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Permalink

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Journal

ACS Nano, 13(1)

ISSN

1936-0851

Authors

Jahromi, Amin Haghghat

Wang, Chao

Adams, Stephen R

et al.

Publication Date

2019-01-22

DOI

10.1021/acsnano.8b04881

Peer reviewed



Published in final edited form as:

ACS Nano. 2019 January 22; 13(1): 143–151. doi:10.1021/acsnano.8b04881.

## Fluorous-Soluble Metal Chelate for Sensitive Fluorine-19 Magnetic Resonance Imaging Nanoemulsion Probes

Amin Haghghat Jahromi<sup>†</sup>, Chao Wang<sup>†</sup>, Stephen R. Adams<sup>‡</sup>, Wenlian Zhu<sup>†</sup>, Kazim Narsinh<sup>†,§</sup>, Hongyan Xu<sup>†</sup>, Danielle L. Gray<sup>||</sup>, Roger Y. Tsien<sup>‡,⊥,#</sup>, and Eric T. Ahrens<sup>\*,†</sup>

<sup>†</sup>Department of Radiology, University of California, San Diego, La Jolla, California 92093, United States

<sup>‡</sup>Department of Pharmacology, University of California, San Diego, La Jolla, California 92093, United States

<sup>§</sup>Department of Radiology, University of Pennsylvania, Philadelphia, Pennsylvania 19104, United States

<sup>||</sup>School of Chemical Sciences, University of Illinois at Urbana–Champaign, Urbana, Illinois 61801, United States

<sup>⊥</sup>Department of Chemistry and Biochemistry, University of California, San Diego, La Jolla, California 92093, United States

<sup>#</sup>Howard Hughes Medical Institute, University of California, San Diego, La Jolla, California 92093, United States

### Abstract

Fluorine-19 MRI is an emerging cellular imaging approach, enabling lucid, quantitative “hot-spot” imaging with no background signal. The utility of <sup>19</sup>F-MRI to detect inflammation and cell therapy products *in vivo* could be expanded by improving the intrinsic sensitivity of the probe by molecular design. We describe a metal chelate based on a salicylidene-tris(aminomethyl)ethane core, with solubility in perfluorocarbon (PFC) oils, and a potent accelerator of the <sup>19</sup>F longitudinal relaxation time ( $T_1$ ). Shortening  $T_1$  can increase the <sup>19</sup>F image sensitivity per time and decrease the minimum number of detectable cells. We used the condensation between the tripodal ligand

\*Corresponding Author: (E. T. Ahrens) Phone: (858) 246-0279. eta@ucsd.edu.

#### ASSOCIATED CONTENT

##### Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: [10.1021/acsnano.8b04881](https://doi.org/10.1021/acsnano.8b04881).

Additional materials and methods associated with chemical synthesis, NMR spectra of compounds and intermediates, MRI sensitivity modeling, biological cell testing, and crystallization methods. (PDF)

X-ray crystallographic data for **5a** POP (CIF)

X-ray crystallographic data for **5a** OOO (CIF)

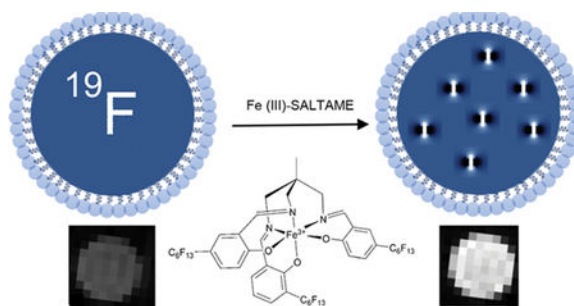
X-ray crystallographic data for **5a** PPP (CIF)

#### Notes

The authors declare the following competing financial interest(s): E.T.A. is founder, consultant, member of the advisory board, and shareholder of Celsense, Inc.

tris-1,1,1-(aminomethyl)ethane and salicylaldehyde to form the salicylidene-tris(aminomethyl)ethane chelating agent (SALTAME). We purified four isomers of SALTAME, elucidated structures using X-ray scattering and NMR, and identified a single isomer with high PFC solubility.  $Mn^{4+}$ ,  $Fe^{3+}$ ,  $Co^{3+}$ , and  $Ga^{3+}$  cations formed stable and separable chelates with SALTAME, but only  $Fe^{3+}$  yielded superior  $T_1$  shortening with modest line broadening at 3 and 9.4 T. We mixed  $Fe^{3+}$  chelate with perfluorooctyl bromide (PFOB) to formulate a stable paramagnetic nanoemulsion imaging probe and assessed its biocompatibility in macrophages *in vitro* using proliferation, cytotoxicity, and phenotypic cell assays. Signal-to-noise modeling of paramagnetic PFOB shows that sensitivity enhancement of nearly 4-fold is feasible at clinical magnetic field strengths using a  $^{19}F$  spin-density-weighted gradient-echo pulse sequence. We demonstrate the utility of this paramagnetic nanoemulsion as an *in vivo* MRI probe for detecting inflammation macrophages in mice. Overall, these paramagnetic PFC compounds represent a platform for the development of sensitive  $^{19}F$  probes.

## Graphical Abstract



## Keywords

MRI;  $^{19}F$ ; metal chelate; perfluorocarbon; nanoemulsion; macrophage; inflammation

For decades, metal chelate chemistry has been the centerpiece in efforts to formulate magnetic resonance imaging (MRI) contrast media. A compelling direction is the use of metal chelates for emerging nonproton imaging approaches such as fluorine-19 MRI.<sup>1</sup>  $^{19}F$  MRI enables hot-spot imaging with no background signal and quantification of spin-density-weighted images.<sup>2,3</sup>  $^{19}F$  MRI using perfluorocarbon (PFC) nanoemulsion (NE) probes has been used to detect cell therapy products *in vivo* (*e.g.*, stem cells and immune cells) that were labeled *ex vivo* prior to delivery to the subject; these methods have recently been translated into human patients.<sup>4</sup> In other uses, PFC probes have been used effectively for imaging leukocyte infiltrates associated with multiple inflammatory diseases.<sup>3,5</sup> In this approach, following intravenous injection, the NE droplets are taken up by monocytes and macrophages *in situ*, and these cells accumulate at sites of inflammation yielding  $^{19}F$  MRI hot-spots.

The utility of this nascent technology could be expanded by improving the sensitivity of  $^{19}F$  detection *via* molecular design.  $^{19}F$  MRI is limited by the total amount and distribution of fluorine atoms introduced into the subject's tissue, as well as the amount of PFC that can be safely internalized into cells of interest. Thus, one must improve the intrinsic sensitivity of

the PFC molecule. A key parameter for boosting sensitivity is decreasing the high  $^{19}\text{F}$  longitudinal relaxation time ( $T_1$ ) of PFC molecules. The  $T_1$  value ultimately limits the rate of  $^{19}\text{F}$  MRI data acquisition. Generally,  $^{19}\text{F}$  imaging requires summation of multiple acquisitions (*i.e.*, signal averaging) to yield a sufficient signal-to-noise ratio (SNR) to gain statistical confidence. A high  $^{19}\text{F}$   $T_1$  value generally requires a longer repetition time, thus limiting the number of acquisitions and amount of signal averaging attainable during a fixed total imaging time. Shortening  $T_1$  allows more signal averages and thus increases SNR and sensitivity and decreases the minimum number of detectable cells per voxel in the same total scan time.

The intermolecular paramagnetic relaxation enhancement (PRE) mechanism<sup>6</sup> can be used to decrease  $T_1$  by incorporating paramagnetic centers such as  $\text{Gd}^{3+}$  and  $\text{Fe}^{3+}$  into or near the fluororous phase.<sup>1,7</sup> The strength of the PRE dipole–dipole interaction is inversely proportional to the sixth power ( $1/r^6$ ) of the fluorine–metal distance. Thus, paramagnetic centers bound to the NE surface can be inefficient, due to the long distance between the relaxation agent and the bulk PFC molecules inside the NE droplet. To yield the optimal  $T_1$  and  $T_2$  (spin–spin relaxation time) with minimal metal added, the paramagnetic center should be dissolved in PFC. However, dissolving the paramagnetic center is challenging due to PFC's highly hydrophobic and lipophobic nature. Free paramagnetic cations are insoluble in PFC; thus, they must be bound to a fluororous-soluble chelating agent. Previous work has shown that metal-binding  $\beta$ -diketones conjugated to linear perfluoropolyether (PFPE) have solubility in PFCs, and optimal  $^{19}\text{F}$ -relaxation occurred when bound to  $\text{Fe}^{3+}$ , whereas  $\text{Gd}^{3+}$  yielded severe line broadening.<sup>1</sup>

In this study, we present synthesis and characterization of a fluorophilic chelating agent based on a salicylidene-tris-(aminomethyl)ethane core, referred to as SALTAME; these chelates are soluble in PFC and serve as potent PRE agents. We describe the SALTAME structure, physical properties, NMR relaxation times, and the impact of different bound metal cations, NE probe formulation, and characterization of probe-labeled macrophages *in vitro*. We demonstrate the utility of paramagnetic SALTAME-PFC NE as an *in vivo*  $^{19}\text{F}$  MRI probe for detecting inflammation-associated macrophages in mice. Overall, the creation of paramagnetic NE probes is a chemical synthesis avenue for advancing the field of  $^{19}\text{F}$  MRI.

## RESULTS AND DISCUSSION

### Molecular Design and Synthesis of the SALTAME Complex.

We describe a synthesis scheme for fluorophilic metal chelates. At the design onset, based on prior work,<sup>1</sup> we assume transition metals, and  $\text{Fe}^{3+}$  in particular, are the best  $T_1$  accelerator for PFCs, while paramagnetic lanthanides (*e.g.*,  $\text{Gd}^{3+}$ ) cause severe line broadening, essentially becoming  $^{19}\text{F}$   $T_2$  agents.<sup>1</sup> Moreover, perfluorooctyl bromide (PFOB) is the preferred  $^{19}\text{F}$ -rich MRI signal media for metal chelate dissolution due to its rapid clearance from the body and well-characterized clinical safety profile.<sup>8–10</sup>

We used (Figure 1) the condensation between tris-1,1,1-(aminomethyl)ethane (TAME) and salicylaldehyde (SAL) to form the tripodal SALTAME, for three reasons: (i) SALTAME is known to be a high-affinity hexadentate chelating agent with three N and three O atoms,

capable of binding different paramagnetic cations;<sup>11</sup> (ii) with a maximum of three negative charges, a chelated trivalent paramagnetic center (*e.g.*, Fe<sup>3+</sup>) gives an overall neutral charge that favors solubility in PFCs; (iii) the geometry of the chelate stabilizes the high-spin state of cations, such as S = 5/2 for Fe<sup>3+</sup>, which maximizes the PRE effect.<sup>11</sup> However, an unsubstituted SALTAME chelate (Figures 1 and 5 R<sub>1</sub>–R<sub>6</sub> = H) is insoluble in PFCs due to the simultaneous hydrophobic and lipophobic nature of all PFCs. As PFCs only dissolve highly fluorinated compounds, we sought to append perfluoroalkyl substituents to the SALTAME complex to increase fluorophilicity. We tried the reported photochemical ring alkylation of SA with perfluoroalkyl iodides under basic conditions,<sup>12</sup> but it gave a mixture of monoalkylated isomers **3a** and **3b** and dialkylated isomer **3c**, which were difficult to isolate (route 1). Instead, alkylation of bromosalicylaldehydes with heating under basic conditions,<sup>13</sup> followed by reductive debromination,<sup>14</sup> gave **3a** and **3b** in much higher yields (route 2). Different isomers of **5a** can be obtained by condensation of pure **3a** or **3b** or their weighed mixtures (1:2 or 2:1 w/w) with TAME and subsequent addition of ferric chloride and separation of the isomers. Although the SALTAME imines tend to dissociate, their iron complexes are very stable and can be readily isolated using chromatographic methods. Confirmation of their structures was achieved by high-resolution liquid chromatography–mass spectrometry (LC-MS) of **5a**, as well as NMR analysis of the respective SALTAME ligands (Figures S1–S3). As expected, incorporation of paramagnetic Fe<sup>3+</sup> into SALTAME greatly increases the intrinsic longitudinal and transverse relaxation rates for fluorine NMR of the **5a** isomers (Table 1) and perturbs their proton and carbon NMR spectra. Definitive structural assignment of three of the four possible **5a** isomers (**5a** POP, OOO, and PPP) is shown by X-ray crystallography (Figures 2, S4, and S5, respectively).

The four **5a** isomers were soluble in various PFC molecules, especially PFOB, and to a lesser extent (<0.5 mM) in perfluoro-15-crown-5-ether (PFCE) and perfluoropolyether (PFPE), which are other PFC compounds previously used for <sup>19</sup>F MRI applications.<sup>8</sup> Unexpectedly, the **5a** isomers have remarkably different solubility in PFOB, with **5a** PPP and **5a** POP having the highest solubility (26 and 102 mM, respectively) among all the isomers (**5a** OOP and **5a** OOO solubilities are 9.4 and 2.0 mM, respectively). Initial studies indicated that more stable PFOB nanoemulsions with **5a** POP were formed with concentrations up to 30 mM of the SALTAME incorporated; thus we explored the properties and applications of this isomer. We also synthesized four additional SALTAME complexes **5c–f** to explore the inclusion of other fluororous substituents, but all showed inferior solubility in PFOB.

### Cation Selection.

We screened the impact of various cations bound to SALTAME **5a** and dissolved in PFOB *via* NMR relaxometry and susceptibility shift measurements (Table 2). Of the period-4 cations tested (V<sup>3+</sup>, Cr<sup>3+</sup>, Mn<sup>2+</sup>, Fe<sup>3+</sup>, Co<sup>2+</sup>, Cu<sup>2+</sup>, Ni<sup>2+</sup>, Zn<sup>2+</sup>, Ga<sup>3+</sup>), only Mn<sup>4+</sup>, Fe<sup>3+</sup>, Co<sup>3+</sup>, and Ga<sup>3+</sup> formed stable and separable (by column chromatography) chelates with SALTAME. These chelates readily dissolve in PFOB, and their intermolecular <sup>19</sup>F NMR relaxivities and susceptibility shift properties at 9.4 T are displayed in Table 2. The diamagnetic Ga<sup>3+</sup> chelate had little relaxation effects or shift for PFOB. Co<sup>3+</sup> has a significant effect on the transverse relaxivity (*r*<sub>2</sub>) but not on the longitudinal relaxivity (*r*<sub>1</sub>),

and the shift was not impacted.  $\text{Mn}^{4+}$  changes  $r_1$  and  $r_2$  and the susceptibility shift, but to a much lesser extent than  $\text{Fe}^{3+}$ .  $\text{Gd}^{3+}$  did not form a stable chelate with SALTAME, presumably because it requires eight to nine coordination sites, whereas SALTAME has a coordination of six.

The longitudinal and transverse  $^{19}\text{F}$  relaxation rates of the  $\text{Fe}^{3+}$  chelate, **5a** POP, was further evaluated as a function of iron concentration at the clinically relevant field strength of 3 T, yielding relaxivities  $r_1$  and  $r_2$  of 0.56 and  $1.67 \text{ s}^{-1} \text{ mM}^{-1}$ , respectively, compared to 0.50 and  $1.07 \text{ s}^{-1} \text{ mM}^{-1}$  values of  $r_1$  and  $r_2$  at 9.4 T, respectively. For neat PFOB,  $R_1/R_2$  values are  $0.79 \text{ s}^{-1}/3.5 \text{ s}^{-1}$  and  $1.4 \text{ s}^{-1}/2.2 \text{ s}^{-1}$  at 3 and 9.4 T, respectively.

Overall, the **5a** POP  $\text{Fe}^{3+}$  chelate served as a representative additive and is used to formulate paramagnetic PFOB (P-PFOB) NE, which was used for further studies.

### Structure of $\text{Fe}^{3+}$ SALTAME.

To characterize the structure and  $\text{Fe}^{3+}$  coordination in chelates, data for **5a** POP, PPP, and OOO were obtained by X-ray crystallography (Figures 2, S4, and S5) and NMR methods. Comparison of these structures with unsubstituted  $\text{Fe}^{3+}$ -SALTAME (**5**,  $\text{R}_1\text{-R}_6 = \text{H}$ ; Figure S6) confirms that the hexadentate coordination of  $\text{Fe}^{3+}$  is not affected by appending fluorine substituents to SALTAME with minimal differences in Fe–O or Fe–N bond lengths and angles compared to its nonfluorinated parent.<sup>11</sup> From these structures, we speculate that the low solubility of **5a** OOO in PFOB (Table S1) may be explained by the observation that all three fluorine substituents are located on one side of **5a** OOO, yielding less favorable van der Waals interactions between substituents and PFOB. Fluorine substituents are more evenly distributed around the chelate in **5a** PPP and POP, yielding higher affinity to PFOB.

Intramolecular  $^{19}\text{F}$  NMR relaxometry measurements of **5a** isomers further confirm the structural analyses. We observe that among  $\text{Fe}^{3+}$  chelates, the **5a** OOO isomer has the highest and the **5a** PPP isomer has the lowest  $R_1$  and  $R_2$  values (Table 1). Ortho-substitution causes a more drastic intramolecular PRE, by  $\sim 3$  orders of magnitude, compared to para-substitution due to the closer proximity of the  $\text{Fe}^{3+}$  paramagnetic center to the ortho position (6.91 nm) compared to the para position (9.53 nm).

### Sensitivity Enhancement of P-PFOB.

We modeled the potential MRI sensitivity enhancement of P-PFOB compared to undoped PFOB based on measured  $^{19}\text{F}$  NMR relaxivities in these materials. Modeling results (Figure 3a) show multifold sensitivity improvement is possible using a gradient-echo (GRE)-based MRI pulse sequence with echo time (TE)  $< 0.8$  ms. Model details are given in the Supporting Information (Section 4).

To support these findings, experimental data were acquired in an MRI phantom (Figure 3b) consisting of two NMR tubes containing PFOB and P-PFOB nanoemulsions. The phantoms were imaged twice at 9.4 T using a GRE chemical shift imaging (CSI) pulse sequence with two different repetition time (TR) parameters set according to  $\text{TR} = 0.5 T_1$  for PFOB or P-PFOB, respectively, with the optimal Ernst angle condition set for each TR value; both images were acquired using the same total imaging time. (See Supporting Information for

additional details.) Image results (Figure 3b) show the P-PFOB tube has a 2.5 times higher SNR than the PFOB tube when comparing the optimal SNR acquisition for each tube, consistent with the model prediction. This phantom imaging example does not represent the absolute value or upper limit to the possible SNR enhancement achievable with these materials; varying  $[\text{Fe}^{3+}]$ , magnetic field strength, pulse sequences, and acquisition parameters may yield different values of SNR enhancement.

### Imaging Probe Formulation.

To create NE-based imaging probes, we formulated colloidal suspensions of P-PFOB. High-shear homogenization of components was used to form NE.<sup>15</sup> To stabilize the NE, egg yolk phospholipids (EYP) was used as a surfactant; P-PFOB has a slight lipophilic character due to PFOB's single bromine, yielding a cohesive tendency between PFOB and the fatty acid chains of EYP.<sup>16,17</sup> Minor surfactant components were also added including  $\text{CH}_3-(\text{CH}_2)_5-(\text{CF}_2)_5-\text{CF}_3$ , which acts as molecular dowel, improving the emulsion stability,<sup>18</sup> Cremophor EL (Polyoxyl 35 hydrogenated castor oil), to decrease NE droplet size and increase shelf life and circulation time,<sup>19</sup> and mannitol as an isotonicizer. After homogenization of P-PFOB ( $[\mathbf{5a} \text{ POP}] = 20 \text{ mM}$ ) with surfactants, the average NE particle size was  $\sim 162 \text{ nm}$  with a polydispersity index (PDI) of 0.27, as measured by dynamic light scattering (DLS). Using similar methods, this particle size is comparable to nanoemulsions made without a chelate.<sup>15</sup> The final P-PFOB NE had  $[\text{F}] = 14.26 \text{ M}$  with a PFOB: $\mathbf{5a}$  POP molar ratio of 192:1. The  $R_1$  and  $R_2$  parameters were  $9.01 \pm 0.03 \text{ s}^{-1}$  and  $16.66 \pm 0.10 \text{ s}^{-1}$  at 9.4 T, respectively. The NE stability was confirmed by longitudinally monitoring the particle size using DLS measurements for an emulsion stored at 4 °C over 60 days (Figure S7). Also, the NE complex is stable for at least 3 days at 37 °C in proteinaceous saline media used to mimic the blood circulation environment (Figure S8).

We also evaluated the iron-binding stability of P-PFOB NE ( $[\text{Fe}^{3+}\text{-SALTAME } \mathbf{5a} \text{ POP}] = 30 \text{ mM}$  in PFOB) following addition of 50 mM ethylenediaminetetraacetate (EDTA), a strong competing  $\text{Fe}^{3+}$  chelating agent, to the aqueous phase; P-PFOB NE shows stable relaxometry parameters over 20 days, indicating that iron is tightly bound in the fluorine phase (Figure S9).

### P-PFOB Labeling Macrophages *in Vitro*.

To test biocompatibility of formulated P-PFOB NE as a cell labeling agent, we performed *in vitro* assays on a labeled murine macrophage cell line derived from blood (RAW 264.7). This cell is a representative phagocyte that would take up NE *in vivo* after intravenous delivery. Following a 16 h incubation of RAW cells with various NE concentrations, the uptake level approached  $\sim 10^{12}$  fluorine atoms per cell, measured by  $^{19}\text{F}$  NMR (Figure S10).

We investigated the potential effect of P-PFOB labeling on RAW cell viability and phenotype (Figure 4). To study cytotoxicity, we used a flow cytometry assay for apoptosis (Figure 4a–c) using 10-*N*-nonyl acridine orange (NAO).<sup>20</sup> As a positive control to induce oxidative stress, 10% ethanol was added to the cell media. No evidence of cytotoxicity is observed in NE-labeled compared to unlabeled cells (Figure 4a,b), whereas ethanol-treated cells display cytotoxicity (Figure 4c) and a characteristic bimodal appearance described



elsewhere.<sup>20</sup> Figure 4b also shows that the labeled cells have increased side-scatter signal due to light scattering from PFC-loaded intracellular vesicles, a common observation. Side-scatter is routinely used in flow cytometry to distinguish cell types by granularity. Labeled cells often display increased granularity with internalization of PFC droplets. To assay cell phenotype, we used flow cytometry to monitor potential changes in cell surface CD86 expression as a marker for macrophage activation, where increased levels indicate a pro-inflammatory response to label. Lipopolysaccharide (LPS) treatment was used as a positive control to induce a pro-inflammatory phenotype. The CD86 expression level in NE-labeled macrophages is comparable to unlabeled cells (Figure 4d), whereas in LPS-treated cells the level is visibly higher, suggesting that P-PFOB NE does not stimulate a pro-inflammatory phenotype in macrophages.

### Visualizing Inflammation *in Vivo* Using P-PFOB NE.

Proof-of-concept *in vivo* imaging of inflammation was performed using P-PFOB NE in a localized inflammation mouse model. Following established protocols,<sup>5,8,21</sup> localized inflammation was induced using an LPS-doped Matrigel solution injected subcutaneously into the posterior neck of a C57BL/6 mouse ( $N = 3$ ). After 2 h, a single bolus of P-PFOB NE (200  $\mu\text{L}$ ,  $[\text{Fe}^{3+}\text{-SALTAME } \mathbf{5a} \text{ POP}] = 20\text{mM}$  in PFOB oil phase), equivalent to 143 mmol F/kg body weight and 0.04 mmol chelated Fe/kg body weight, was injected intravenously. No adverse reactions were observed. Mice were imaged 24 h after injection to allow for nanoemulsion uptake by monocytes/macrophages *in situ*. Scans were performed at 11.7 T. The  $^{19}\text{F}$  images were acquired using a two-dimensional CSI pulse sequence along with conventional anatomical  $^1\text{H}$  images. A representative coronal image in the neck region is displayed in Figure 5, where the Matrigel plug appears as a hyperintense, subcutaneous structure in the dorsal region of the  $^1\text{H}$  image (Figure 5). A robust  $^{19}\text{F}$  signal signifying macrophage uptake is seen co-registered with the Matrigel plug.  $^{19}\text{F}$  is also found in an area in the anterior neck, presumably at a lymph node (asterisk, Figure 5). Overall, these preliminary *in vivo* data show the feasibility of using P-PFOB NE as an inflammation imaging agent.

### Susceptibility Shifts by $\text{Fe}^{3+}$ SALTAME.

In unemulsified P-PFOB oil we observe significant  $^{19}\text{F}$  shifts in all PFOB peaks with the addition of  $\mathbf{5a}$  POP  $\text{Fe}^{3+}$  chelate (Figure 6a). This shift is linear with concentration of  $\mathbf{5a}$  POP in PFOB, with a slope of  $-0.060$  ppm/mM at 9.4 T. Generally, a  $^{19}\text{F}$  shift reagent can be used for multispectral (color-coded) MRI. To demonstrate multispectral MRI with P-PFOB oil, a phantom was prepared using varying concentrations of  $\mathbf{5a}$  POP in three NMR tubes containing PFOB dissolved with  $[\mathbf{5a} \text{ POP}] = 0, 14.5, \text{ and } 25$  mM. It was imaged using a CSI pulse sequence. The red–green–blue overlays (Figure 6a, right) represent the shift images generated at the characteristic  $\text{CF}_3$  resonance frequencies of each  $[\mathbf{5a} \text{ POP}]$  concentration. The linear relationship between shift change and  $[\mathbf{5a} \text{ POP}]$  provides the potential for multiplexed imaging using different amounts of additive  $\mathbf{5a}$  POP to achieve  $^{19}\text{F}$  multichromicity.

However, we attribute the observed field shift to bulk magnetic susceptibility (BMS) effects,<sup>22</sup> where shift is dependent on sample geometry (*i.e.*, cylindrical NMR tube). Upon



formulating the P-PFOB oil into a colloidal suspension of spherical NE droplets (described above), the P-PFOB shift compared to PFOB is strongly attenuated (Figure 6b,c).

Overall, we describe the design and characterization of SALTAME, a stable hexadentate chelating agent for iron(III). We used this moiety to formulate a nanoemulsion MRI probe that can be used for “hot-spot” detection *in vivo*. Incorporation of iron-bound SALTAME into the fluorous phase of NE causes a profound reduction of the  $^{19}\text{F}$   $T_1$  value and only mild line broadening, thus offering improved sensitivity of  $^{19}\text{F}$  MRI due to increased signal averaging and/or reduced MRI scan time.

One of the primary challenges in the emerging field of  $^{19}\text{F}$  MRI is sensitivity, and improvement of the intrinsic MRI sensitivity of the PFC molecule employed could lower the barriers for wider use in biomedical applications. Generally, PFCs are often employed due to their high F-density and safety profile. PFOB has been shown to be a promising molecular candidate for the translation of  $^{19}\text{F}$  MRI because of its low toxicity and rapid clearance from the body due to its modest lipophilicity.<sup>8,23</sup> In fact, PFOB has already been used in patients as an oxygen transporting media.<sup>9,16</sup> Importantly, it has a short half-life (3–8 days), *in vivo*. By comparison, PFCE, another PFC molecule commonly used for  $^{19}\text{F}$  MRI, has a body half-life of >100 days.<sup>16</sup> Generally, clearance of PFC NE agents from the body occurs *via* uptake into cells and organs of the reticuloendothelial system, followed by lung exhalation.<sup>24</sup> However, PFOB has some undesirable properties as a  $^{19}\text{F}$  probe; it has a complex multipeak  $^{19}\text{F}$  spectrum and intrinsically high  $T_1$  values common for many PFC molecules.<sup>25</sup> In fact, among the most common PFCs, PFOB has the highest  $T_1$  value of ~1.2 s, compared to ~0.9 ms for PFCE, for example, at 3 T.<sup>26</sup> Generally,  $^{19}\text{F}$  MRI requires signal averaging, and the high  $T_1$  value of PFOB limits the rate of  $^{19}\text{F}$  MRI data acquisition within a constrained imaging time. Shortening  $T_1$  can increase the SNR per time and decrease the minimum number of detectable cells per voxel.

$T_1$  can be profoundly altered by paramagnetic cations due to PRE. In prior studies, PRE has been used for  $^{19}\text{F}$  probes using lanthanide macrocycles with fluorinated sites.<sup>27–30</sup> These chelates<sup>29</sup> yield a reduction of ~2–3 orders of magnitude in intramolecular  $T_1$ , similar to  $\text{Fe}^{3+}$ -SALTAME **5a** isomers (Table 1). However, the relatively low  $^{19}\text{F}$  content of macrocyclic chelates makes it challenging to achieve detectable intracellular labeling levels, compared to fluorine-dense PFC oils. Gd-macrocyclic chelates tethered to an NE surface provides mild  $^{19}\text{F}$   $T_1$  enhancement, but are unstable inside cells.<sup>31</sup>

$\text{Fe}^{3+}$ -SALTAME retains the distorted octahedral geometry and, presumably, the high-spin state of the previously reported  $\text{Fe}^{3+}$  chelate without fluororous substituents.<sup>11</sup> Its high stability and affinity for  $\text{Fe}^{3+}$  permit easy isolation of the complex and formulation into PFOB NE without the requirement of the subsequent iron loading step necessary for prior compounds.<sup>1</sup> We observed that  $\text{Fe}^{3+}$  is the most effective cation for intermolecular  $^{19}\text{F}$   $T_1$  acceleration in PFOB, consistent with our prior work.<sup>1</sup>  $\text{Gd}^{3+}$  and  $\text{Fe}^{3+}$  ions form the basis of  $T_1$ - and  $T_2$ - based  $^1\text{H}$  contrast agents, respectively, but for  $^{19}\text{F}$  MRI, the roles these ions play are reversed.<sup>1</sup>

The *in vivo* inflammation mouse model results (Figure 5) are preliminary data showing *in vivo* compatibility and MRI detectability of P-PFOB NE. The method of using intravenously delivered  $^{19}\text{F}$  MRI for “*in situ*” labeling of macrophages to elucidate inflammation hot-spots is well established and has been used preclinically in a wide range of disease models.<sup>3,23,32</sup> Alternative PFC NE formulations facilitate *ex vivo* cell uptake in culture,<sup>33</sup> for example during the preparation of stem cell or immunotherapeutic cytotherapies, and following transfer to the patient, these cells are detected *in vivo* using  $^{19}\text{F}$  MRI.<sup>4</sup> The fluorine inside the cells yields positive-signal hot-spot images that can be quantified to measure apparent cell numbers at sites of accumulation, thereby enabling “*in vivo* cytometry”.<sup>34</sup> The sensitivity limits of detection are on the order of  $10^4$ – $10^5$  cells/voxel.<sup>3</sup>

The relaxation rates of P-PFOB are predictably tuned by varying Fe concentration. Tests were performed using a range of concentrations less than the maximum because these are the most useful for MRI experiments using the conventional (*i.e.*, spin echo or gradient echo) pulse sequences most used in MRI practice (see SI Section 4). However, advanced MRI pulse sequences, such as zero time to echo (ZTE) and ultrashort time to echo (UTE), which are used by early adoptors in MRI practice,<sup>29</sup> can predictably reap the benefits of the maximum iron concentration when  $T_1$  and  $T_2$  are very short; a detailed comparison of the performance of P-PFOB with different pulse sequences is beyond the scope of this article.

The total amount of Fe delivered *via* the P-PFOB injection is miniscule compared to innate Fe levels in mice. The total dose of SALTAME-bound Fe in the inoculant is approximately  $50\ \mu\text{g}$  of chelated Fe. The amount of Fe contained in mouse blood tissue, which represents roughly 2/3 of the total iron stores in the body, is approximately 0.75 mg of Fe, 15-fold higher. Thus, Fe metabolism of the organism should not be impacted by addition of P-PFOB, which contains small amounts of saturated, SALTAME-bound Fe.

The cell toxicity data presented (Figure 4) is preliminary, but shows no overt toxicity induced by intracellular labeling; the flow-based NAO assay is very sensitive, and both positive and negative controls were included in the experimental design. Generally, the cell safety of PFC cell labeling has been demonstrated exhaustively in numerous studies and cell types,<sup>35–38</sup> including primary immune cells for a human clinical trial<sup>4</sup> using a panel of *in vitro* cell assays of phenotype and function, including NAO.

We observe significant linear  $^{19}\text{F}$  NMR field shifts in all PFOB peaks with the addition of a SALTAME  $\text{Fe}^{3+}$  chelate (Figure 6a) in the unemulsified oil. SALTAME as a  $^{19}\text{F}$  shift reagent could be used for multicolor MRI tags composed of neat P-PFOB oils, for example, to image catheter tips<sup>39</sup> and the gastrointestinal track<sup>38</sup> or for external  $^{19}\text{F}$  fiduciary capsules.

## CONCLUSIONS

Here, we describe the design and characterization of SALTAME, a stable hexadentate chelating agent for iron(III). We used this moiety to formulate a nanoemulsion MRI probe that can be used for “hot-spot” detection *in vivo*. Incorporation of iron-bound SALTAME into the fluorous phase of NE causes a profound reduction of the  $^{19}\text{F}$   $T_1$  value and only mild

line broadening, thus offering improved sensitivity of  $^{19}\text{F}$  MRI due to increased signal averaging and/or reduced MRI scan time.

Overall, the use of fluorinated molecules is emerging as an option for cellular imaging probe design. This probe has the potential to enable longitudinal, noninvasive quantification of inflammation and therapeutic cell delivery and aid in the monitoring of therapeutic test articles.

## METHODS

### Materials.

Chemicals were purchased from Sigma-Aldrich (St. Louis, MO, USA) or other reputable suppliers, and organic solvents (liquid chromatography-grade) were purchased from Fisher Scientific (Pittsburgh, PA, USA) and used as received. Anhydrous dimethylformamide (DMF) and acetonitrile were stored over activated 4 Å molecular sieves. Fluorous solvents were purchased from Perseptive Biosystems (Framingham, MA, USA). Precoated silica gel plates (60 F-254, Merck Millipore, Billerica, MA, USA) were used for thin-layer chromatography (TLC), and silica gel 60 (230–400 mesh) was used for column chromatography.

### Analytic Methods.

All reactions were carried out under  $\text{N}_2$  unless otherwise noted. Reactions were monitored by TLC and LC-MS using an Agilent (Santa Clara, CA, USA) 1100 HPLC with MSD Trap XLT using a Phenomenex Luna C18(2) 100 Å, 5  $\mu\text{m}$ , 4.6 mm  $\times$  250 column, MeCN/ $\text{H}_2\text{O}$  linear gradients with constant 0.05% v/v  $\text{CF}_3\text{CO}_2\text{H}$  additive, 1 mL/min flow, and ESI positive or negative ion mode. Reaction products were purified by flash chromatography on silica gel eluted with ethyl acetate and hexane. High-resolution mass spectrometry was performed by the Molecular Mass Spectrometry Facility at the University of California San Diego. Ultraviolet–visible (UV–vis) absorption spectra were recorded on a Shimadzu (Kyoto, Japan) spectrophotometer. Solubility of  $\text{Fe}^{3+}$ -SALTAME compounds in perfluorocarbons was determined by absorbance at 450 nm following dilution in ethyl acetate using experimentally determined extinction coefficients of 700  $\text{M}^{-1} \text{cm}^{-1}$  for the 5a isomers.

### NMR Measurements.

NMR spectra and relaxation rates were recorded on a Bruker Ascend 400 MHz (9.4 T) spectrometer (Billerica, MA, USA). Additional relaxation rate measurements were performed at 3 T using a GE750 (General Electric, Milwaukee, WI, USA) clinical MRI system equipped with a custom dual-channel  $^{19}\text{F}/^1\text{H}$  72 mm diameter volume coil (RAPID Biomedical, Rimpf, Germany). The  $^{19}\text{F}$  NMR spectra were referenced to an internal standard, sodium trifluoroacetate (NaTFA, 0.1 wt %,  $-76.00$  ppm, Sigma-Aldrich, T6508), which was placed in a separate sealed capillary tube within the NMR tube. Relaxation measurements were performed using a standard inversion recovery pulse sequence and a Carr–Purcell–Meiboom–Gill sequence with echo-time values in 12 linear increments. The  $T_1$  and  $T_2$  values were obtained by nonlinear fitting using MNova 6.0.2 software (Mestrelab,

Escondido, CA, USA). Fit errors were less than 5% of  $T_1$  and  $T_2$  values. For quantitative peak measurements, spectra were acquired using calibrated  $90^\circ$  pulses, 32 000 complex points, spectral window 0 to  $-200$  ppm, relaxation delay 2.5 s, and 128 averages. Spectra were processed in MNova with manual phase correction and Whittaker smoother baseline correction (filter = 1 ppm). Relaxation parameters were measured for six middle  $CF_2$  peaks of PFOB, spanning  $\sim 10$  ppm, which account for 12 out of 17 F atoms in the molecule. To assay nanoemulsion uptake ( $^{19}F$ /cell) using NMR, an aliquot of  $1 \times 10^6$  cells was pelleted and resuspended in 0.1 mL of lysis solution (0.5% Triton X, 100mM NaCl, 20mM Tris, 1 mM EDTA), and reference compound NaTFA (0.745 mg/mL)/ $D_2O$  was added to a 5 mm NMR tube. The integrated ratio of PFOB fluorine peaks to NaTFA singlet fluorine peak at  $-76.00$  ppm determined the amount of  $^{19}F$  in the cell pellet, and the  $^{19}F$ /cell was calculated.

### Phantom MRI.

A phantom consisted of two 5 mm NMR tubes containing PFOB and P-PFOB ([5a POP] = 20 mM in PFOB oil) nanoemulsions side-by-side. Images were acquired using a 9.4 T Bruker Avance III HD Nanobay spectrometer equipped for microimaging with a double-tuned 10mm  $^1H/^{19}F$  coil and ParaVision 6 software. Two sets of  $^{19}F$  images were acquired using a two-dimensional CSI method to yield optimal SNR for each tube. Imaging parameters were TE = 0.8 ms, matrix size  $32 \times 32$ , field of view  $9.5 \times 9.5 \times 3$  mm<sup>3</sup>, and acquisition time 49 min for each. To optimize SNR for each tube, the TR was set to 103 ms with 28 averages, and then TR = 721 ms and four averages, with the optimal Ernst angle condition set for each TR value.

### In Vivo MRI.

Live mouse images were acquired using an 11.7 T horizontal-bore Bruker BioSpec MRI system equipped with a double-tuned  $^1H/^{19}F$  mouse volume coil and ParaVision 6 software. Mice were anesthetized using 1.5% isoflurane in  $O_2$  and maintained at 37 °C during acquisitions. The  $^{19}F$  images were acquired using a two-dimensional CSI method with TR/TE = 13.3/0.53 ms, matrix size  $32 \times 32$ , field of view  $20 \times 20 \times 6$  mm<sup>3</sup>, 134 averages, and acquisition time  $\sim 30$  min. For anatomical data,  $^1H$  spin-echo images were acquired with TR/TE = 550/14 ms, matrix size  $128 \times 96$ , field of view  $20 \times 20 \times 1$  mm<sup>3</sup>, and acquisition time  $\sim 3$  min. CSI visualization was performed using the CSI Visualization Tool inside the ParaVision software. The calculation and display of the phantom was performed by selecting all PFOB resonance peaks from  $-60$  to  $-130$  ppm. Notably, examination of the CSI spectral data showed no peaks at  $-82.8$  and  $-89.9$  ppm, corresponding to isoflurane anesthesia.

### Shift MRI.

Images to demonstrate POP  $Fe^{3+}$  chelate dissolved in PFOB oil induces bulk magnetic susceptibility shifts were acquired using the above 11.7 T MRI system. A CSI pulse sequence was used with TR/TE = 26.4/0.53 ms, matrix size  $64 \times 64$ , FOV  $30 \times 30 \times 3$  mm<sup>3</sup>, 1024 complex points, spectrum width 40.8 kHz, six averages, and acquisition time 10.8 min. CSI visualization was performed using the CSI Visualization Tool in ParaVision software. The calculation and display of the P-PFOB map was performed by selecting the  $CF_3$  resonance peak only.

The crystallographic data have been deposited at the Cambridge Crystallographic Data Centre (CCDC) as CCDC 1572939 (**5a** POP at 100 K), CCDC 1572845 (**5a** OOOat 100 K), and CCDC 1572938 (**5a** PPP at 100 K) and can be obtained free of charge from the CCDC via [www.ccdc.cam.ac.uk/getstructures](http://www.ccdc.cam.ac.uk/getstructures).

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## ACKNOWLEDGMENTS

Funding for E.T.A. was provided by National Institutes of Health (NIH) grants R01-EB017271, R01-EB024015, and R01 CA139579 and the California Institute for Regenerative Medicine LA1-C12-06919. A.H.J. is funded by a Cook Medical Cesare Gianturco/RSNA Research Resident grant and NIH T32 4T32EB005970.

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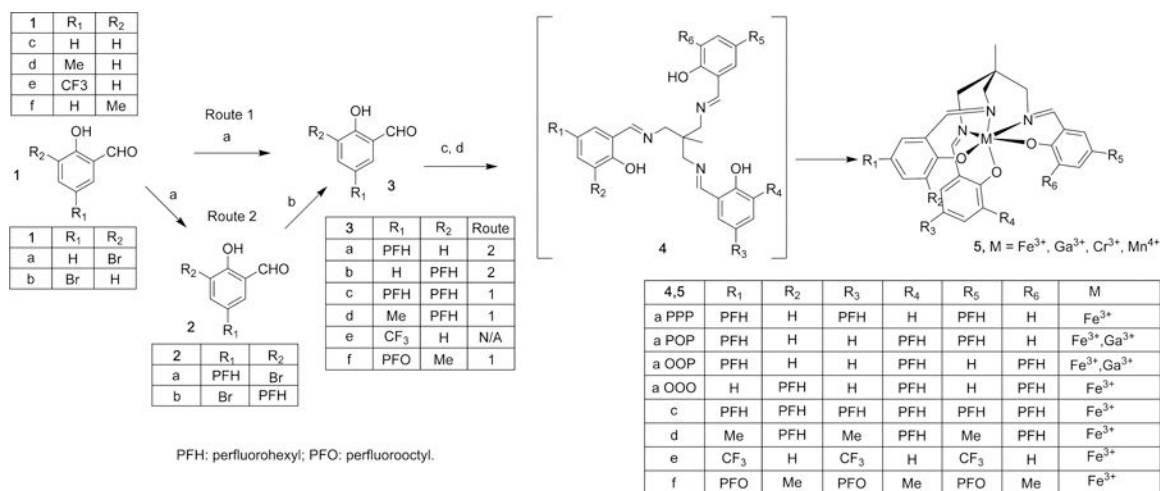
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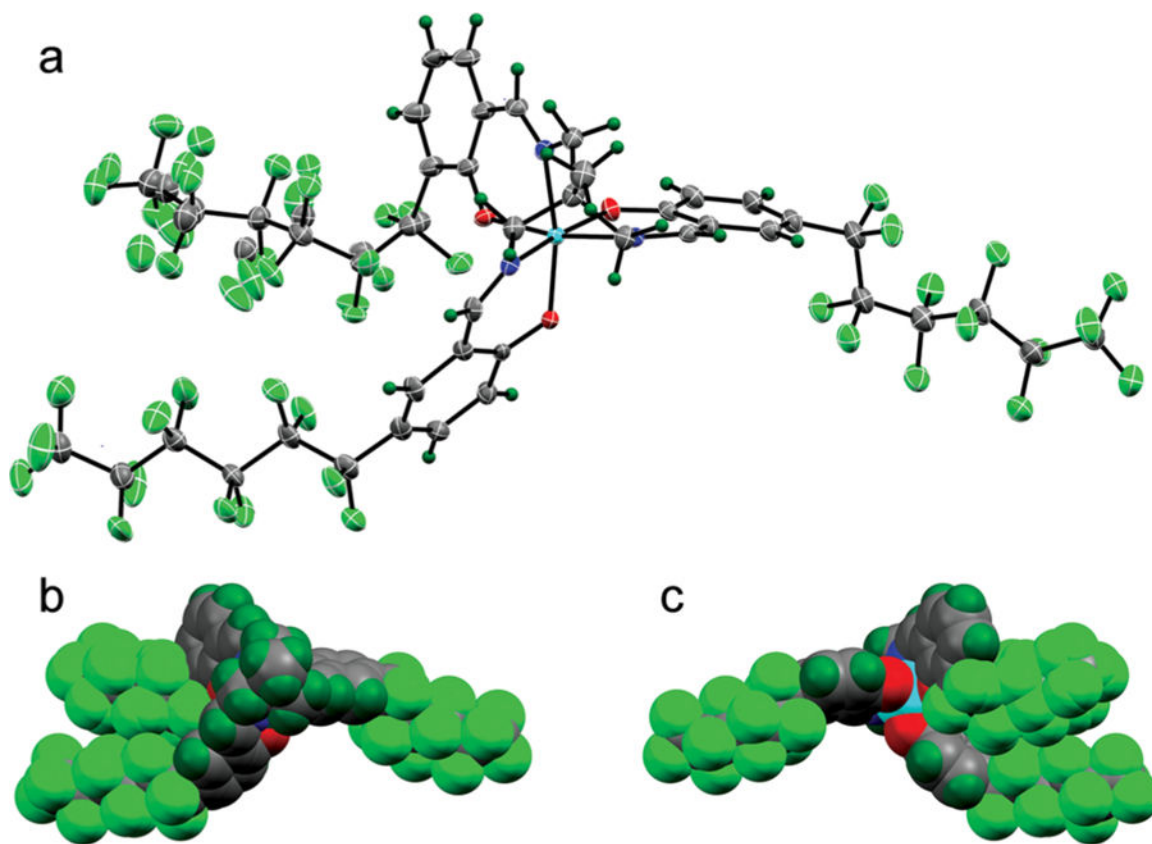


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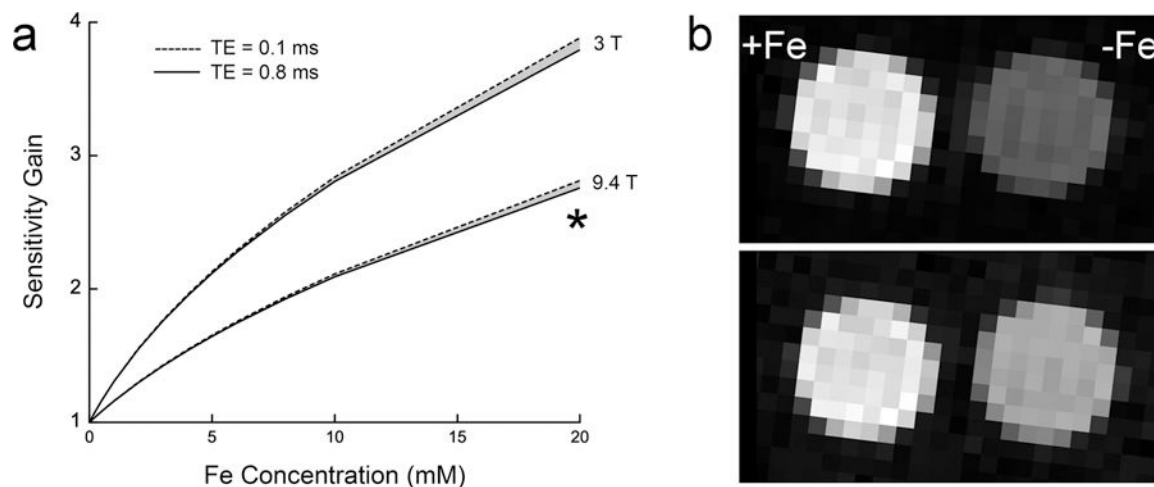


**Figure 1.**

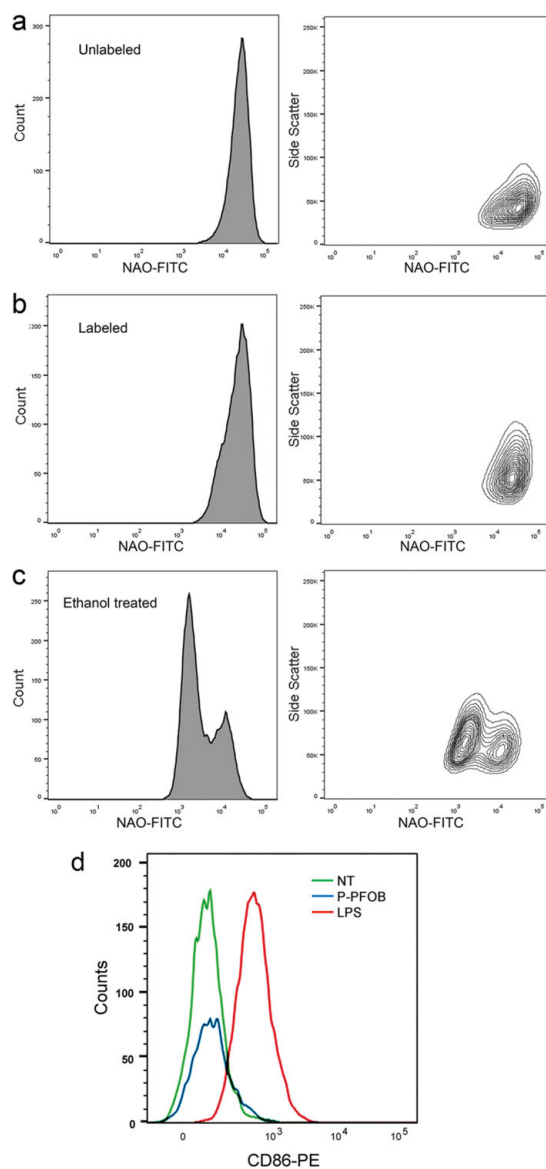
Synthetic scheme for chelating agents. Fe<sup>3+</sup>-SALTAME chelate 5a POP is used to form P-PFOB. Reagents: (a) perfluoroalkyl iodide, CsCO<sub>3</sub>, DMF, 100 °C; (b) 1 atm of H<sub>2</sub>, Pd-C, NaOAc, MeOH; (c) TAME, NEt<sub>3</sub>, EtOH, 80 °C; (d) FeCl<sub>3</sub>, NaOAc, EtOAc-EtOH.



**Figure 2.** Crystal structure of 5a POP using X-ray crystallography. (a) Structure confirms the hexadentate coordination of the chelated  $\text{Fe}^{3+}$ ; (b) space-filling rendering of the same view and (c) after a 180 deg rotation to reveal the solvent-exposed face of the bound ferric ion. Iron is colored cyan for clarity.

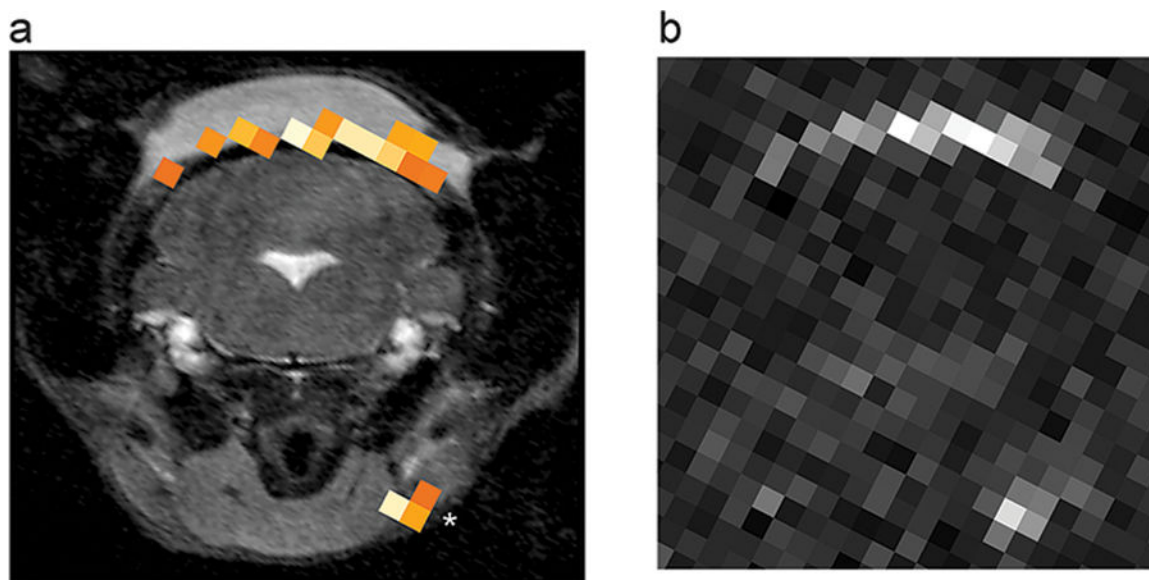


**Figure 3.** P-PFOB  $^{19}\text{F}$  MRI signal intensity enhancement. Panel (a) shows simulated sensitivity gain of P-PFOB *versus* PFOB at 3 and 9.4 T as a function of nanoemulsion-bound iron. Sensitivity gain is defined as  $\text{SNR}(\text{P-PFOB})/\text{SNR}(\text{PFOB})$  for constant time imaging. The model uses the empirically measured  $r_1$  and  $r_2$  relaxivities. The shaded regions represent a 0.1–0.8 ms TE parameter range. See Supporting Information Section 4 for model details. (b) Phantom  $^{19}\text{F}$  images (9.4 T) for P-PFOB (+Fe) and PFOB (–Fe). The top and bottom panels show the tubes optimized separately for P-PFOB ( $[\text{5a POP}] = 20 \text{ mM}$  in oil) and PFOB SNR, respectively. From these images P-PFOB displays a 2.5-fold increase in SNR over PFOB when both are imaged optimally, and the asterisk in panel (a) shows approximate experimental agreement ( $\sim 7.5\%$  error) with the model. CSI parameters for (b) were 28 averages,  $\text{TR} = 103 \text{ ms}$ , and  $\text{TE} = 0.8 \text{ ms}$  for the top panel P-PFOB and four averages,  $\text{TR} = 721 \text{ ms}$ , and  $\text{TE} = 0.8 \text{ ms}$  for the bottom panel PFOB.



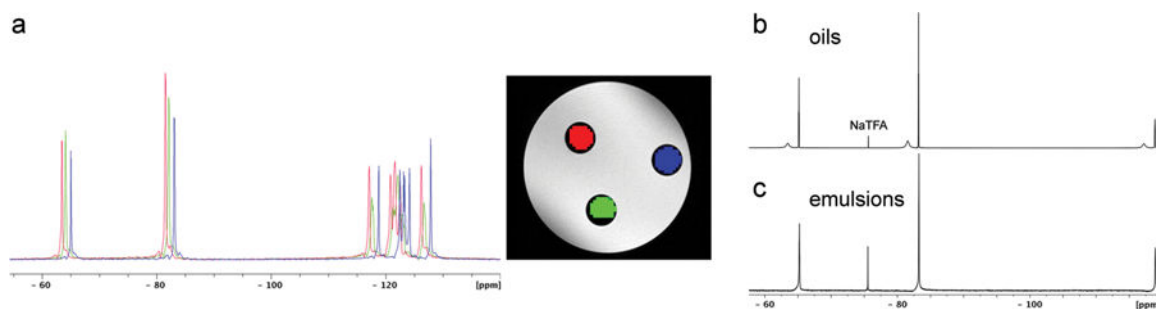
**Figure 4.**

*In vitro* cytotoxicity and cell phenotype in P-PFOB-labeled RAW cells. Cells were incubated with P-PFOB NE ( $[\text{Fe}^{3+}\text{-SALTAME 5a POP}] = 20 \text{ mM}$  in PFOB) in culture for 16 h. The NAO flow cytometry assay was used to measure cytotoxicity (apoptosis). (a) Unlabeled (negative) control cells, (b) labeled cells, and (c) 10% ethanol treated cells as positive control. (d) Flow cytometry results for CD86 expression in P-PFOB-labeled RAW cells. LPS-treated cells serve as a positive control.



**Figure 5.**

*In vivo*  $^1\text{H}/^{19}\text{F}$  MRI using P-PFOB NE to visualize inflammation in mice. An LPS-doped Matrigel plug implanted in the subcutaneous tissue of the neck was used to induce localized inflammation. One day after intravenous injection of P-PFOB NE ( $[\text{Fe}^{3+}\text{-SALTAME 5a POP}] = 20\text{mM}$  in PFOB oil phase, yielding 4.4 mM in inoculant after P-PFOB NE formulation in water), coronal  $^1\text{H}/^{19}\text{F}$  MRI data of the plug region were acquired. (a) Grayscale  $^1\text{H}$  showing the hindbrain, cervical spine, and isointense Matrigel in the subcutaneous tissue of the neck. The  $^{19}\text{F}$  signal, rendered in pseudocolor, is present in the periphery of the Matrigel plug and nearby lymph node (asterisk), consistent with the presence of inflammatory infiltrates. (b) Raw, unthresholded  $^{19}\text{F}$  image. The peak voxel  $^{19}\text{F}$  SNR at the Matrigel boundary and lymph node is 17.6 and 16.4, respectively. For  $^{19}\text{F}$ , CSI imaging parameters were as follows: 134 averages, TR = 13.3 ms, TE = 0.53 ms, and matrix size is  $32 \times 32$ . For  $T_2^*$ -weighted  $^1\text{H}$ , TR/TE = 550/14 ms and matrix size is  $128 \times 96$ . For additional details see the Supporting Information.



**Figure 6.**

Bulk magnetic susceptibility (BMS) shifts for P-PFOB. (a, left)  $^{19}\text{F}$  CSI spectra of NMR tubes containing PFOB with  $[5\text{a POP}] = 25\text{ mM}$  (red), PFOB  $[5\text{a POP}] = 14.5\text{ mM}$  (green), and pure PFOB (blue) oils. The NMR tubes were embedded in agarose and imaged using the CSI method. Resulting  $^{19}\text{F}$  MRI images are shown (a, right), where shifted  $\text{CF}_3$  peaks are assigned red, green, or blue color channels and superimposed along with the  $^1\text{H}$  grayscale image. Here, red is  $-81.535\text{ ppm}$  with  $[5\text{a POP}] = 25\text{ mM}$ , green is  $-82.178\text{ ppm}$  with  $[5\text{a POP}] = 14.5\text{ mM}$ , and blue is  $-83.048\text{ ppm}$  for pure PFOB. The CSI data were acquired with  $\text{TR/TE} = 26.4/0.53\text{ ms}$ , matrix size  $64 \times 64$ , and 6 averages. (See Supporting Information for additional details.) (b, c) P-PFOB emulsification, forming spheroidal oil droplets, eliminates observed BMS shift. Displayed are  $^{19}\text{F}$  NMR spectra of PFOB and P-PFOB, prepared with dual-capillary NMR samples inside a 5 mm NMR tube, where (b) are oils and (c) formed nanoemulsions. In (b), two sets of peaks are observed, where P-PFOB displays a downfield BMS shift of  $\sim 1.6\text{ ppm}$  with peak broadening. After emulsification (c), only one set of peaks is observed. Capillaries are immersed in  $\text{D}_2\text{O}$  with 0.1% NaTFA as an internal standard.

**Table 1.**

Intramolecular Fluorine-19 Relaxation Rates ( $R_1$  and  $R_2$ ) at 9.4 T of  $\text{Fe}^{3+}$ -SALTAME Chelates **5a**,  $\text{Ga}^{3+}$ -SALTAME Chelate **5a** POP, and Nonchelated Perfluorinated Salicylaldehydes **3a** and **3b**<sup>a</sup>

	isomer	$R_1$ ( $\text{s}^{-1}$ )	$R_2$ ( $\text{s}^{-1}$ )
nonchelated precursors	<b>3b</b>	$0.87 \pm 0.01$	$1.12 \pm 0.01$
	<b>3a</b>	$1.33 \pm 0.01$	$1.75 \pm 0.01$
$\text{Ga}^{3+}$ -SALTAME	<b>5a</b> PPP	$2.36 \pm 0.06$	$5.18 \pm 0.01$
$\text{Fe}^{3+}$ -SALTAME isomers	<b>5a</b> PPP	$146.7 \pm 4.4$	$177.6 \pm 3.9$
	<b>5a</b> POP	$176.9 \pm 3.9$	$219.2 \pm 0.3$
	<b>5a</b> OOP	$227.5 \pm 7.3$	$333.9 \pm 12.0$
	<b>5a</b> OOO	$414.7 \pm 13.7$	$653.0 \pm 14.5$

<sup>a</sup>Chelation with diamagnetic  $\text{Ga}^{3+}$  has only a small effect, whereas chelation with paramagnetic  $\text{Fe}^{3+}$  causes a 2–3 orders of magnitude increase in  $R_1$  and  $R_2$  values. Relaxation rates correspond to 5  $\text{CF}_2$  groups for precursors and  $\text{Ga}^{3+}$  chelate and the 5'  $\text{CF}_2$  group in all  $\text{Fe}^{3+}$  chelates. The  $\pm$  is standard error of the mean (SEM).



**Table 2.**

Intermolecular Fluorine-19 Relaxivities ( $r_1$  and  $r_2$ ) and Susceptibility Shifts of SALTAME 5a POP Chelates with  $\text{Fe}^{3+}$ ,  $\text{Mn}^{4+}$ ,  $\text{Co}^{3+}$ , and  $\text{Ga}^{3+}$  at 9.4 T in PFOB<sup>a</sup>

	<b>Mn<sup>4+</sup></b>	<b>Fe<sup>3+</sup></b>	<b>Co<sup>3+</sup></b>	<b>Ga<sup>3+</sup></b>
$r_1$ (s <sup>-1</sup> mM <sup>-1</sup> )	0.03	0.50	0.001	0.002
$r_2$ (s <sup>-1</sup> mM <sup>-1</sup> )	0.13	1.07	0.15	0.02
shift change (ppm/mM)	0.035	0.060	0.001	0.001

<sup>a</sup>Relaxivities were measured for (CF<sub>2</sub>)<sub>6</sub> fluorine atoms of PFOB, and susceptibility shifts were measured for CF<sub>3</sub> representative fluorine atoms of PFOB. Overall, the SALTAME 5a POP Fe<sup>3+</sup> chelate has the highest relaxivity and susceptibility.

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