Synthesis Of PEG-Coated, Ultrasmall, Manganese-Doped Iron Oxide Nanoparticles With High Relaxivity For $T_1$/$T_2$ Dual-Contrast Magnetic Resonance Imaging

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**Background:** Beyond magnetic resonance imaging (MRI), which has been widely used clinically, molecular MRI (mMRI) can further provide qualitative and quantitative information at the cellular and molecular levels. However, the diagnostic accuracy may not be satisfactory via single-contrast mMRI due to some interferences in vivo. $T_1$/$T_2$ dual-contrast MRI using the same contrast agent (CA) could significantly improve the detection accuracy. Therefore, in this study, we fabricated poly(ethylene glycol) (PEG)-coated, manganese-doped iron oxide nanocomposites (Mn-IONPs@PEG) as $T_1$/$T_2$ dual-contrast CA, and evaluated its feasibility of $T_1$/$T_2$ dual-contrast MRI in vitro and in vivo.

**Methods:** Mn-IONPs were prepared by the thermal decomposition of iron-erucate and manganese-oleate complexes and were coated with 1,2-Distearoyl-sn-glycero-3-phosphoethanolamine-N-(methoxy[polyethylene glycol]-2000) (DSPE-PEG 2000). The physico-chemical properties and cytotoxicity of the Mn-IONPs were fully characterized, followed by MRI in vitro and in vivo.

**Results:** Ultrasmall 3 nm-sized nanoparticles were successfully prepared and were identified using transmission electron microscopy (TEM), high-resolution TEM, and X-ray diffraction. After coating with DSPE-PEG, the Mn-IONPs@PEG displayed excellent hydrophilicity and good biocompatibility. Due to the manganese-doping and PEG coating, the Mn-IONPs@PEG showed good relaxivity in vitro. Especially, the Mn-IONPs@PEG coated with DSPE-PEG following a mass ratio to Mn-IONPs of 1:2 showed harmonious longitudinal relaxivity ($r_1 = 7.1$ mM$^{-1}$s$^{-1}$) and transversal relaxivity ($r_2 = 120.9$ mM$^{-1}$s$^{-1}$), making it a better candidate for $T_1$/$T_2$ dual-contrast mMRI. After administrated via a caudal vein, the Mn-IONPs@PEG can induce significant enhancement in both $T_1$-weighted and $T_2$-weighted MR images and the time at 10 mins after injection was regarded as a suitable time for imaging because both the $T_1$ and $T_2$ enhancement were optimum at that time.

**Conclusion:** The obtained Mn-IONPs@PEG exhibited good $r_1$ and $r_2$ and was a reasonable candidate for $T_1$/$T_2$ dual-contrast mMRI.

**Keywords:** magnetic resonance imaging, manganese-doped iron oxide nanoparticles, $T_1$/$T_2$ dual-contrast, DSPE-PEG coating, high relaxivity

**Introduction**

In recent decades, magnetic resonance imaging (MRI) has played an irreplaceable role in medical diagnosis because of its efficient soft tissue contrast, high spatial resolution, low of ionizing radiation, and unrestricted signal penetration depth.\(^1\)\(^2\)
Particularly, molecular magnetic resonance imaging (mMRI) has extended beyond traditional MRI technology. mMRI can obtain qualitative and quantitative information about biological processes in vivo at the cellular and molecular levels, greatly expanding the research and application fields of MRI. In recent years, mMRI has been intensively studied in gene imaging, cell tracing, drug screening, and evaluating early diagnoses and therapeutic effects of disease. Contrast agent (CA) serving as the signal source of mMRI is the most important ingredient and has reasonably become a hot spot in the field of mMRI.

Although many types of CA based on magnetic nanoparticles (NPs) with high longitudinal relaxivity ($r_1$) or transverse relaxivity ($r_2$) have been developed, single-contrast mMRI still possesses insufficient sensitivity and specificity due to its innate properties. Transverse relaxation time ($T_2$) single-contrast CA usually results in MR signal loss and susceptibility artifacts. Additionally, the darker areas in MR $T_2$-weighted imaging ($T_2$WI) induced by the $T_2$ single-contrast CA may be confused with other physiological or pathological low-signal regions, such as bleeding, calcification, and metal deposition, which will significantly reduce the sensitivity and specificity of mMRI. Similarly, some normal tissues (such as fatty tissue) with hyperintensity in MR longitudinal relaxation time ($T_1$)-weighted imaging ($T_1$WI) may be misunderstood as bright lesions enhanced by $T_1$ CA. $T_1/T_2$ dual-contrast MRI using the same CA provides complementary information about a lesion to self-confirm and provide fault-free MR images, proving that it can significantly improve the detection accuracy.

Thus far, multifarious CAs have been reported for $T_1/T_2$ dual-contrast MRI. A common strategy for the synthesis of $T_1/T_2$ dual-contrast CA is to integrate $T_1$ and $T_2$ signal moieties as hybrid nanoparticles. However, due to the magnetic coupling effect, the enhancement generated by the $T_1$ and $T_2$ moieties will interact with each other, leading to the compromise of both $r_1$ and $r_2$. Embedding the $T_1$ species into the $T_2$ species can parallel the electron spins of paramagnetic $T_1$ CA with the direction of the magnetic field induced by the $T_2$ CA, and reasonably enhance both the $T_1$ and $T_2$ relaxivity of the $T_1/T_2$ dual-contrast CA.

Because of good biocompatibility, high relaxivity, and easy and versatile functionalization, superparamagnetic iron oxide nanoparticles (IONPs) have been extensively studied and have emerged as promising $T_2$ CA. Benefiting from the size-dependent contrast, the ultrasmall IONPs also have demonstrated satisfactory $r_1$, making them alternative $T_1/T_2$ dual-contrast CA. Nevertheless, the increased $T_1$ relaxivity of the ultrasmall IONPs is the result of the consumption of the size, resulting in a relatively small $r_2$. Thus, ultrasmall IONPs may be not optimal as $T_1/T_2$ dual-contrast CA.

Divalent manganese ions (Mn$^{2+}$) possess five unpaired electrons and are of higher paramagnetism, and Mn$^{2+}$ doping has been proven to be an effective route to improve the $r_1$ of ultrasmall IONPs. Additionally, doped Mn$^{2+}$ with a higher magnetic moment ($\mu_{B}=5.92$) can occupy both the tetrahedral ($T_d$) and octahedral holes ($O_h$) in the crystal lattices, resulting in a special mixed spinel structure, a larger saturation magnetization (Ms), and a high $r_2$ of manganese-doped IONPs (Mn-IONPs). Moreover, based on the embedding rationale, the doped Mn$^{2+}$ and ultrasmall IONPs show synergetic enhancement, which will further improve both $r_1$ and $r_2$ of Mn-IONPs. Therefore, Mn-IONPs could be better candidates as dual-contrast CA.

Usually, the hydrophobic inorganic nanoparticles need to be modified with either natural macromolecules or synthetic polymers for hydrophilicity and further functionализation for biomedical applications. 1,2-Distearoyl-sn-glycero-3-phosphoethanolamine-N-[methoxy (polyethylene glycol)] (DSPE-PEG), with various advantages such as a low critical micellar concentration (CMC), excellent bio-compatibility, and long blood circulation time has been diffusely employed for surface modification. More importantly, it was reported that DSPE-PEG can significantly improve $r_2$ of the IONPs via the immobilization of nearby water molecules by forming hydrogen bonds.

Herein, Mn-IONPs were prepared through the dynamic simultaneous thermal decomposition (DSTD) method. Strikingly, after coating with DSPE-PEG, the obtained hydrophilic manganese-doped iron oxide nanocomposites (Mn-IONPs@PEG) showed both good $r_1$ and $r_2$ in vitro and in vivo, demonstrating that these nanocomposites comprising ultrasmall Mn-IONPs and DSPE-PEG could be ideal candidates for use as $T_1/T_2$ dual-contrast CA.

**Materials And Methods**

**Materials**

Oleyl alcohol and dibenzyl ether were purchased from Aldrich Chemical Co. (Saint Louis, MO, USA). Absolute ethyl alcohol was obtained from Chongqing Chuandong Chemical Co. Ltd. (Chongqing, China). Hexane and tetrahydrofuran (THF) were purchased from Adams Reagent Co. Ltd. (Shanghai, China). DSPE-PEG2000 was obtained from Pharmaion Co. Ltd. (Shanghai, China).
from Shanghai Ponsure Biotechnology Co. Ltd. (Shanghai, China). Dulbecco’s Modified Eagle’s Medium (DMEM) was purchased from Life Technologies Co. (Saint Louis, MO, USA). Fetal bovine serum (FBS) was obtained from Zhejiang Tianhang Biotechnology Co. Ltd. (Zhejiang, China). Penicillin/streptomycin solution and L-glutamine were purchased from Beyotime Biotechnology Co. Ltd. (Shanghai, China). Cell Count Kit-8 (CCK-8) was purchased from Boster Biological Technology Co. Ltd. (Wuhan, China). All of these reagents were used as received.

Synthesis Of Mn-IONPs

The Mn-IONPs were prepared following a previously reported procedure with modifications.12 The iron-erucate and manganese-oleate were first synthesized and described in the supporting information (SI) in detail. Next, 2 mmol of iron-erucate, 1 mmol of manganese-oleate and 6 mmol of oleyl alcohol were dissolved in 10 mL of dibenzyl ether at room temperature. The mixture was then heated to 100 °C and maintained for 30 min under a constant argon flow and magnetic stirring. Subsequently, the mixture was heated to 265 °C at a rate of 5 °C/min and was maintained for 1 h. After cooling to room temperature, the synthesized Mn-IONPs were washed with ethanol and centrifuged and finally dispersed in 20 mL of hexane for storage.

Surface Modification

By coating with the DSPE-PEG-2000, the Mn-IONPs were characterized with hydrophilicity. Following the previous method,32 50 mg of Mn-IONPs was first dispersed in 50 mL of THF to form 1 mg/mL of Mn-IONPs solution. Next, DSPE-PEG-2000 was mixed with 1 mL of Mn-IONPs solution with different mass ratios (NPs/PEG = 1:1, 1:5, 1:20, or 1:40) and then was added dropwise into 5 mL of deionized water under ultrasonication. After shaking overnight, the supernatants, including Mn-IONPs@PEG (1:1, 1:5, 1:20, or 1:40), were collected via filtration through a 0.22-μm syringe filter and was stored at 4 °C. Additionally, the hydrophilic Mn-IONPs-TMAH were obtained through ligand exchange with tetramethylammonium hydroxide aqueous solution (TMAH) and were described in the SI in detail.

Characterization

Transmission electron microscopy (TEM; Hitachi HT-7700, Japan) was used to observe the lattice spacing of the nanoparticles. Powder X-ray diffraction (XRD; PANalytical X’Pert Powder, Netherlands) was performed to detect the structure of the nanoparticles. Inductive-coupled plasma optical emission spectrometry (ICP-OES; Thermo Scientific Icap6300 Duo, USA) was carried out to determine the elemental composition of the nanoparticles. The hysteresis loops of the nanoparticles were recorded using a superconducting quantum interference device (SQUID; VSM-Versalab, USA) magnetometer.

Cytotoxicity

The human liver carcinoma cell line HepG2, purchased from the Chinese Academy of Sciences Cell Bank, was used for cytotoxicity analysis in this study. The cells were routinely cultured in DMEM containing 10% FBS, 2 mM L-glutamine, and 100 U/mL penicillin/streptomycin solution in a humidified incubator with 5% CO2 at 37 °C. CCK-8 was used for the analysis of the cytotoxicity of Mn-IONPs@PEG with serial metal (Fe + Mn) concentrations (0, 1, 5, 10, 20, or 50 μg/mL) were added to replace the previous medium, followed by incubation for an additional 24 h. Each concentration of the samples was repeated five times. Next, the medium was replaced with 100 μL of fresh medium including 10 μL of WST-8 solution. Following 1.5 h of coculture, a spectral scanning multimode reader (Thermo Scientific Varioskan Flash, USA) was used to determine the optical density at the wavelength of 450 nm.

MR Imaging In Vitro

In vitro MR imaging was carried out using a clinical 3.0 T MRI scanner (GE, Signa HDx 3.0T, USA) with a head coil. The Mn-IONPs@PEG and Mn-IONPs-TMAH (shown in SI) solutions with [Fe+Mn] concentrations from 0 to 0.5 mM were imaged using the following parameters: fast spin echo sequence (FSE) T1,W1: repetition time (TR) = 240 ms, echo time (TE) = 7.5 ms, field of view (FOV) = 160×160 mm², matrix = 320×192, slice thickness/spacing =2.0 mm/0.1 mm, number of excitations (NEX) = 4; T1,W1: mapping: TE = 9 ms, TR = 150, 300, 600, 900, 1200 ms, FOV = 160×160 mm², matrix = 256×256, slice thickness/spacing = 2.0 mm/0.1 mm, NEX = 1; FSE T1,W1: TR = 2000 ms, TE = 46.5 ms, FOV = 160×128 mm², matrix = 192×160, slice thickness/spacing
= 2.0 mm/0.1 mm, NEX = 4; T₂ mapping: TR = 1500 ms, TE = 7.8, 15.6, 23.5, 31.3, 39.1, 46.9, 54.8, 62.6 ms, FOV = 160×128 mm², matrix = 160×128, slice thickness/spacing = 2.0 mm/0.4 mm, and NEX = 1.

**MR Imaging In Vivo**

Animal experiments were conducted in accordance with the National Institutes of Health guidelines on the use of animals in research and were approved by the Laboratory Animal Welfare and Ethics Committee of the Army Medical University. Female C57BL/6 mice with a body weight of approximately 20 g at the age of 8 weeks were purchased from Chongqing TengXin Biotechnology Co. and were housed at the Experimental Animal Center of XinQiao Hospital. The mice were anesthetized with isoflurane and were maintained in a stereotaxic frame (RS10IP; RWD Life Science). The T₁WI (TE=9.3 ms, TR=580 ms, FOV = 60×60 mm², matrix = 288×192, slice thickness/spacing =2.0 mm/0 mm, and NEX = 4) and T₂WI (TE=69.6 ms, TR=2820 ms, FOV = 60×60 mm², matrix = 288×192, slice thickness/spacing =2.0 mm/0 mm, and NEX = 6.) Images of mice were acquired using a special rat coil (MS40; Suzhou Medcoil Healthcare Co., Ltd) and a 3.0-T clinical GE MRI scanner before and after injection of the Mn-IONPs@PEG (1:20) with 5 mg [Fe+Mn] per kilogram body weight.

**Results**

**Synthesis And Characterization Of Mn-IONPs**

Mn-IONPs were prepared via thermal decomposition of iron-eruciate and manganese-oleate complexes in the presence of oleyl alcohol in dibenzyl ether. Iron-eruciate and manganese-oleate complexes were manufactured through a simple method and are shown in Figure S1. The characteristic bands of iron-eruciate appeared at 1582, 1557, and 1433 cm⁻¹, while those of manganese-oleate were at 1606, 1549, and 1423 cm⁻¹, which were consistent with the data of a previous report. Coupled with the results of ICP-OES showing that the Mn/Fe molar ratio was approximately 0.20, Mn-IONPs were revealed to be successfully prepared. The hysteresis loop recorded by a SQUID magnetometer using a magnetic field up to 3.0 T at the temperature of 300 K showed that the saturation magnetization (Ms) of Mn-IONPs was 24.45 emu g⁻¹ (Figure 1F). Meanwhile, the negligible coercivity and remanence when the external magnetic field was absent indicate their superparamagnetic behavior. Furthermore, after coating with DSPE-PEG-2000 or TMAH, the obtained hydrophilic Mn-IONPs@PEG or Mn-IONPs-TMAH still maintained size uniformity and showed good stability in water.

**Relaxivity Of Mn-IONPs**

To assess T₁ and T₂ enhancement, the Mn-IONPs@PEG and Mn-IONPs-TMAH solutions were imaged using a 3.0 T clinical MRI scanner with FSE T₁WI, FSE T₂WI, T₁ mapping and T₂ mapping sequences. The two types of Mn-IONPs solutions
all showed concentration-dependent signal enhancement (Figure 2A–D). With the concentration increasing, the signal intensity tended to be brighter in T₁WI but gradually grew darker in T₂WI. The linear fitting of the concentration and 1/T₁ or 1/T₂ showed that r₁ values of Mn-IONPs-TMAH, Mn-IONPs@PEG (1:1), Mn-IONPs@PEG (1:5), Mn-IONPs@PEG (1:20), and Mn-IONPs@PEG (1:40) were 3.7, 2.4, 3.5, 7.1, and 7.8 mM⁻¹s⁻¹, respectively (Figure 2E), and the r₂ values were 36.9, 199.5, 149.7, 120.9, and 94.8 mM⁻¹s⁻¹, respectively (Figure 2F).

**In Vitro Cytotoxicity Assay**

Excellent biocompatibility was considered to be an important requirement of Mn-IONPs@PEG for in vivo

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**Figure 2**: In vitro MR images of the Mn-IONPs-TMAH and Mn-IONPs@PEG (1:1, 1:5, 1:20, and 1:40) at 3.0 T: (A) T₁-weighted, (B) T₁ mapping, (C) T₂-weighted and (D) T₂ mapping. (E, F) Linear fitting of 1/T₁ and 1/T₂ over different (Fe+Mn) concentrations of the nanocomposites.
application. The in vitro cell cytotoxicity assay of Mn-IONPs@PEG (1:20) was performed on HepG2 cells using CCK-8. Figure 3 shows the viability of HepG2 cells cocultured with the Mn-IONPs@PEG of [Fe+Mn] concentration ranging from 1 to 50 µg/mL at 37 °C for 12 hrs and 24 hrs. All the cells survived under the current experimental conditions. All Mn-IONPs@PEG samples showed insignificant toxicity with a cell viability greater than 80% at a concentration of less than 50 µg/mL for 12 hrs and 24 hrs. These results demonstrated that Mn-IONPs@PEG have a good biocompatibility.

In Vivo Contrast-Enhanced MRI

Based on the above results, we further explored the potential of the Mn-IONPs as dual-contrast CA in vivo. Due to the high relaxivity, Mn-IONPs@PEG (1:20) were employed for contrast-enhanced MRI of the liver. Figure 4A shows the axial T1WI and T2WI MR images acquired preinjection and postinjection (0, 10, 30, 60, and 120 min) of Mn-IONPs@PEG. The significantly brighter T1WI MR images and darker T2WI MR images suggest good r1 and r2 values of Mn-IONPs@PEG in vivo. The contrast-enhancement ratios of the MR signal intensity ΔSI (ΔSI = (SI\text{post} − SI\text{pre})/SI\text{pre} × 100%) are shown in Figure 4B and C. The ΔSI peak of 133% in T1WI appeared at 10 min after injection followed by a gradual decrease, unexpectedly becoming negative at 120 min. By contrast, the T2WI images showed obvious enhancement at 10 min after injection but increased to a plateau at 30 min and maintained for a long period.

Discussion

As is well known, T1 enhancement of superparamagnetic NPs depends on the interaction of the paramagnetic ions on the surface and nearby proton. Doping paramagnetic ions into the NPs has proven to be an effective route to improve r1.29 Many kinds of metal ions have been reported to be paramagnetic because of their unpaired electrons. Unlike lanthanide ions, the transition metal ions show both excellent biocompatibility and comparative paramagnetism and should be considered better ions for doping. Mn2+ with five unpaired electrons shows the highest paramagnetism compared with other transition metal ions.29 Additionally, based on the quantum mechanical inner-sphere theory, the T1 relaxation for NPs-based T1 CA can be given by

\[
\frac{1}{T_1} = \frac{qP_M}{T_{1M} + \tau_M} 
\]

where \(P_M\) is the mole fraction of metal ions, \(q\) is the number of water molecules bound per metal ion, \(\tau_M\) is the residence lifetime of the bound water, and \(T_{1M}\), which is related to \(\tau_M\) and the rotational correlation time \(\tau_R\), is the relaxation time of the bound water protons. It was revealed that the \(\tau_M\) of Mn (H2O)62+ is \(4.3 \times 10^4\) ns and is 2 orders of magnitude smaller than that of Fe(H2O)63+ (\(5.3 \times 10^6\) ns) and other transition metal ions.33 Thus, Mn2+ doping will effectively accelerate spin-lattice relaxation and increase \(r_1\). Moreover, embedding paramagnetic ions into the superparamagnetic NPs significantly boosts both \(r_1\) and \(r_2\) of superparamagnetic hybrid NPs by synergetic interaction between the T1 and T2 species.16

In addition to Mn2+ doping, decreasing the size of NPs into the ultrasmall region has also been proven to be an effective approach to boost \(r_1\). The smaller the size of
superparamagnetic NPs is, the larger the surface-to-volume ratio is, and the more superficial paramagnetic ions are, and the better \( r_1 \) the NPs possess.\(^\text{12}\) Thus, sufficiently small Mn\(^{2+}\) doping nanoparticles must be synthesized to guarantee good \( r_1 \). Theoretically, the key to successfully obtain ultrasmall Mn\(^{2+}\) doping nanoparticles is to decompose both the precursors at a similar temperature to form a multiple metal-containing monomer before nucleation to obtain smaller nanoparticles instead of a single-metal-containing monomer, resulting in the formation of a nondopant-containing nucleus initially and then another metal serving as the host to grow nanoparticles.\(^\text{12}\) Thus, the Mn-IONPs in this study were prepared by DSTD of the iron-erucate and manganese-oleate complexes, which had a similar decomposition temperature in the presence of oleyl alcohol in dibenzyl ether instead of iron (III) acetylacetonate and manganese (II) acetylacetonate. From the results of TEM, HRTEM, XRD and ICP, Mn-IONPs with an ultrasmall size of 3.0 ± 0.28 nm were successfully synthesized and the Mn\(^{2+}\) was fully proven to be doped successfully.

It is very important to have both good \( r_1 \) and \( r_2 \) to be good \( T_1/T_2 \) dual-contrast CA. However, we decreased the size of the NPs to the ultrasmall range to reach an optimal \( r_1 \) of the NPs in this study, resulting in a relative thicker spin-canted layer and a much lower saturation magnetization (\( M_s \)) and, finally, a much lower \( r_2 \) as expected. Unlike \( r_1 \), the relaxation of \( T_2 \) CA is dominated by the outer-sphere mechanism and can be given by\(^\text{34}\)

\[
\frac{1}{T_2} = \frac{256\pi^2\gamma_1M_s}{1215\rho} \left( \frac{1}{1+L/a} \right)^3 M_s^2 \tau_D \tag{2}
\]

where \( \gamma_1 \), \( M_s \), \( \rho \), \( a \) and \( L \) are the proton gyromagnetic ratio, molar mass, density, radius of NPs and impermeable thickness of the surface coating layer, respectively. \( \tau_D \) is the translational diffusion time, which is determined by \( r \) and \( D \) according to the equation \( \tau_D = r^2/D \), where \( r \) and \( D \) are the effective radius and diffusion coefficient, respectively. It is obvious that \( r_2 \) is proportional to \( a^3 \), while \( M_s^2 \) and \( \tau_D \) are inversely proportional to \( L^3 \). To increase \( M_s \), we doped Mn\(^{2+}\) into ultrasmall supermagnetic magnetite. Because the doped Mn\(^{2+}\) can occupy both the tetrahedral (\( T_0 \)) and octahedral holes (\( O_6 \)) in the crystal lattices, Mn-IONPs form a special mixed spinel structure, finally resulting in larger \( M_s \) and \( r_2 \); this finding has been confirmed in this study by showing that the \( r_2 \) of Mn-IONPs-TMAH (36.9 mM\(^{-1}\)s\(^{-1}\)) is larger than IONPs with the same size (\( r_2 = 29.2 \text{ mM}^{-1}\text{s}^{-1} \)).\(^\text{15}\) Using another approach by coating Mn-IONPs with DSPE-PEG, we further enhance its \( r_2 \).

According to Equation (2), \( r_2 \) can be improved by decreasing \( D \). The PEG molecules on the surface of the Mn-IONPs can immobilize surrounding water molecules in a large region by forming hydrogen bonds, resulting in a small \( D \) and a large \( \tau_D \).\(^\text{30,31}\) Thus, it is reasonable that all the DSPE-PEG-coated Mn-IONPs showed a much higher \( r_2 \) than THAM-coated Mn-IONPs. However, a decreasing trend for \( r_2 \) was noted with the increase in the molar ratio of Mn-IONPs: DSPE-PEG. This phenomenon can be easily explained because the hydrogen–bond interactions between PEG and water protons have a critical molar ratio. Too many coating molecules may exclude water protons from the induced magnetic field from the Mn-IONPs, forming an impermeable coating layer on the surface of the Mn-IONPs and leading to a lower \( r_2 \). Interestingly, we also found that \( r_1 \) can be changed by varying the coating molecules and the molar ratio of these molecules. Mn-IONPs-TMAH showed a higher \( r_1 \) than Mn-IONPs-PEG (1:1) and Mn-IONPs-PEG (1:5) but possessed a lower \( r_1 \) than Mn-IONPs-PEG (1:20) and Mn-IONPs-PEG (1:40). Unlike \( r_2 \), \( r_1 \) increased significantly with the increase in the molar ratio of Mn-IONPs: DSPE-PEG. This phenomenon can be attributed to \( \tau_R \) and \( q \) based on Equation (1). As is well known, \( T_1 \) relaxation relies on a chemical exchange between the paramagnetic center and water molecules in the first coordination sphere. The more water molecules located in the first coordination sphere, the better \( r_1 \) is. Although NPs become hydrophilic by coating with DSPE-PEG, there is still a DSPE-oleic acid layer that is impermeable on the surface of the NPs, resulting in much fewer water molecules in the first coordination sphere than Mn-IONPs-TMAH, in which the impermeable oleic acid layer has been exchanged. Thus, the \( r_1 \) of Mn-IONPs-TMAH was higher than that of Mn-IONPs-PEG (1:1) and Mn-IONPs-PEG (1:5). However, why do Mn-IONPs-PEG (1:20) and Mn-IONPs-PEG (1:40) show a higher \( r_1 \) than Mn-IONPs-TMAH? The cause may be that \( r_1 \) can also be remarkably influenced by \( \tau_R \), which is proportional to the molecular weight. With the increase in the molar ratio of Mn-IONPs: DSPE-PEG, the molecular weight of NPs increases notably, leading to a higher \( r_1 \). Over the critical ratio, the shielding effect of the impermeable DSPE-oleic acid layer will be counteracted and \( \tau_R \) will be dominant, making the \( r_1 \) of Mn-IONPs-PEG (1:20) and Mn-IONPs-PEG (1:40) higher than that of Mn-IONPs-TMAH.
Excellent biocompatibility is considered a key point for inorganic NPs application in vivo. To improve the biocompatibility, we selected DSPE-PEG, a well-established nonimmunogenic and nontoxic material, for Mn-IONPs coating. By CCK-8 analysis, the acquired Mn-IONPs@PEG (1:20) showed negligible cytotoxicity and excellent biocompatibility, which were consistent with the results of previous studies. Thus, we further explored the potential of Mn-IONPs@PEG (1:20) for T1/T2 dual-contrast MR imaging in vivo. Obviously, clear signal intensity changes were noted in both T1WI and T2WI over time. The signal intensity changes in the liver may be the result of the biological distribution and metabolism of Mn-IONPs@PEG in vivo. After injection, Mn-IONPs@PEG gradually accumulated in the liver within 10 mins, leading to increased ΔSI of both T1 and T2. However, the ΔSI of T1WI decreased over time because of passive aggregation driven by the phagocytosis of Kupffer cells, resulting in stable and individual Mn-IONPs@PEG gradually aggregating into clusters and causing the larger T2 contrast enhancement compared with the smaller T1 contrast enhancement. Particularly, the ΔSI even turned out to be negative at 120 min because the dominated T2 contrast enhancement significantly affected the T1 contrast. By contrast, the ΔSI of T2WI kept increasing until it reached a plateau and there was no downward trend within the observation period because of no degradation. The results obtained from the in vivo MRI verified that Mn-IONPs@PEG are highly sensitive T1 and T2 dual CA for imaging of the liver.

Conclusion

In summary, we successfully synthesized 3-nm-sized Mn-IONPs via facile thermal decomposition of iron-eruciate and manganese-oleate complexes in the presence of oleyl alcohol in dibenzyl ether. After encapsulation in DSPE-PEG-2000, the Mn-IONPs@PEG displayed good colloidal stability and excellent biocompatibility. More importantly, due to Mn2+ doping and DSPE-PEG-2000 coating, the synthesized NPs showed unexpected high r1 and r2. Confirmed by in vitro and in vivo MR imaging, the Mn-IONPs@PEG are ideal candidates for use as T1/T2 dual-contrast CA.

Acknowledgment

This research was supported by National Science Foundation of China (No. 81471635 and 81501521), Natural Science Foundation of Chongqing (No. cstc2018jcyjAX0321 and cstc2017jcyjBX0038).

Disclosure

The authors report no conflicts of interest in this work.

References

1. Smith BR, Gambhir SS. Nanomaterials for in vivo imaging. Chem Rev. 2017;117(3):901–986. doi:10.1021/acs.chemrev.6b00073
2. Ni D, Bu W, Ehlerding EB, et al. Engineering of inorganic nanoparticles as magnetic resonance imaging contrast agents. Chem Soc Rev. 2017;46(23):7438–7468. doi:10.1039/c7cs00316a
3. Weissleder R. Molecular imaging in cancer. Science. 2006;312:1168–1171.
4. Weissleder R. Molecular imaging: exploring the next frontier. Radiology. 1999;212(3):609–614.
5. Herschman HR. Molecular imaging: looking at problems, seeing solutions. Science. 2003;302(5645):605–608.
6. Louie AY, Hüber MM, Ahrens ET, et al. In vivo visualization of gene expression using magnetic resonance imaging. Nat Biotechnol. 2000;18:321.
7. Liu J, Wang L, Cao J, et al. Functional investigations on embryonic stem cells labeled with clinically translatable iron oxide nanoparticles. Nanoscale. 2014;6(15):9025–9033.
8. Willmann JK, van Bruggen N, Dinkelborg LM, Gambhir SS. Molecular imaging in drug development. Nat Rev Drug Discov. 2008;7(7):591–607.
9. Yang Z, Ren J, Ye Z, et al. Bio-inspired synthesis of PEGylated polypyrrole/polydopamine nanocomposite as theranostic agent for T1-weighted MR imaging guided photothermal therapy. J Mater Chem B. 2017;5:1108–1116.
10. Zhang F, Ni Q, Jacobson O, et al. Polymeric nanoparticles with a glutathione-sensitive heterodimeric multifunctional prodrug for in vivo drug monitoring and synergistic cancer therapy. Angew Chem Int Ed Engl. 2018;57(24):7066–7070. doi:10.1002/anie.201801984
11. Na HB, Song IC, Hyeon T. Inorganic nanoparticles for MRI contrast agents. Adv Mater. 2009;21(21):2133–2148. doi:10.1002/adma.2011033169
12. Zhang H, Li L, Liu XL, et al. Ultrasmall ferrite nanoparticles synthesized via dynamic simultaneous thermal decomposition for high-performance and multifunctional T1 magnetic resonance imaging contrast agent. ACS Nano. 2017;11(4):3614–3631. doi:10.1021/acsnano.6b07684
13. Jang JT, Nah H, Lee JH, et al. Critical enhancements of MRI contrast and hyperthermic effects by dopant-controlled magnetic nanoparticles. Angew Chem Int Ed Engl. 2009;48(7):1234–1238. doi:10.1002/anie.200805149
14. Shin T-H, Choi J-S, Yun S, et al. T1 and T2 dual-mode MRI contrast agent for enhancing accuracy by engineered nanomaterials. ACS Nano. 2014;8:3393–3401. doi:10.1021/nm4059777
15. Kim BH, Lee N, Kim H, et al. Large-scale synthesis of uniform and extremely small-sized iron oxide nanoparticles for high-resolution T1 magnetic resonance imaging contrast agents. J Am Chem Soc. 2011;133(32):12624–12631. doi:10.1021/ja203340u
16. Zhou Z, Huang D, Bao J, et al. A synergistically enhanced T1/T2 dual-modal contrast agent. Adv Mater. 2012;24(46):6223–6228. doi:10.1002/adma.201203169
17. Li J, You J, Wu C, et al. T1-T2 molecular magnetic resonance imaging of renal carcinoma cells based on nano-contrast agents. Int J Nanomedicine. 2018;13:4607–4625. doi:10.2147/IJN.S168660
18. Im GH, Kim SM, Lee DG, et al. Fe3O4/MnO hybrid nanocrystals as a dual contrast agent for both T1- and T2-weighted liver MRI. Biomaterials. 2013;34(8):2069–2076. doi:10.1016/j.biomaterials.2012.11.054
19. Tegafaw T, Xu W, Ahmad MW, et al. Dual-mode T1 and T2 magnetic resonance imaging contrast agent based on ultrasmall mixed gadolinium-dysprosium oxide nanoparticles: synthesis, characterization, and in vivo application. Nanotechnology. 2015;26(36):365102. doi:10.1088/0957-4484/26/36/365102
20. Yang M, Gao L, Liu K, et al. Characterization of Fe$_3$O$_4$/SiO$_2$/Gd$_2$O$_3$(CO$_3$)$_2$ core/shell/shell nanoparticles as T$_1$ and T$_2$ dual mode MRI contrast agent. *Talanta*. 2015;131:661–665. doi:10.1016/j.talanta.2014.08.042

21. Sun X, Du R, Zhang L, et al. A pH-responsive yolk-like nanoplatform for tumor targeted dual-mode magnetic resonance imaging and chemotherapy. *ACS Nano*. 2017;11(7):7049–7059. doi:10.1021/acsnano.7b02675

22. Cabrera-Garcia A, Checa-Chavarria E, Pacheco-Torres J, et al. Engineered contrast agents in a single structure for T$_1$-T$_2$ dual magnetic resonance imaging. *Nanoscale*. 2018;10(14):6349–6360. doi:10.1039/C7NR07948F

23. Gong M, Yang H, Zhang S, et al. Targeting T$_1$ and T$_2$ dual modality enhanced magnetic resonance imaging of tumor vascular endothelial cells based on peptides-conjugated manganese ferrite nanomicelles. *Int J Nanomedicine*. 2016;11:4051–4063.

24. Lee N, Hyeon T. Designed synthesis of uniformly sized iron oxide nanoparticles for efficient magnetic resonance imaging contrast agents. *Chem Soc Rev*. 2012;41(7):2575–2589.

25. Wang G, Zhang X, Skallberg A, et al. One-step synthesis of water-dispersible ultra-small Fe$_3$O$_4$ nanoparticles as contrast agents for T$_1$ and T$_2$ magnetic resonance imaging. *Nanoscale*. 2014;6(5):2953–2963.

26. Pellico J, Ruiz-Cabello J, Fernandez-Barahona I, et al. One-step fast synthesis of nanoparticles for MRI: coating chemistry as the key variable determining positive or negative contrast. *Langmuir*. 2017;33(39):10239–10247.

27. Li Z, Wang SX, Sun Q, et al. Ultrasmall manganese ferrite nanoparticles as positive contrast agent for magnetic resonance imaging. *Adv Healthc Mater*. 2013;2(7):958–964.

28. Zhang M, Cao Y, Wang L, et al. Manganese doped iron oxide theranostic nanoparticles for combined T$_1$ magnetic resonance imaging and photothermal therapy. *ACS Appl Mater Interfaces*. 2015;7(8):4650–4658.

29. Lee JH, Huh YM, Jun YW, et al. Artificially engineered magnetic nanoparticles for ultra-sensitive molecular imaging. *Nat Med*. 2007;13(1):95–99.

30. Tong S, Hou S, Zheng Z, et al. Coating optimization of superparamagnetic iron oxide nanoparticles for high T$_2$ relaxivity. *Nano Lett*. 2010;10(11):4607–4613.

31. LaConte LE, Nitin N, Zurkiya O, et al. Coating thickness of magnetic iron oxide nanoparticles affects R$_2$ relaxivity. *J Magn Reson Imaging*. 2007;26(6):1634–1641.

32. Lu J, Ma S, Sun J, et al. Manganese ferrite nanoparticle micellar nanocomposites as MRI contrast agent for liver imaging. *Biomaterials*. 2009;30(15):2919–2928.

33. RB LAUFFER. Paramagnetic metal complexes as water proton relaxation agents for NMR imaging theory and design. *Chem Rev*. 1987;87:901–927.

34. Zeng J, Jing L, Hou Y, et al. Anchoring group effects of surface ligands on magnetic properties of Fe$_3$O$_4$ nanoparticles: towards high performance MRI contrast agents. *Adv Mater*. 2014;26(17):2694–2698. 2016.

35. Johnson NJ, He S, Nguyen Huu VA, Almutairi A. Compact micellization: a strategy for ultrahigh $T_1$ magnetic resonance contrast with gadolinium-based nanocrystals. *ACS Nano*. 2016;10(9):8299–8307.

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