclen-Based Ligands: Synthesis, Solid-State and Solution Structure, and Fluoride Interaction, *Inorg. Chem.*, 2011, **50**, 12508–12521.
- 26 T. Liu, A. Nonat, M. Beyler, M. Regueiro-Figueroa, K. Nchimi Nono, O. Jeannin, F. Camerel, F. Debaene, S. Cianférani-Sanglier, R. Tripier, C. Platas-Iglesias and L. J. Charbonnière, Supramolecular Luminescent Lanthanide Dimers for Fluoride Sequestering and Sensing, *Angew. Chem., Int. Ed.*, 2014, **126**, 7387–7391.
- 27 M. L. Lepage, H.-T. Kuo, Á. Roxin, S. Huh, Z. Zhang, R. Kandasamy, H. Merckens, J. O. Kumlin, A. Limoges, S. K. Zeisler, K.-S. Lin, F. Bénard and D. M. Perrin, Toward 18F-Labeled Theranostics: A Single Agent that Can Be Labeled with 18F, 64Cu, or 177Lu, *ChemBioChem*, 2020, **21**, 943–947.
- 28 T. Yang, L. Peng, J. Qiu, X. He, D. Zhang, R. Wu, J. Liu, X. Zhang and Z. Zha, A radiohybrid theranostics ligand labeled with fluorine-18 and lutetium-177 for fibroblast activation protein-targeted imaging and radionuclide therapy, *Eur. J. Nucl. Med. Mol. Imaging*, 2023, **50**, 2331–2341.
- 29 A. Wurzer, D. DiCarlo, A. Schmidt, R. Beck, M. Eiber, M. Schwaiger and H.-J. Wester, Radiohybrid ligands: a novel tracer concept exemplified by 18F- or 68Ga-labeled rhPSMA-inhibitors, *J. Nucl. Med.*, 2020, **61**(5), 735–742.
- 30 Á. Roxin, C. Zhang, S. Huh, M. Lepage, Z. Zhang, K.-S. Lin, F. Bénard and D. M. Perrin, A Metal-Free DOTA-Conjugated 18F-Labeled Radiotracer: [18F]DOTA-AMBF3-LLP2A for Imaging VLA-4 Over-Expression in Murine Melanoma with Improved Tumor Uptake and Greatly Enhanced Renal Clearance, *Bioconjugate Chem.*, 2019, **30**, 1210–1219.
- 31 J. Lozada, W. Xuan Lin, R. M. Cao-Shen, R. Astoria Tai and D. M. Perrin, Salt Metathesis: Tetrafluoroborate Anion Rapidly Fluoridates Organoboronic Acids to give Organotrifluoroborates, *Angew. Chem., Int. Ed.*, 2023, **62**, e202215371.
- 32 K. H. Chalmers, E. De Luca, N. H. M. Hogg, A. M. Kenwright, I. Kuprov, D. Parker, M. Botta, J. I. Wilson and A. M. Blamire, Design Principles and Theory of Paramagnetic Fluorine-Labelled Lanthanide Complexes as Probes for ¹⁹F Magnetic Resonance: A Proof-of-Concept Study, *Chem.–Eur. J.*, 2010, **16**, 134–148.
- 33 K. H. Chalmers, M. Botta and D. Parker, Strategies to enhance signal intensity with paramagnetic fluorine-labelled lanthanide complexes as probes for ¹⁹F magnetic resonance, *Dalton Trans.*, 2011, **40**, 904–913.
- 34 K. Srivastava, E. A. Weitz, K. L. Peterson, M. Marjańska and V. C. Pierre, Fe- and Ln-DOTAm-F12 Are Effective Paramagnetic Fluorine Contrast Agents for MRI in Water and Blood, *Inorg. Chem.*, 2017, **56**, 1546–1557.
- 35 J. Blahut, K. Bernásek, A. Gálisová, V. Herynek, I. Císařová, J. Kotek, J. Lang, S. Matějková and P. Hermann, Paramagnetic ¹⁹F Relaxation Enhancement in Nickel(II) Complexes of *N*-Trifluoroethyl Cyclam Derivatives and Cell Labeling for ¹⁹F MRI, *Inorg. Chem.*, 2017, **56**, 13337–13348.
- 36 R. Pujales-Paradela, T. Savić, I. Brandariz, P. Pérez-Lourido, G. Angelovski, D. Esteban and C. Platas-Iglesias, Reinforced Ni(II)-cyclam derivatives as dual 1H/19F MRI Probes, *Chem. Commun.*, 2019, **55**, 4115–4118.
- 37 M. Yu, D. Xie, K. P. Phan, J. S. Enriquez, J. J. Luci and E. L. Que, A Co(II) complex for 19F MRI-based detection of reactive oxygen species, *Chem. Commun.*, 2016, **52**, 13885–13888.
- 38 J. S. Enriquez, M. Yu, B. S. Bouley, D. Xie and E. L. Que, Copper(II) complexes for cysteine detection using 19F magnetic resonance, *Dalton Trans.*, 2018, **47**, 15024–15030.
- 39 J.-F. Morfin, R. Tripier, M. L. Baccon and H. Handel, Bismuth(III) coordination to cyclen and cyclam bearing four appended groups, *Polyhedron*, 2009, **28**, 3691–3698.
- 40 N. Bernier, J. Costa, R. Delgado, V. Félix, G. Royal and R. Tripier, trans-Methylpyridine cyclen versus cross-bridged trans-methylpyridine cyclen. Synthesis, acid-base and metal complexation studies (metal = Co²⁺, Cu²⁺, and Zn²⁺), *Dalton Trans.*, 2011, **40**, 4514–4526.
- 41 K. R. Wilson, D. J. Cannon-Smith, B. P. Burke, O. C. Birdsong, S. J. Archibald and T. J. Hubin, Synthesis and structural studies of two pyridine-armed reinforced cyclen chelators and their transition metal complexes, *Polyhedron*, 2016, **114**, 118–127.
- 42 Z. Liu, M. Pourghiasian, M. A. Radtke, J. Lau, J. Pan, G. M. Dias, D. Yapp, K.-S. Lin, F. Bénard and D. M. Perrin, An Organotrifluoroborate for Broadly Applicable One-Step 18F-Labeling, *Angew. Chem., Int. Ed.*, 2014, **53**, 11876–11880.
- 43 Deposition numbers 2314190 (for H₂Do2py2BF₃), 2314191 (for H₄Do2py2BF₃⁺), and 2314192 (for Zn(Do2py2BF₃)) contain the ESI† data for this paper.
- 44 G. Tircsó, E. Tircsóné Benyó, Z. Garda, J. Singh, R. Trokowski, E. Brücher, A. D. Sherry, É. Tóth and Z. Kovács, Comparison of the equilibrium, kinetic and water exchange properties of some metal ion-DOTA and DOTA-bis(amide) complexes, *J. Inorg. Biochem.*, 2020, **206**, 111042.
- 45 R. Ting, C. W. Harwig, J. Lo, Y. Li, M. J. Adam, T. J. Ruth and D. M. Perrin, Substituent Effects on Aryltrifluoroborate Solvolysis in Water: Implications for Suzuki–Miyaura Coupling and the Design of Stable 18F-Labeled Aryltrifluoroborates for Use in PET Imaging, *J. Org. Chem.*, 2008, **73**, 4662–4670.
- 46 Z. Liu, D. Chao, Y. Li, R. Ting, J. Oh and D. M. Perrin, From Minutes to Years: Predicting Organotrifluoroborate Solvolysis Rates, *Chem.–Eur. J.*, 2015, **21**, 3924–3928.
- 47 M. Butters, J. N. Harvey, J. Jover, A. J. J. Lennox, G. C. Lloyd-Jones and P. M. Murray, Aryl Trifluoroborates in Suzuki–Miyaura Coupling: The Roles of Endogenous Aryl Boronic Acid and Fluoride, *Angew. Chem., Int. Ed.*, 2010, **49**, 5156–5160.
- 48 D. M. Perrin, [18F]-Organotrifluoroborates as Radioprosthetic Groups for PET Imaging: From Design



- Principles to Preclinical Applications, *Acc. Chem. Res.*, 2016, **49**, 1333–1343.
- 49 Z. Liu, K.-S. Lin, F. Bénard, M. Pourghiasian, D. O. Kiesewetter, D. M. Perrin and X. Chen, One-step ^{18}F labeling of biomolecules using organotrifluoroborates, *Nat. Protoc.*, 2015, **10**, 1423–1432.
- 50 H.-T. Kuo, M. Lepage, K.-S. Lin, J. Pan, Z. Zhang, Z. Liu, A. Pryma, C. Zhang, H. Merkens, A. Roxin, D. Perrin and F. Benard, One-step ^{18}F -labeling and preclinical evaluation of prostate specific membrane antigen trifluoroborate probes for cancer imaging, *J. Nucl. Med.*, 2019, **60**, 1160–1166.
- 51 W. J. McBride, R. M. Sharkey and D. M. Goldenberg, Radiofluorination using aluminum-fluoride (Al^{18}F), *EJNMMI Res.*, 2013, **3**, 36.
- 52 R. Colotti, J. A. M. Bastiaansen, A. Wilson, U. Flögel, C. Gonzales, J. Schwitter, M. Stuber and R. B. van Heeswijk, Characterization of perfluorocarbon relaxation times and their influence on the optimization of fluorine-19 MRI at 3 Tesla, *Magn. Reson. Med.*, 2017, **77**, 2263–2271.

