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## The internal structure of gadolinium and perfluorocarbon-loaded polymer nanoparticles affects $^{19}\text{F}$ MRI relaxation times†

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$^{19}\text{F}$  magnetic resonance imaging ( $^{19}\text{F}$  MRI) is an emerging technique for quantitative imaging in novel therapies, such as cellular therapies and theranostic nanocarriers. Nanocarriers loaded with liquid perfluorocarbon (PFC) typically have a (single) core–shell structure with PFC in the core due to the poor miscibility of PFC with organic and inorganic solvents. Paramagnetic relaxation enhancement acts only at a distance of a few angstroms. Thus, efficient modulation of the  $^{19}\text{F}$  signal is possible only with fluorophilic PFC-soluble chelates. However, these chelates cannot interact with the surrounding environment and they might result in image artifacts. Conversely, chelates bound to the nanoparticle shell typically have a minimal effect on the  $^{19}\text{F}$  signal and a strong impact on the aqueous environment. We show that the confinement of PFC in biodegradable polymeric nanoparticles (NPs) with a multicore structure enables the modulation of longitudinal ( $T_1$ ) and transverse ( $T_2$ )  $^{19}\text{F}$  relaxation, as well as proton ( $^1\text{H}$ ) signals, using non-fluorophilic paramagnetic chelates. We compared multicore NPs *versus* a conventional single core structure, where the PFC is encapsulated in the core(s) and the chelate in the surrounding polymeric matrix. This modulated relaxation also makes multicore NPs sensitive to various acidic pH environments, while preserving their stability. This effect was not observed with single core nanocapsules (NCs). Importantly, paramagnetic chelates affected both  $T_1$  and  $T_2$   $^{19}\text{F}$  relaxation in multicore NPs, but not in single core NCs. Both relaxation times of the  $^{19}\text{F}$  nucleus were enhanced with an increasing concentration of the paramagnetic chelate. Moreover, as the polymeric matrix remained water permeable, proton enhancement additionally was observed in MRI.

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## Introduction

$^{19}\text{F}$  Magnetic Resonance Imaging ( $^{19}\text{F}$  MRI) is a promising noninvasive and quantitative technique which is in experimental clinical use.<sup>1–3</sup>  $^{19}\text{F}$  MRI is able to unambiguously loca-

lize and also quantify the imaging agent, without an endogenous background, combined with anatomic proton imaging.<sup>1,2,4,5</sup> These features make  $^{19}\text{F}$  MRI very powerful in a broad range of applications, particularly in tracking of cellular therapies or therapeutic nanocarriers, and for the design of activatable probes.<sup>2,3,6–9</sup> The combination of liquid perfluorocarbons (PFCs) and paramagnetic chelates in nanocarriers can provide multifunctional imaging probes. Paramagnetic chelates on the surface of nanocarriers can be used to sense the biological environment of the probe.<sup>10–12</sup> Conversely, fluorophilic chelates that enhance the longitudinal  $^{19}\text{F}$ -relaxation rate increase the sensitivity of  $^{19}\text{F}$  MR signals.<sup>13–16</sup> However, it remains difficult to develop nano-systems in which the paramagnetic chelate modulates both the longitudinal  $^{19}\text{F}$ -relaxation and the proton signal.

Paramagnetic Relaxation Enhancement (PRE) acts over very short distances in a range of angstroms and decreases with the radius  $r$ ,<sup>6</sup> as described by the Solomon–Bloembergen–Morgan equations.<sup>10,14,16,17</sup> Hence, the paramagnetic chelate and the nucleus that should be modulated have to be very close to

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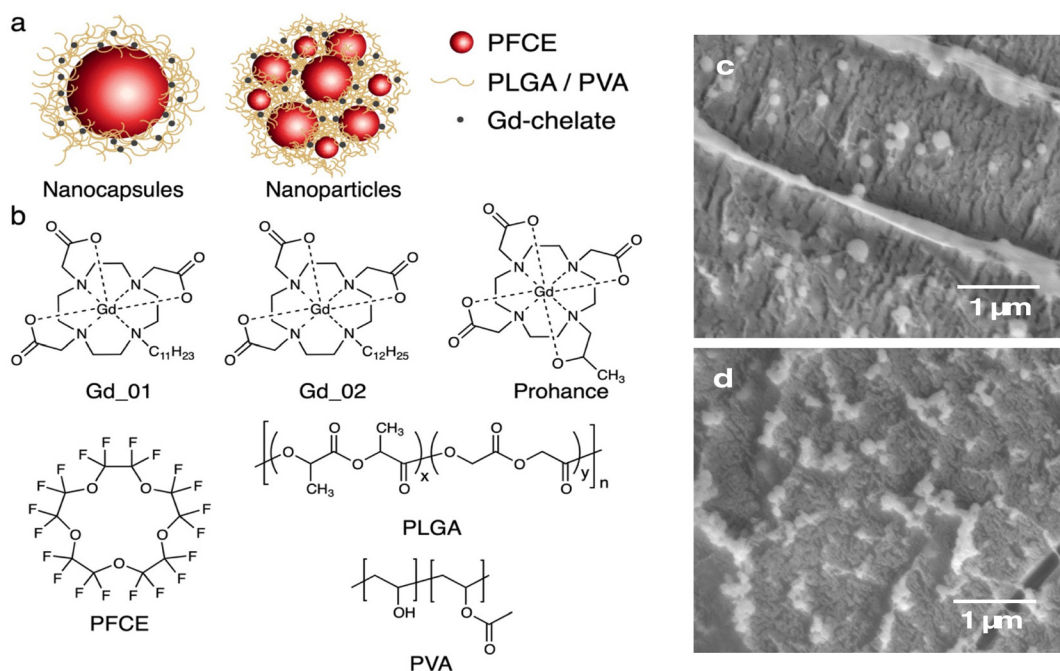
each other, in the range of angstroms.<sup>10,14,16</sup> However, liquid PFC is both hydrophobic and lipophobic.<sup>18–20</sup> As a result, common nanosized <sup>19</sup>F MRI imaging agents are emulsions or single core capsules with a liquid PFC in the core that is stabilized either by a surfactant, polymer or a silica shell.<sup>1,2,4,5,21,22</sup> The core sizes are usually at least several 10s of nanometers.<sup>1,2,4,5</sup> As a result of the single core structure, currently, there are two strategies for the modification with paramagnetic chelates: (1) either covalent attachment or encapsulation of the hydrophilic or hydrophobic chelates in the organic shell of the agent,<sup>12,23,24</sup> and (2) encapsulation of fluorophilic chelates in the fluororous core.<sup>13–15,25</sup>

When a paramagnetic chelate is encapsulated or covalently bound to the shell, it typically affects only a small fraction of <sup>19</sup>F-nuclei that are close to the shell.<sup>23,24</sup> Thus, only modest effects on <sup>19</sup>F longitudinal relaxation rates can be achieved;<sup>23,24</sup> moreover, the signal can become inhomogeneous due to different relaxation rates of <sup>19</sup>F nuclei, depending on their distance from the chelate.<sup>25</sup> On the other hand, this approach enables the modulation of protons in the surrounding medium and detection of other ions.<sup>12</sup> Conversely, the fluorophilic chelates are soluble in the fluororous core; thus, the distance between <sup>19</sup>F nuclei and paramagnetic ions decreases, altering the longitudinal relaxation properties of <sup>19</sup>F nuclei.<sup>13–15,25</sup> Depending on the ion, this strategy can be used to increase the imaging sensitivity of PFC-emulsions.<sup>13,14</sup> However, in such emulsions the paramagnetic nuclei are iso-

lated from the aqueous environment, making simultaneous modulation of the proton signal impossible.

Our group has introduced polymeric PFC-loaded NPs that display a structure different from conventional single core systems, namely the multicore structure (Fig. 1a).<sup>26,27</sup> These NPs contain multiple small cores of perfluoro-15-crown-5 ether (PFCE) with a radius of 10–12 nm that are encapsulated in a matrix of biocompatible and biodegradable poly(lactic-co-glycolic acid) (PLGA) (Fig. 1a and b).<sup>26</sup> This structure is advantageous for *in vivo* and clinical use, as it results in about 15-fold faster clearance of PFCE compared to the PFCE emulsions and it generates stable ultrasound.<sup>28–30</sup> Conversely, emulsions and other single core systems can display organ accumulation times up to several months.<sup>28,29</sup> These NPs can be produced at clinical grade and received approval for a clinical trial.<sup>26,30–33</sup> Thus, combining these systems with paramagnetic probes could open up new opportunities in the development of probes for combined imaging, biosensing and delivery of therapeutics.

Here, we explored how the formation of a multicore structure and the resulting confinement of PFCE affect the PRE. Therefore, we used lipophilic gadolinium (Gd) chelates which were encapsulated in the polymer matrix of the multicore NPs. We compared their relaxation properties with single core NCs. The changes of <sup>19</sup>F-relaxation in core-shell systems were minor, as expected. Conversely, in multicore nanoparticles, we detected enhancement of both *T*<sub>1</sub> and *T*<sub>2</sub> relaxation times,



**Fig. 1** (a) Schematic of nanoparticles used in this study. Left: single core NCs, middle: multicore PFCE-PLGA NPs. (b) Chemical structures of components of NPs. Top row lipophilic chelates Gd\_01 and Gd\_02, and hydrophilic gadoteridol (Prohance). Bottom row: PFCE, PLGA and poly(vinyl alcohol) (PVA). (c and d) Cryogenic scanning electron microscopy (cryoSEM) micrographs of NPs (c) and NCs (d). See also Fig. S1 and S2† for larger images. Scale bar 1  $\mu$ m.



depending on the concentration of encapsulated gadolinium. Furthermore, unlike conventional emulsions, the proton signal could be modulated as well. A further interesting feature shown by multicore NPs was their  $T_1$  and  $T_2$  relaxation variation when the NPs were solubilized in various acidic solutions when compared to neutral pH. This variation was not present in single core NCs. In conclusion, this approach can be adopted for the development of dual  $^1\text{H}/^{19}\text{F}$  MR imaging agents co-loaded with PFC and paramagnetic compounds for different applications, particularly biosensing and enhanced relaxation probes.

## Results and discussion

### Lipophilic chelates can be efficiently encapsulated in both multicore NPs and single core NCs

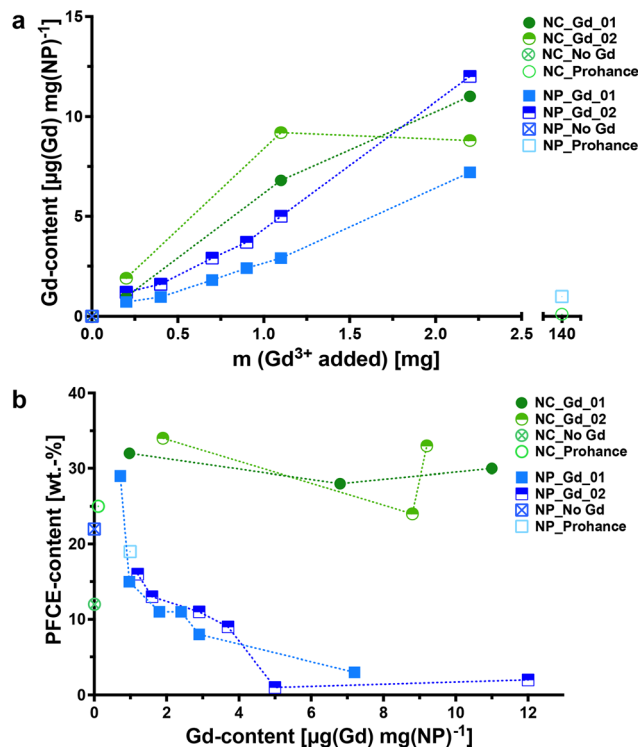
To investigate how structural properties affect the MR properties of NPs co-encapsulating Gd chelates and PFCE, we compared multicore PFCE-PLGA NPs with single core NCs (Fig. 1a). As gadolinium-agents we used two different hydrophobic Gd chelates that differ only in the length of a hydrophobic linker (Fig. 1b).<sup>34</sup> These chelates are hydrophobic but not fluorophilic, ensuring they remain within the hydrophobic PLGA matrix.

Both multicore NPs and single core NCs can be produced by a miniemulsion formulation approach,<sup>26</sup> adjusting the concentration of the lipophilic Gd chelate in the PLGA matrix. This approach varies between NCs and NPs, and this distinction has been thoroughly investigated in previous research.<sup>26</sup> Notably, the internal structure resulting from these distinct approaches was extensively characterized and confirmed using Small-Angle Neutron Scattering (SANS).

Various amounts of the lipophilic chelates, ranging from 0 to 2.4 mg Gd, were employed. Additionally, we included a hydrophilic clinical contrast agent, gadoteridol (Fig. 1b), as a control (Prohance; 140 mg Gd). Post-synthesis, repetitive centrifugation purification steps were performed to remove any free chelate.

Co-encapsulation of the Gd chelates did not impact the NP size (Fig. 1c and d, Tables S1 and S2†). Cryogenic Scanning Electron Microscopy (cryoSEM) images (Fig. 1c and d and Fig. S1 and S2†) revealed spherical NPs with relatively smooth surfaces. Gadolinium-loaded multicore NPs displayed diameters of around 200 nm and a monomodal size distribution (PDI  $\sim 0.1$ ), as determined by Dynamic Light Scattering (DLS, Table S1†). Capsules were slightly smaller with diameters ranging from 150 to 180 nm (Table S2†). Both size and PDI are similar to a control without a Gd chelate, and the variation between different samples is within a typical batch-to-batch variation range.<sup>26</sup>

When lipophilic chelates were used for the synthesis, the concentration of encapsulated Gd increased in both multicore and single core particles with the added amount of the chelate, as shown by inductively coupled plasma-mass spectrometry (ICP-MS, Fig. 2a). The encapsulation of Gd<sub>02</sub>, that



**Fig. 2** Characterization of multicore PFCE-PLGA-NPs and single core PFCE-PLGA-NCs loaded with different Gd chelates (Gd<sub>01</sub>, Gd<sub>02</sub> and Prohance). (a) Gd-content versus mass ( $m$ ) of Gd used for the preparation of NPs determined by ICP-MS. (b) PFCE-content determined by  $^{19}\text{F}$  NMR spectroscopy as a function of the Gd content and type of Gd used. Multicore nanoparticles and single core NCs are indicated in the graphs, respectively, as NP and NC.

has a longer hydrophobic chain, was slightly higher compared to that of Gd<sub>01</sub>. In contrast, the encapsulation of hydrophilic gadoteridol was very low, despite using a nearly 50-fold higher Gd-concentration than the lipophilic chelates (Tables S3 and S4†). PLGA, being a hydrophobic polymer, is well-suited for encapsulating hydrophobic cargo like lipophilic chelates. In contrast, hydrophilic compounds like gadoteridol tend to remain in the aqueous phase during the miniemulsion process. Thus, it is possible to adjust the concentration of hydrophobic Gd chelates during the miniemulsion formulation.

Different trends were observed in the encapsulation of PFCE in multicore NPs and single-core NCs, as revealed by quantitative  $^{19}\text{F}$  NMR spectroscopy (Fig. 2b; indicated as NP\_Gd<sub>01</sub>, NP\_Gd<sub>02</sub>, NP\_No Gd, and NP\_Prohance). In multicore NPs, the PFCE content appeared to decrease with increasing Gd( $m$ ) concentrations, while in NCs, it was even higher compared to capsules without a Gd chelate. It is important to note that the encapsulation of paramagnetic chelates can have varying effects on the MR properties in both particle types, including line broadening, which can affect quantification (discussed further in the next section).

Finally, all NPs and NCs did not show any toxic effects, as demonstrated by a standard viability assay using the same



dose as typically applied for labeling the cells for  $^{19}\text{F}$  MRI with multicore NPs (MTT-assay with RAW macrophages, Fig. S3†).<sup>27,30</sup> The viability values were slightly higher compared to that of a live cell control. This behavior might originate from an increased phagocytic activity of the cells in the presence of NPs and was already observed with other cell types using PFCE-PLGA-NPs without Gd chelates.<sup>30</sup> Hence, the encapsulation of Gd chelates did not affect the toxicity of NPs.

### Confinement of PFC in multicore NPs enables the modulation of fluorine-19 and proton MR signals

After synthesizing Gd-loaded NPs and NCs, we assessed their impact on  $^{19}\text{F}$  nucleus MR properties by measuring  $^{19}\text{F}$  relaxation times using NMR spectroscopy. The relaxation times without Gd chelates were consistent with prior studies.<sup>35</sup> We then examined the effects of Gd encapsulation on longitudinal and transverse relaxation times ( $T_1$  and  $T_2$ ), with the results presented in Fig. 3 (refer to Tables S5–S8† for detailed data).

NMR measurements revealed that the internal structure of NPs significantly influenced  $^{19}\text{F}$  nucleus relaxation properties. In multicore NPs (Fig. 3a and b), longitudinal ( $T_1$ ) relaxation times decreased by nearly 8-fold at the highest Gd-loading

compared to Gd-free NPs. Additionally, transverse relaxation ( $T_2$ ) was notably affected, exhibiting approximately a 255-fold decrease. In single core NCs (Fig. 3c and d), there was a decrease in  $T_2$  relaxation time, albeit much less pronounced, approximately 2-fold, compared to multicore NPs. Furthermore, the encapsulation of Gd chelates induced a modest enhancement in  $T_1$  relaxation time in the single core capsules.

The stark decrease in transverse relaxation time observed in multicore NPs could explain the differences in PFCE quantification between multicore NPs and single core NCs. A shorter  $T_2$  leads to line broadening of the NMR signal (Fig. S4 and S5†), making it challenging to accurately integrate broad peaks within the baseline. As the concentration of encapsulated Gd increases,  $T_2$  relaxation time shortens further, broadening the PFCE peak and causing a decrease in detected PFCE concentration within multicore NPs. This effect did not occur in single core NCs.

What is the reason behind the altered relaxation properties? The effect of paramagnetic ions on the relaxation times of the  $^{19}\text{F}$  nucleus depends on the distance between the ion and the  $^{19}\text{F}$  nucleus.<sup>14,23,24</sup> Previously, we reported the structural

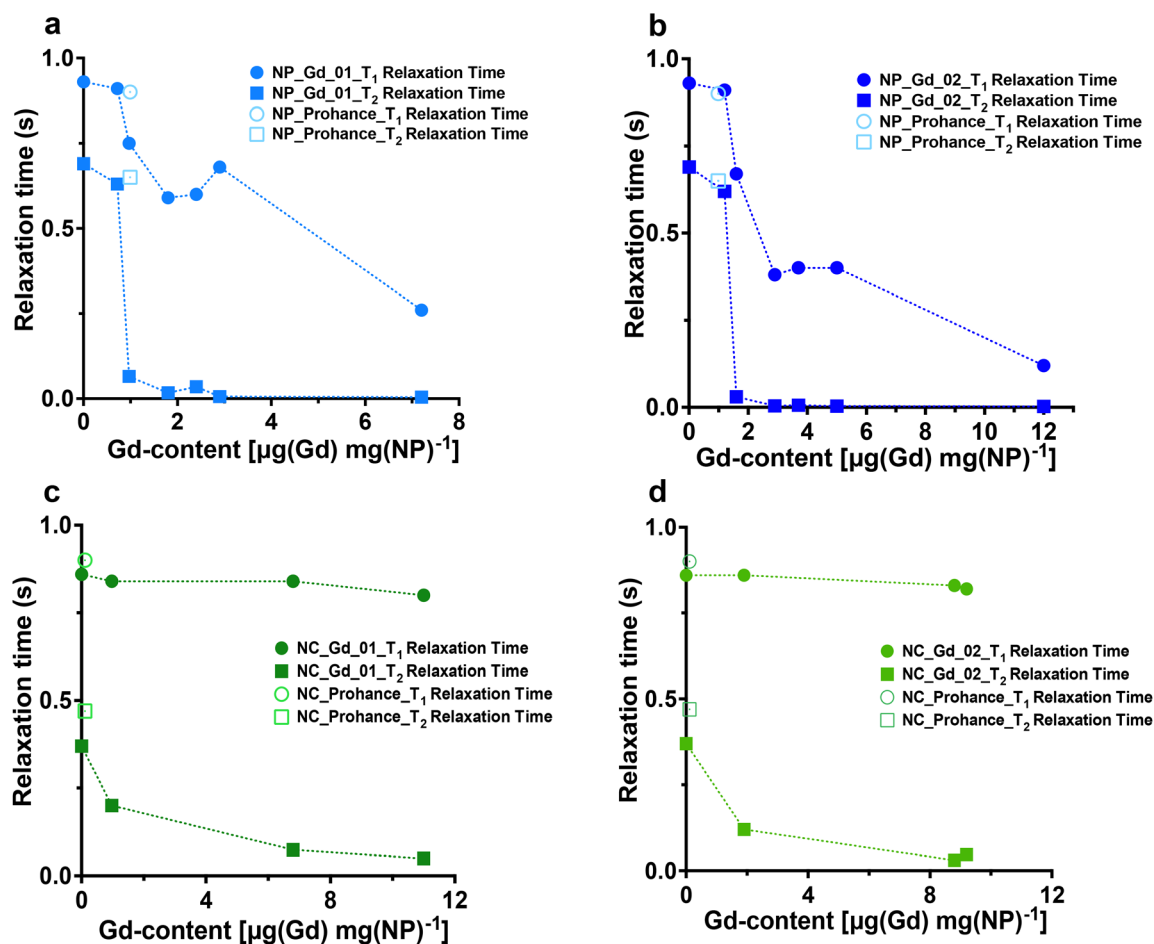


Fig. 3  $^{19}\text{F}$  MR relaxation properties of multicore NPs loaded with increased amount of paramagnetic Gd-chelates compared with single core NCs. Longitudinal and transverse relaxation times of (a) Gd\_01 and (b) Gd\_02-loaded multicore NPs and (c) Gd\_01 and (d) Gd\_02-loaded single core NCs.



characterization of Gd chelate-free and gadoteridol-loaded multicore NPs and single core NCs by Small-Angle Neutron Scattering (SANS).<sup>26</sup> Note that the radii obtained with SANS are typically smaller than the hydrodynamic sizes from DLS, as reported previously.<sup>26</sup> In multicore NPs, each PFCE core typically displays a radius of 9–12 nm and a shell thickness of 4 nm, with some batch-to-batch variation.<sup>26</sup> The single core capsules usually display a core radius of around 20 nm that is surrounded by a shell of a similar thickness.<sup>26</sup>

The distance between Gd and PFCE is shorter in multicore NPs compared to that in NCs, if we assume that Gd chelates are distributed homogeneously within the PLGA matrix. Thus, the farthest possible distance between a PFCE molecule that is located in the center of the core and the Gd chelate in the outer part of the shell is approximately 15 nm. The distance between PFCE in the center of the core and the Gd chelate in the inner part of the shell is 9–12 nm. This distance decreases the closer a PFCE molecule is located to the shell. For the capsules, the farthest distance is 40 nm and the distance between PFCE in the center and the Gd molecule in the inner part of the shell is approximately 20 nm. Thus, this distance is almost twice as high compared to the distance in the multicore NPs. However, PRE typically decreases with the distance  $r^6$ .<sup>14,16,17,10</sup> Hence, shortening the distance alone would not be sufficient to modulate the PFCE molecules located in the center of the core. Thus, the diffusion of PFCE and its confinement in the multicore structure appear to affect the PRE.

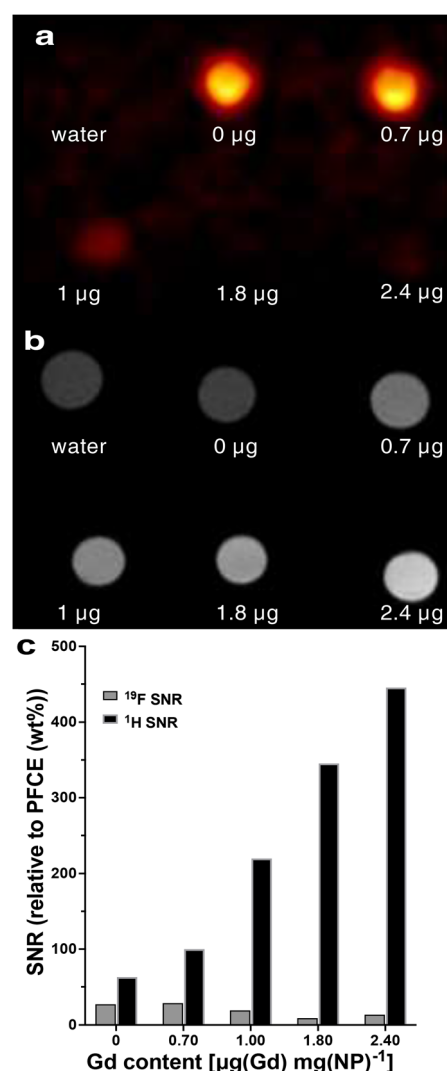
Indeed, other reports also suggested that diffusion of PFCE can influence the PRE. Particularly, several reports showed that conjugation of Gd chelates to the surface of PFCE-loaded silica capsules results in the enhancement of transverse relaxation time.<sup>36–38</sup> These studies did not report the PRE of the longitudinal relaxation time. Importantly, other groups did not observe PRE of longitudinal relaxation in <sup>19</sup>F upon the incorporation of Gd in the surfactant layer of emulsions.<sup>12,23,24</sup> Similarly, in this study we detected only a moderate effect on the longitudinal relaxation times in the control single core NCs, compared with multicore NPs. Thus, the formation of the confining multicore structure affects the enhancement of the longitudinal relaxation time. Typically, the diffusion speed of confined liquids is reduced compared to bulk.<sup>39</sup> However, liquid PFCs display a very low intermolecular cohesiveness,<sup>40</sup> which may invert these effects. Additionally, the distance that a PFCE molecule has to diffuse to relocate from the core center to the shell becomes lower due to the formation of the confining multicore structure. As a result, the PRE of both longitudinal and transverse relaxation times can be observed.

Furthermore, the concentration of Gd within the polymeric matrix influences the PRE. Higher Gd concentrations result in more Gd nuclei in the PLGA shell closer to the PFCE phase, leading to further decreases in both longitudinal and transverse relaxation times in multicore NPs.

Overall, our data show that the structural properties of NPs affect the paramagnetic relaxation of the <sup>19</sup>F nucleus. The PRE effect enhances the PFC nano-emulsion imaging sensitivity and reduces the imaging time. It also enables the development

of stimuli-responsive fluorinated nanoemulsions for on/off switching in response to various stimuli. This switch is achieved through the presence of paramagnetic metal ions, which significantly decrease the  $T_2$  relaxation time, leading to line broadening effects and modulation of the <sup>19</sup>F MR signal for responsive imaging.

Finally, <sup>1</sup>H/<sup>19</sup>F MRI demonstrated that the presence of paramagnetic chelates affected both the fluorine (Fig. 4a) and proton signals in multicore particles (Fig. 4b). As the concentration of Gd increased, the <sup>19</sup>F signal intensity decreased due to the reduction in  $T_2$  relaxation time (Fig. 4a and c). Conversely, the proton signal increased with higher chelate encapsulation (Fig. 4c). The signal-to-noise-ratio of the proton signal increased from 17 to 49, when the Gd content increased from 0 to 2.4  $\mu\text{g}$  per mg NPs. In multicore particles, Gd is



**Fig. 4** (a) <sup>19</sup>F and (b) <sup>1</sup>H MRI of multicore NPs that contain a different amount of Gd chelates. The Gd content in  $\mu\text{g}$  per mg of NPs is indicated under each sample. Water and NPs loaded only with PFCE were used as controls. (c) The signal-to-noise ratio of <sup>19</sup>F and <sup>1</sup>H MRI signals for different Gd contents. (See Table S9† for the signal-to-noise values).



encapsulated within the PLGA matrix, which retains hydration and permeability to water and small molecules.<sup>26</sup> This property allows for the modulation of the proton signal.

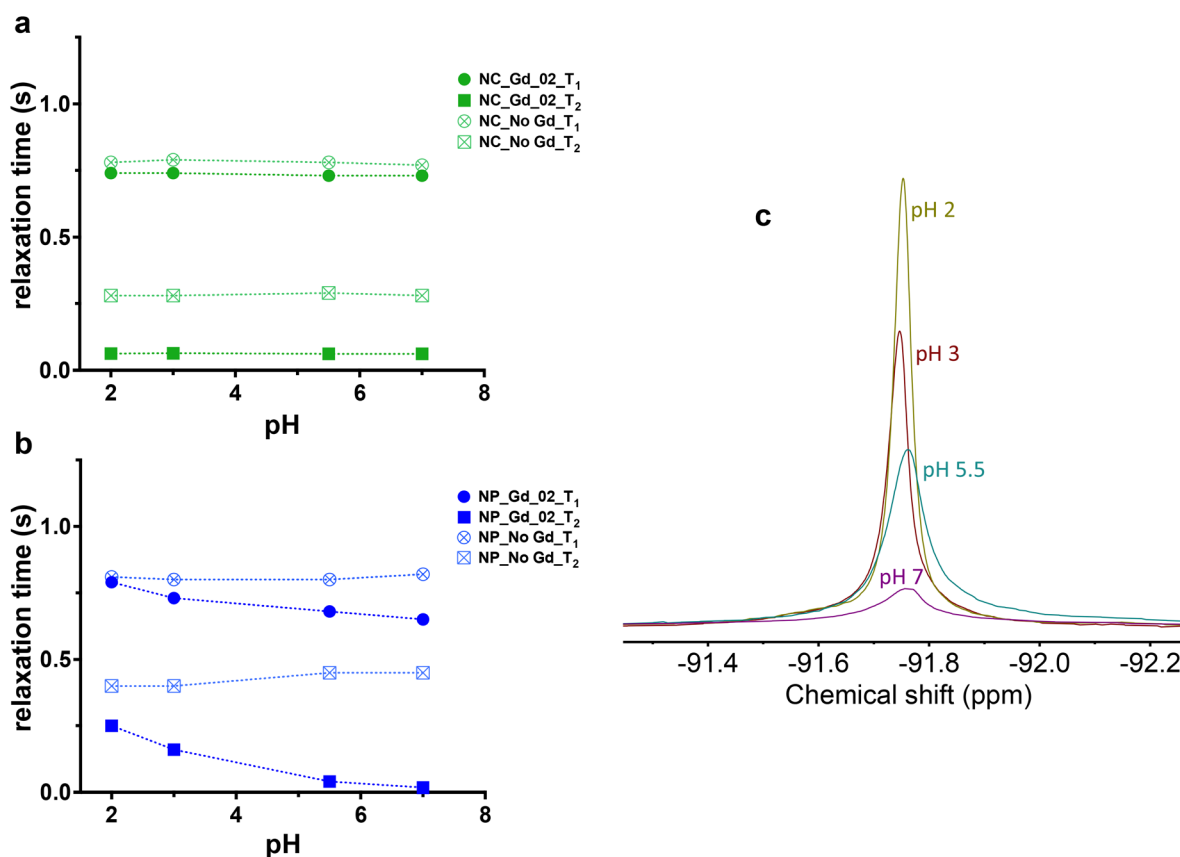
The encapsulation of paramagnetic chelates in multicore NPs allows for the modulation of both  $T_1$  and  $T_2$  relaxation times of the  $^{19}\text{F}$  nucleus, along with the proton signals from the aqueous environment of the NPs. In emulsions, it is typically possible to efficiently modulate either the proton or the fluorine relaxation, but not both simultaneously. When paramagnetic chelates are attached to the shell, they primarily alter  $^1\text{H}$  relaxation, with a minor effect on the PRE of  $^{19}\text{F}$  nuclei in the fluorinated phase.<sup>23</sup>

In contrast, the fluorophilic chelates are dispersed in the PFC phase, shortening the  $^{19}\text{F}$  relaxation times, but remain inaccessible to water.<sup>14</sup> Hence, only the PRE of the  $^{19}\text{F}$ -signal is obtained. In contrast to emulsions, co-encapsulation of PFC and Gd chelates in multicore NPs enables modulation of both  $^{19}\text{F}$  and  $^1\text{H}$  signals. Here we used gadolinium(III) chelates, which are known to cause a strong PRE of transverse relaxation. In future studies, these effects could be reduced by using other paramagnetic ions, similarly as Ahrens' group reported for fluorophilic chelates.<sup>13,15</sup>

### Acidic environment impacts the relaxation times in multicore NPs but not in single core NCs

As  $^{19}\text{F}$  paramagnetic relaxation enhancement (PRE) has been shown to be an efficient tool for the development of nano-systems with an on/off switching mechanism based on specific stimuli, we explored if multicore NPs can respond to a pH variation.<sup>36,38,41,42</sup> Considering that the PRE-effect is concentration dependent, the Gd content for both single core and multicore NPs was kept constant for all measurements and samples were analyzed in environments with varied pH.

Multicore NPs showed a more pronounced effect of pH compared with single core capsules (Fig. 5). Indeed, in addition to the negligible influence of Gd-loaded single core NCs on  $T_2$  at pH 7 when compared with the control NPs (Gd-free NCs), there was no impact on  $T_1$  and  $T_2$  relaxation times as a function of different pH used for both Gd\_01 and Gd\_02 (Fig. 5a; Tables S12 and S13†). While  $^{19}\text{F}$  relaxation properties of NCs remained almost constant, in multicore NPs, the relaxation times varied (Fig. 5b) from neutral to the most acidic environment. Thus, the transverse relaxation time increased gradually from pH 7 to pH 2 approximately 15-fold (Tables S10



**Fig. 5**  $^{19}\text{F}$  MR relaxation properties in the acidic environment of multicore NPs encapsulated with Gd-chelates compared to single core capsules. Longitudinal and transverse relaxation time of (a) Gd\_02-loaded single core and (b) multicore NPs and related (c)  $^{19}\text{F}$  spectral line profile of multicore NPs under different pH environments,  $c = 10 \text{ mg mL}^{-1}$ , 500 MHz. When both type of NPs are loaded with Gd\_01, the effect is the same (Table S10† for multicore nanoparticles loaded with Gd\_01 and Table S12† for core-shell nanocapsules loaded with Gd\_01, results not included in the graphs for clarity).



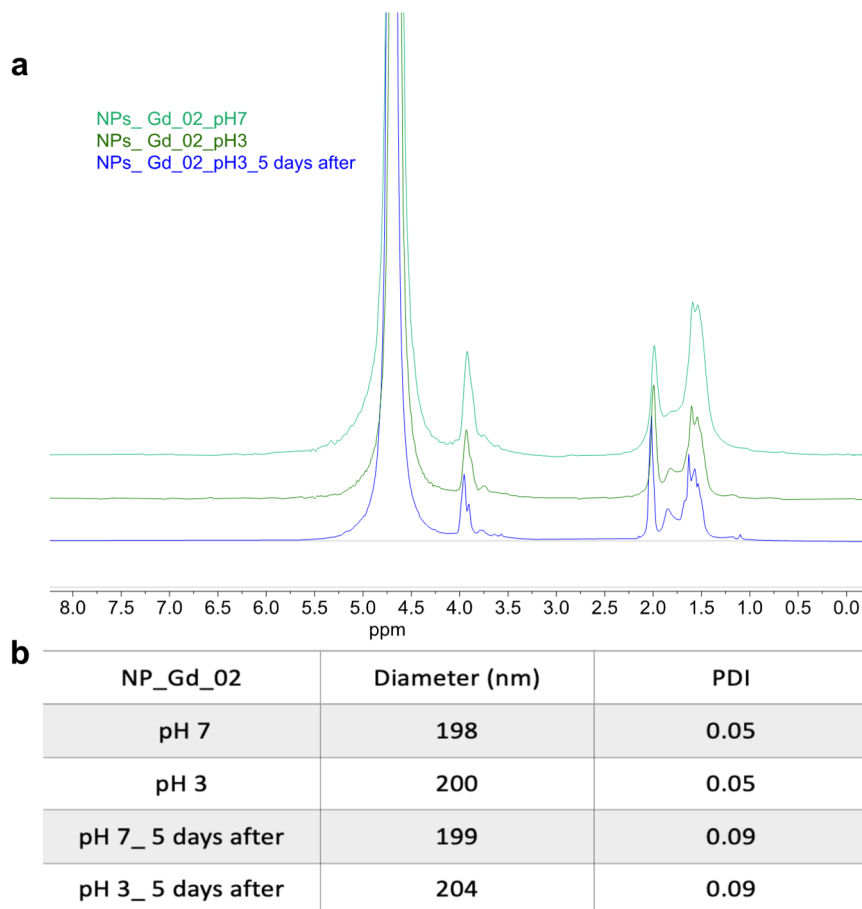
and S11†) for both Gd\_01 and Gd\_02, indicating that the acidic solution can penetrate more easily into the multicore NP matrix as compared to single core NCs. This result is manifested by the sharp increase in the  $^{19}\text{F}$ 's line-width as expected with an increase in  $T_2$  relaxation time (Fig. 5c). A minor change was observed in the  $T_1$  relaxation time of Gd\_01 and Gd\_02 encapsulated multicore NPs. This is attributed to the fact that Gd chelates are primarily  $^{19}\text{F}$   $T_2$  relaxation agents.<sup>14</sup>

**Table 1** Transverse relaxation time of solvent protons and fluorine-19.  $^1\text{H}$  and  $^{19}\text{F}$  transverse relaxation times at pH 7 and pH 3 of NPs not encapsulated by Gd (NPs\_No Gd), NPs loaded with Gd (NPs\_Gd) and NPs without Gd but supplemented with  $\text{GdCl}_3$  salt to mimic Gd linkage

	$^1\text{H}$ $T_2$ (S)		$^{19}\text{F}$ $T_2$ (S)	
	pH 7	pH 3	pH 7	pH 3
D <sub>2</sub> O (Deuterium oxide)	2.6	4.2	n/a	n/a
NPs_No Gd	1.1	2.7	0.45	0.49
NPs_Gd	0.18	0.36	0.02	0.21
NPs_No Gd + $\text{GdCl}_3$	0.12	0.36	0.12	0.50

It is well known from the literature<sup>43</sup> that in strongly acidic solutions protons compete with  $\text{Gd}^{3+}$  to bind to a carboxylate ligand, thus increasing the dissociation rate of  $\text{Gd}^{3+}$  complexes. Therefore, it can be hypothesized that such a substantial change in  $T_2$  from pH variation is explained by the fact that a low pH environment might trigger the release of Gd from the chelate into the aqueous phase. Such release can be problematic, as the primary reason for toxicity caused by Gd-based contrast agents has been identified to be the dissociation of Gd from the chelated complex *in vivo*.<sup>44</sup> As  $\text{Gd}^{3+}$  ions have a comparable ionic radius to  $\text{Ca}^{2+}$ , they can interfere with calcium-mediated signaling pathways, thus making them exceedingly toxic.

To understand the mechanism of  $T_2$  variation of multicore NPs at different pH, we studied the transverse relaxation time of the residual solvent protons and fluorine-19 at pH 7 and 3 (Table 1). Three different samples were analyzed: (i) multicore NPs with no gadolinium (NPs\_No Gd), (ii) NPs loaded with Gd chelates (NPs\_Gd), and (iii) NPs without Gd but supplemented with gadolinium chloride (NPs\_No Gd +  $\text{GdCl}_3$ ), used as a positive control to mimic Gd leakage from the NPs. Proton  $T_2$  relaxation time emerged to be equal in both NPs loaded with



**Fig. 6** (a)  $^1\text{H}$  NMR spectrum of multicore NPs loaded with Gd\_02 at pH 7 and 3, immediately after dispersing them in D<sub>2</sub>O (pH 7) and acidic solvent (pH 3), and five days after, 700 MHz,  $c = 10 \text{ mg mL}^{-1}$ . (b) The stability of the NPs is also demonstrated by the DLS values at pH 7 and 3. The stability was also confirmed for the multicore NPs loaded with Gd\_01. (Results not included for clarity, Fig. S7†).



Gd and the ones supplied with salt, indicating that the PLGA matrix of the NPs, which additionally incorporates PVA,<sup>26</sup> is permeable to water or at least to water protons. The exchange must be quite fast as the impact of the encapsulated Gd and the one supplemented with GdCl<sub>3</sub> is negligible on  $T_2$  relaxation time. However, it is very unlikely that Gd<sup>3+</sup> is permeable to the NPs as the <sup>19</sup>F  $T_2$  does not change notably in the presence of the added Gd-based salt. Only encapsulated Gd has an enhanced impact (10-fold), which in turns further proves that encapsulation additionally provides a clue that Gd<sup>3+</sup> does not dissociate under acidic conditions. If this was the case, then one would expect that <sup>19</sup>F transverse relaxation time of NPs encapsulating Gd equals the one of NPs supplemented with gadolinium salt. Additionally, it is interesting to note that the <sup>19</sup>F  $T_2$  value of NPs\_No Gd + GdCl<sub>3</sub> at pH 3 is really close to that of NPs\_No Gd than to NPs\_Gd demonstrating that in the acidic environment Gd<sup>3+</sup> is not released from the chelate. These results suggest that the changes in the internal conformational structure change of NPs leads to a change in  $T_2$  relaxation time in the acidic environment.

A further possible mechanism which drives the variation of  $T_2$  values at different pH might be related to the stability of the NPs: at low pH the ester bonds in the polymer backbone typically hydrolyze. However, it can be distinctly observed that following dispersion of NPs, there is no difference in the <sup>1</sup>H spectrum (Fig. 6a) between NPs in water (pH 7) compared to the ones in acidic solution (pH 3), even after 5 days, suggesting slow and negligible hydrolysis during this period. Moreover, the stability of the NPs was also confirmed by DLS. Indeed, size and PDI remain constant at neutral and acidic pH (Fig. 6b).

These data might be explained by a morphology change inducing an increase in  $T_2$  relaxation time and signal intensity in acidic medium. Hence, it has been previously shown that in acidic pH PLGA particles have a different morphology than in alkaline and neutral pH.<sup>45,46</sup> Both articles state that PLGA particles incubated in the acidic environment present fewer pores on the surface and inside compared to those under neutral pH. However, the particles at acidic pH keep having a smooth surface for a long period of time with a minimal loss in mass of PLGA particles. Therefore, we can hypothesize that this evident morphology alteration might push the Gd chelates farther from fluorine-19 atoms than the original morphology and hence increase the <sup>19</sup>F  $T_2$  relaxation time.

The stability of the nanoprobe was also studied in terms of  $T_2$  values over time. For this purpose, the  $T_1$  and  $T_2$  variations of the system at pH 3 were measured right after and 3 hours after the solubilization (Fig. 7b). Fig. 7a shows the system stability in terms of signal intensity.

When multicore NPs were dispersed in a basic environment (pH 14), the PLGA was hydrolyzed and in the presence of Gd<sup>3+</sup>, it resulted in a complete line broadening of the <sup>19</sup>F NMR resonances (Fig. S6†). However, for NPs prepared without Gd<sup>3+</sup>, the high pH environment did not significantly impact the <sup>19</sup>F line-shape (Fig. S6†), in agreement with our earlier results.<sup>35</sup> Similarly, the DLS analysis showed an increase in size from

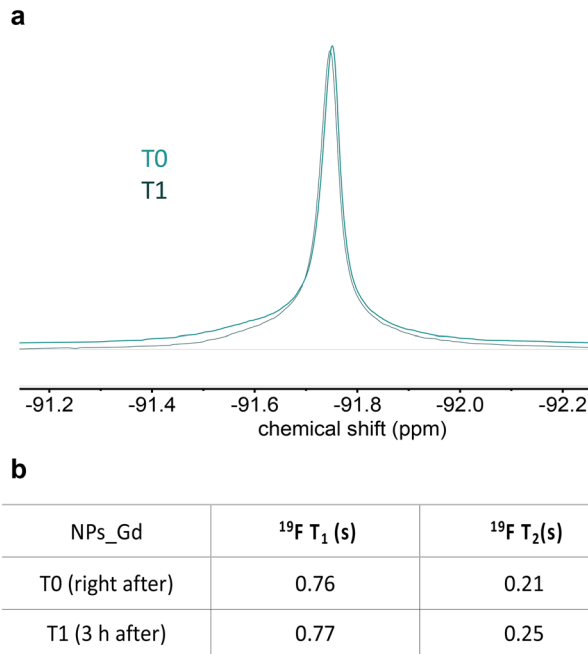


Fig. 7 (a) <sup>19</sup>F spectral line profile of multicore NPs at pH 3 in two time-points and (b) related <sup>19</sup>F MR relaxation properties. Multicore NPs:  $c = 10 \text{ mg mL}^{-1}$ , 500 MHz.

164 nm at pH 7 to 1300 nm at basic pH and in the PDI value from 0.05 to 0.4. As reported previously, NPs degrade at alkaline pH leading to the formation of agglomerates of the remaining surfactant and PFC.<sup>35</sup>

## Conclusion

A co-encapsulation of PFC and a paramagnetic chelate in NPs can lead to probes suitable for high-sensitive <sup>19</sup>F MRI or sensing application. Yet, the modulation of both the <sup>19</sup>F-signal of PFC inside the NPs and the proton signal from the aqueous environment of the probe remained not possible. We explored the impact of the nanoparticle structure on paramagnetic relaxation enhancement (PRE) and compared liquid PFC-loaded PLGA nanoparticles with multicore and single-core structures. Our findings highlight the crucial role of structural properties in <sup>19</sup>F MR relaxation. In single-core capsules, the effect of the paramagnetic chelate was primarily observed as an increase in  $T_2$  relaxation time. Interestingly, multicore nanoparticles exhibited decreases in both <sup>19</sup>F  $T_1$  and  $T_2$  relaxation time with increasing Gd chelate encapsulation, and they also allowed for the modulation of the proton signal. This approach holds promise for fine-tuning both <sup>19</sup>F and <sup>1</sup>H MR characteristics in agents containing liquid PFC and paramagnetic chelates, particularly for high-sensitivity probes and environmentally responsive agents. Finally, we showed that the multicore and single core structures of the NPs lead to different <sup>19</sup>F relaxation times under different pH conditions. The <sup>19</sup>F MR relaxation was not affected in single core NCs irre-





spective of the pH. However, an increase in  $T_2$  relaxation time has been observed in multicore NPs at the most acidic pH analyzed despite maintaining the physiochemical stability and integrity. In conclusion, multicore NPs can also be applied as a pH probe for acidic environment detection.

## Materials and methods

### Materials

All chemicals were used as received: PLGA (Resomer 502H) from Evonik Industries, Germany; dichloromethane (DCM) from EMSURE, Merck Millipore, Darmstadt, Germany; polyvinyl alcohol (PVA), deuterium oxide and trifluoroacetic acid (TFA) were purchased from Sigma-Aldrich (St Louis, MO, USA); perfluoro-15-crown-5 ether (PFCE) from Exflour, Texas, United States, and Prohance from Bracco, Amsterdam, The Netherlands. Lipophilic gadolinium chelates were synthesized as described previously.<sup>34</sup> Water was purified by a Synergy water purification system from Merck.

### Methods

**Synthesis of multicore nanoparticles.** PVA (0.5 g) was dissolved in ultrapure water (25 g) under magnetic stirring. PLGA (0.1 g) was dissolved in dichloromethane (3 mL) and mixed with PFCE (0.9 mL) by pipetting it up and down with a glass pipette. In parallel, stock solutions of Gd-chelates in dichloromethane were prepared ( $c = 0.0166 \text{ mg } \mu\text{L}^{-1}$ ). Different amounts of these chelates were added to the organic phase. For preparation of particles with gadoteridol, Prohance (1.78 mL, corresponds to 497 mg gadoteridol and 140 mg Gd) was added to the organic phase and sonicated for 15 s. Afterwards, the sonication of the aqueous surfactant solution was started, and the organic phase was added rapidly with a glass pipette (microtip Branson digital Sonifier s250, Missouri, United States). The miniemulsion was sonicated in an ice-water bath for 3 minutes at 40% amplitude. As PFCE is both hydrophobic and lipophobic, the organic phase has to be constantly premixed by pipetting during the addition to ensure that the PFCE and PLGA phases are added simultaneously. After sonication, DCM was evaporated overnight at room temperature under stirring to achieve solidification of the particles. The dispersions were transferred to 50 mL tubes and washed four times by centrifugation at 16 000g (4 °C, 35 min) and supernatant replacement. After the second washing step, the samples were sonicated in an ultrasonic bath (sonication bath, Diagenode Bioruptor, Seraign, Belgium) and the washing cycle was repeated. After washing, the particles were resuspended in water (5 mL), frozen with liquid nitrogen and freeze-dried. The resulting powder was stored at  $-20 \text{ }^\circ\text{C}$ .

**Synthesis of single core nanocapsules.** For the synthesis of NCs, sodium cholate (0.375 g) that was used as a surfactant in the sonication step was dissolved in water (24.625 g, 1.5 wt% solution). PLGA (0.1 g) was dissolved in dichloromethane (2 mL) and mixed with PFCE (0.9 mL) by pipetting it up and down with a glass pipette. Different amounts of these chelates

were added to the organic phase (corresponding to 1 mg, 5 mg and 10 mg of chelate). Some samples were prepared with the addition of Prohance (1.78 mL). The organic phase was then added with a glass pipette to the sodium cholate solution. As PFCE is both hydrophobic and lipophobic, the organic phase had to be premixed before sonication to ensure that PFCE and PLGA phases are added simultaneously. The miniemulsion was sonicated in an ice bath for 3 minutes at 40% amplitude (microtip Branson digital sonifier s250, Missouri, United States). After sonication, DCM was evaporated overnight at room temperature under stirring to achieve solidification of the particles. To exchange the surfactant, 10 g of PVA solution (1.96 wt%) was added to the suspension, followed by continuous stirring at 4 °C for 4 days. After 4 days, the capsules were washed 4 times with deionized water through centrifugation (16 098g, 35 minutes, 4 °C), resuspended in 5 mL of water, frozen (liquid  $\text{N}_2$ ) and freeze-dried. The NCs in a powder form were stored at  $-20 \text{ }^\circ\text{C}$ .

### Inductively coupled plasma-mass spectrometry (ICP-MS).

The Gd content was determined by ICP-MS using an X series I with a quadrupole mass spectrometer. Samples were prepared by adding 10 mg of the nanoparticles to 145  $\mu\text{L}$   $\text{HNO}_3$  (1%) and left overnight at room temperature in order to let the particles fully degrade. Afterwards, the samples were filled up to a total volume of 10 mL prior to the measurement.

**Dynamic light scattering (DLS).** The particle size and the polydispersity index (PDI) were measured by DLS on a Malvern Zetasizer ZS nano instrument. The particles were dispersed in ultrapure water at a concentration of  $0.01 \text{ mg mL}^{-1}$ .

**$^{19}\text{F}$  NMR experiments.** They were performed either with a Bruker Avance III HD 400 MHz nanobay spectrometer equipped with a 5 mm BBFO + probe at 298 K (Gd<sup>3+</sup> loading relaxation experiments) or a JEOL ECZ-R 500 MHz spectrometer equipped with a RoyalHFX probe (pH effects on relaxation experiments).

Determination of PFCE content was done by quantitative  $^{19}\text{F}$  NMR. The NPs and NCs were resuspended in 400  $\mu\text{L}$  of  $\text{D}_2\text{O}$  and 100  $\mu\text{L}$  of 0.1 vol% trifluoroacetic acid (TFA) as an internal reference. The interscan delay d1 was set to  $25 (5 \times T_1)$ .

For relaxation time measurements approximately 5 mg of particles were resuspended in  $\text{D}_2\text{O}$  and transferred to an NMR tube. The  $^{19}\text{F}$  90° pulse was calibrated prior to the measurement. The same method was used for the pH effect experiments except for the solvent which was a  $\text{D}_2\text{O}$  solution of hydrogen chloride with the desired pH. For the latter experiment, the nanoparticles were resuspended prior to the measurement and  $T_1$  and  $T_2$  measured as described below.

$T_1$  measurements were done using the inversion recovery method of a 180° pulse followed by a variable delay period for transverse magnetization to relax longitudinally followed by a 90° detection pulse. The  $T_1$  was first estimated using a simple 1D sequence and then a full pseudo-2D acquisition was acquired with a list of delays with an interscan delay set to  $5 \times T_1$ , the slowest. The recovery was fit to an exponential growth function which revealed the  $T_1$ . The  $^{19}\text{F}$   $T_1$  experiments were performed without  $^1\text{H}$  decoupling.



$T_2$  measurements were performed using the CPMG method of a  $90^\circ$  pulse followed by a train of tau-180-tau, where tau is a variable delay. For rapid  $T_2$ -relaxing nuclei ( $T_2 < 30$  ms), a tau of 0.5 ms was used. For other, slower  $T_2$ -relaxing nuclei, a tau of 1.2 ms was used.  $T_2$  was first estimated using a simple 1D sequence. Afterwards, a full pseudo-2D acquisition was acquired with a list of refocusing pulse repetitions ranging from 2 to 1750 and an interscan delay set to  $5 \times T_1$ , the slowest. The recovery was fit to a mono-exponential decay function which revealed the  $T_2$ .

The same setting was used to measure  $T_1$  and  $T_2$  relaxation times in an acidic environment. In this regard, roughly 5 mg of nanoparticles from the same sample were resuspended in 500  $\mu\text{L}$  of several acidic solutions with pH 2, 3, 5.5 and 7. Gd-free PFCE nanoparticles were used as a control.  $T_1$  and  $T_2$  measurements were performed right after the resuspension.

**Cryogenic scanning electron microscopy (CryoSEM).** The images were obtained with a JEOL 6300F field emission scanning electron microscope. The samples (8  $\mu\text{L}$ , 10 mg  $\text{mL}^{-1}$ ) were pipetted in 2 rivets that are placed together. Then the sample was frozen in liquid nitrogen slush and with a cryo transfer device placed in an Oxford Alto 2500 cryo-station. There the top rivet was broken and the sample was heated to  $-95^\circ\text{C}$  for 5 minutes. Then a coating of 60/40 Au/Pd was done and the sample was transferred to the cryoSEM.

**$^1\text{H}$  and  $^{19}\text{F}$  magnetic resonance imaging.** The imaging was carried out on a Bruker Biospec 11.7 T. The samples had Gd contents of 0, 0.7, 0.9, 1.8 and 2.5  $\mu\text{g mg}^{-1}$ . The concentration of nanoparticles was 10 mg  $\text{mL}^{-1}$ , which is also often used for *in vivo* imaging.  $^1\text{H}$  MRI was acquired with the  $T_1$ -weighted 2D fast low angle shot (FLASH) sequence: TR 104 ms, TE 2.25 ms, flip angle  $40^\circ$ , 2 averages, matrix  $256 \times 256$ , field of view (FOV)  $32 \times 32$  mm, 12 slices with a slice thickness of 1 mm, scan time 53 s.  $^{19}\text{F}$  MRI was performed in a 3D RARE sequence with the following parameters: TR 1 s, TE 6.5 ms, turbo factor 8, averages, matrix  $32 \times 32 \times 12$ , FOV  $32 \times 32 \times 24$  mm, scan time 6 : 20 min.

**Cell labelling and viability.** A murine macrophage cell line (RAW cells) was cultured in RPMI medium (Gibco, Merelbeke, Belgium) enriched with 1% antimycotic-antibiotic (AA) (Gibco, Merelbeke, Belgium) + 10% foetal bovine serum (FBS). Cells were incubated at  $37^\circ\text{C}$  for 24 hours, with nanoparticles at the concentration of 2 mg/ $1.0 \times 10^6$  cells. To investigate the influence of labelling on cell viability, an MTT assay was performed. For this purpose, the excess of label was removed, and cells were gently washed and resuspended in 100  $\mu\text{L}$  of PBS. For the negative control, 100  $\mu\text{L}$  of DMSO was added to cells instead of PBS and all the samples were placed in a 96-well plate. Next, 60  $\mu\text{L}$  of complete medium and 10  $\mu\text{L}$  of thiazolyl blue tetrazolium bromide (MTT) solution (4 mg MTT in 1 mL sterile water) (Sigma-Aldrich, Missouri, USA) was added to each well, followed by 1-hour incubation at  $37^\circ\text{C}$ . After incubation, the plate was centrifuged for 2 minutes and 100  $\mu\text{L}$  lysis buffer (9 mL isopropanol, 12.5  $\mu\text{L}$  SDS, 200  $\mu\text{L}$  of 2 N HCL, and 787.5  $\mu\text{L}$  of deionized water) was added per well and the plates were incubated for 15 minutes in the dark at RT.

The absorbance was then measured using an iMark<sup>TM</sup> microplate absorbance reader (Bio-rad, California, USA) at 590 nm.

## Author contributions

The experiments were designed by Alvja Mali, Margot Verbeelen, Paul B. White, Olga Koshkina and Mangala Srinivas. The experiments were conducted by Alvja Mali, Margot Verbeelen, Paul B. White, Alexander H.J. Staal, N. Koen van Riessen, and Cyril Cadiou. All the authors assisted with the data analysis and interpretation of the results. All authors contributed to the final draft. Alvja Mali and Margot Verbeelen contributed equally to this work.

## Conflicts of interest

The authors have no conflicts of interest to declare.

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