**Highly Optimized Iron Oxide Embedded Poly (Lactic Acid) Nanocomposites for Effective Magnetic Hyperthermia and Biosecurity**

Chiseon Ryu1
Hwangjae Lee1
Hohyeon Kim2
Seong Hwang1
Yaser Hadadian2,3
Ayeskanta Mohanty4
In-Kyu Park4
Beongki Cho1
Jungwon Yoon2,3
Jae Young Lee6

1School of Materials Science and Engineering, Gwangju Institute of Science and Technology, Gwangju, 61005, Republic of Korea; 2School of Integrated Technology, Gwangju Institute of Science and Technology, Gwangju, 61005, Republic of Korea; 3Research Center for Nanorobotics in Brain, Gwangju Institute of Science and Technology, Gwangju, 61005, Republic of Korea; 4Department of Biomedical Sciences, Chonnam National University Medical School, Hwasun-gun, Jeollanam-do, Republic of Korea

**Introduction:** Iron oxide magnetic nanoparticles (IONPs) have attracted considerable attention for various biomedical applications owing to their ease of synthesis, strong magnetic properties, and biocompatibility. In particular, IONPs can generate heat under an alternating magnetic field, the effects of which have been extensively studied for magnetic hyperthermia therapy. However, the development of IONPs with high heating efficiency, biocompatibility, and colloidal stability in physiological environments is still required for their safe and effective application in biomedical fields.

**Methods:** We synthesized magnetic IONP/polymer nanocomposites (MNCs) by embedding IONPs in a poly(L-lactic acid) (PLA) matrix via nanoemulsion. The IONP contents (Fe: 9–22 [w/w]%) in MNCs were varied to investigate their effects on the magnetic and hyperthermia performances based on their optimal interparticle interactions. Further, we explored the stability, cytocompatibility, biodistribution, and in vivo tissue compatibility of the MNCs.

**Results:** The MNCs showed enhanced heating efficiency with over two-fold increase compared to nonembedded bare IONPs. The relationship between the IONP content and heating performance in MNCs was nonmonotonous. The highest heating performance was obtained from MNC2, which contain 13% Fe (w/w), implying that interparticle interactions in MNCs can be optimized to achieve high heating performance. In addition, the MNCs exhibited good colloidal stability under physiological conditions and maintained their heating efficiency during 48 h of incubation in cell culture medium. Both in vitro and in vivo studies revealed excellent biocompatibility of the MNCs.

**Conclusion:** Our nanocomposites, comprising biocompatible IONPs and PLA, display improved heating efficiency, good colloidal stability, and cytocompatibility, and thus will be beneficial for diverse biomedical applications, including magnetic hyperthermia for cancer treatment.

**Keywords:** iron oxide nanoparticle, hyperthermia, inter-particle interactions, nanomedicine
magnetic and heat generation properties of IONPs are highly influenced by multiple factors, such as composition, size, shape, and interparticle interactions.\textsuperscript{14–16} These individual parameters have been experimentally and theoretically studied to improve the heat generation efficiency of IONPs. Therefore, small amounts of IONPs with low magnetic fields are favored to avoid their potential toxicity in various biomedical applications.\textsuperscript{17–19} Hence, extensive efforts have been made to synthesize IONPs with high heat generation efficiency.\textsuperscript{20–22} IONPs mainly generate heat in response to AMF via relaxation loss and/or hysteresis losses. Superparamagnetic single-domain magnetic nanoparticles at a low concentration in an AMF typically generate heat via Néel relaxation and Brownian relaxation.\textsuperscript{23,24} In addition, interparticle interactions between IONPs can play dominant roles in the overall magnetic and hyperthermia properties of interacting IONPs (eg, clusters and concentrated ferrofluids).\textsuperscript{25–27} The close proximity of multiple IONPs can affect the local dipole and anisotropic energy, resulting in an alteration of their magnetic susceptibility and heating efficiency.\textsuperscript{28–30} Hence, several attempts have been made to modulate the interparticle interactions of IONPs to enhance the magnetic or hyperthermia characteristics. Several studies have reported that interparticle interactions (dipole–dipole interactions) of IONPs can increase or decrease magnetic relaxation events and heat conversion, for which clear mechanisms have not been suggested.\textsuperscript{31–33} Nevertheless, it has been reported that optimization of interparticle interactions (such as spacing and density) can enhance the overall heating efficiencies of various magnetic nanoparticle clusters and ensembles.\textsuperscript{26,27,34} Evans et al investigated the effects of magnetic nanoparticle (magnetite) concentrations in silicone composites on heat generation.\textsuperscript{35} They found that heat generation was maximal at 1 [v/v]% magnetic nanoparticle concentration, and the optimal concentration was affected by the applied magnetic field. In addition, the colloidal stability of MNPs in biological environments is pivotal because MNPs readily aggregate in vivo and lead to substantial problems, such as reduction of heating performance, cytotoxicity, and thrombosis.\textsuperscript{36–38} Altogether, it is highly desirable to develop MNPs with high heat generation efficiency and colloidal stability in physiological environments for effective and biocompatible applications of MNPs.\textsuperscript{39,40} In this study, we produced IONPs/poly lactic acid (PLA) magnetic nanocomposites (MNCs) by incorporating different amounts of IONPs into the PLA matrix, in which the IONPs within the solid polymeric matrices were expected to have different interparticle distances (interactions) depending on their contents. Specifically, we synthesized hydrophobic IONPs (oleic acid-coated IONPs) and embedded them in PLA using an oil-in-water emulsion while varying the contents of IONPs and PLA (Scheme 1). We hypothesized that specific IONP content in the PLA matrix could
enhance efficient hyperthermia properties via optimized interparticle interactions. Furthermore, PLA-based IONP encapsulation (ie, MNCs) was expected to offer high stability and biocompatibility in biological environments. Notably, PLA is a biocompatible and biodegradable synthetic polymer that has been extensively used for various biomedical applications. Hence, given that the main components (ie, PLA and IONPs) are generally recognized as biocompatible, our MNCs could be expected to be biocompatible. Our IONP/polymer nanocomposite system may provide insights for improving the heating efficiency and stability of other magnetic nanoparticles by simply controlling the particle content in biocompatible polymer matrix.

**Materials and Methods**

**Synthesis of Oleic Acid-Coated Iron Oxide Nanoparticles**

Oleic acid-coated IONPs were synthesized according to a previously reported co-precipitation method. Briefly, iron (II) chloride tetrahydrate (FeCl$_2$·4H$_2$O) (4.3 g), and Iron (III) chloride hexahydrate (FeCl$_3$·6H$_2$O) (11.6 g) were completely dissolved in deionized water (DIW) (350 mL) in a three-neck round-bottom flask. The flask was then purged with nitrogen gas (N$_2$) and heated to 80 °C with vigorous stirring. Next, ammonium hydroxide solution (NH$_4$OH, ~25 wt%) (20 mL) was added, and the mixture was stirred for 5 min. Subsequently, oleic acid solution (3 mL) was added to the flask and then stirred for an additional 25 min. The mixture was cooled to room temperature, and a black precipitate settled at the bottom. The black precipitate was washed three times with a mixture of DIW and ethanol (1:1, v/v) to remove excess oleic acid and subsequently dried in an oven at 60 °C. FeCl$_2$·4H$_2$O, FeCl$_3$·6H$_2$O, NH$_4$OH, and oleic acid were purchased from Sigma-Aldrich, USA.

**Synthesis of Magnetic Nanocomposites**

The MNCs were prepared based on the modified methods described in the literature. Briefly, the oleic acid-coated IONPs (30, 60, 90, or 120 mg) were first suspended in chloroform (7.5 mL). Then, 180 mg of poly(L-lactide) (PLA; 260 kDa, Sigma-Aldrich) was added and dissolved in IONP-containing solution. Next, 36 mL of 8% (w/v) polyvinyl alcohol (PVA; 30–70 kDa, Sigma-Aldrich) was added and sonicated for 3 min to form an oil/water emulsion, which was then stirred for 24 h to evaporate the chloroform. The solution was subsequently centrifuged at 17,000 rpm for 30 min. The particles were collected and dispersed in DIW. The samples were washed three times by centrifugation, resuspended in DIW, and finally stored at 4 °C for further use. The individual MNCs synthesized with IONPs of 30, 60, 90, and 120 mg during emulsion were denoted as MNC1, MNC2, MNC3, and MNC4, respectively.

**Iron Content Quantification**

The iron content of each MNC was quantified using a colorimetric method. A colorimetric standard was first plotted using FeCl$_3$·6H$_2$O and FeCl$_2$·4H$_2$O (2:1, w/w). Briefly, various concentrations of Fe were prepared in

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**Scheme 1** Schematic for the synthesis of iron oxide nanoparticles (IONPs)-embedded poly(lactic acid) (PLA) magnetic nanocomposites (MNCs) via oil-in-water emulsion by differing the contents of IONPs in the MNCs.
hydrochloric acid (HCl, Duksan, Republic of Korea) solution, and 50 μL of each solution was transferred into a 96-well plate. Subsequently, 50 μL of 1% ammonium phosphate (AP, Sigma-Aldrich) and 100 μL of 0.1 M potassium thiocyanate (KSCN, Sigma-Aldrich) were added to the solution. After 5 min of reaction, the absorbance of the solution was measured at 490 nm using a microplate reader. The MNC sample was first lyophilized. Subsequently, the dehydrated sample was dissolved in HCl and reacted with the reagents using the same procedures as the standard samples. Finally, the iron content of each sample was calculated using the standard curve.

Materials Characterization
The morphologies and shapes of the synthesized nanomaterials were characterized using transmission electron microscopy (TEM, Technai G2S Twin) at 300 keV. Dynamic light scattering and zeta potential measurements were performed using a zeta-potential and particle-size analyzer (ELS Z2 Plus; Photal Otsuka Electronics). The crystal structure of the IONPs was analyzed by X-ray diffraction (XRD, Rigaku SmartLab). The magnetic properties of the particles were measured using a vibrating sample magnetometer (VSM, Lake Shore Cryotronics 7400 series) in a magnetic field ranging from −9 kOe to 9 kOe. From the hysteresis loop of each sample, a hysteresis area was obtained for comparison.

Hyperthermia Test
Magnetic heat generation was studied using IONPs, MNCs, and Resovist® (Meito Sangyo Co., Ltd. Nagoya, Japan). Solutions containing different concentrations (0.5, 1, 2, and 4 mg Fe/mL) were prepared for each sample. Before the hyperthermia experiment, the initial temperature of the sample solution was maintained at 25 °C using a water-circulating jacket. An AMF (OSH-120-B; Osung Hitech, Republic of Korea) with an amplitude (H, 3.25 kA/m) and frequency (f, 191 kHz) was applied for 10 min. In addition, to study the effects of the AMF conditions (ie, amplitude and frequency) on the heat generation of the prepared MNCs, a lab-made AMF (GIS) was employed. During AMF application, the temperature was monitored using an IR laser camera (FLIR SC300-series). The heat generation efficiency of the magnetic nanoparticles, which is typically referred to as specific loss power (SLP), was calculated using the following formula:

$$\text{Specific Loss Power (SLP)} = - \frac{c}{m_{Fe}} \times \frac{dT}{dt} \bigg|_{t=0}$$

where $C$ is the specific heat capacity of water, $m_{Fe}$ is the weight of iron, $dT$ is the temperature change, and $dt$ is the change in time.

Stability Test
For the stability tests, we performed both size analysis and hyperthermia analysis after incubating the samples in various solutions. First, MNC2 and Resovist® were incubated in PBS and phenol red-free cell culture medium (DMEM). The hydrodynamic diameters of individual samples were measured by DLS during incubation at 0 h, 6 h, 12 h, 24 h, and 48 h. SLP changes in the samples were measured after incubation in DMEM. MNC2 and Resovist® samples were diluted to 1 mg Fe/mL. The SLP value of each sample was obtained with AMF application for 10 min at 0 h and 48 h after incubation at 20 °C.

In vitro Cytocompatibility Test
The cytocompatibility of MNC2 and Resovist® was tested by in vitro NIH-3T3 (Korean Cell Line Bank) cell culture using a live/dead assay. The cells were seeded at a density of $2 \times 10^4$ cells/well in a 24-well plate and cultured in the Dulbecco’s modified Eagle’s medium (DMEM) for 12 h. DMEM was supplemented with 10% fetal bovine serum (FBS) and 1% antibiotic-antimycotic. Subsequently, the culture medium (1 mL) was replaced with the sample solution containing various concentrations of MNC2 or Resovist® in the culture medium and then incubated for an additional 24 h. Cells with no particles were used as a positive control. After 24 h, the sample solution was removed and washed with Dulbecco’s phosphate-buffered saline (DPBS). For live/dead staining of the cells, the sample was stained with a staining solution (DMI3000B; Leica, Germany). At least nine images from each sample were randomly selected for cytotoxicity analysis. Green- and red-stained cells were counted as live and dead cells, respectively. Cell viability was calculated as a percentage of the number of live cells from the total number of cells (live + dead cells). Experiments were conducted in triplicate (n = 3).
In vivo Biodistribution and Toxicity Study
All animal experiments were performed in accordance with the permission of the Institutional Animal Care and Use Committee (IACUC) of Chonnam National University, Republic of Korea (approval number: CNU IACUC-H-2020-25). Five-week-old male BALB/C nude mice (Orient Bio Inc., Korea) were anesthetized by vaporized isoflurane (Terrell; Piramal Critical Care Inc.) using a matrix vip 3000 vaporizer (Midmark Co.). For the biodistribution study, Cyanine 7 fluorescence dye (Cy 7) was loaded in MNC during O/W emulsion. In brief, 800 μg of the dye was added in oil phase while the other nano-emulsion procedure was the same as described above. The Cy7-loaded MNC was intravenously injected via tail vein with 6 mg/kg dose. In vivo fluorescence images of the animals were acquired at 0, 4, 8, 12, 24, and 48 h after injection using FOBI (Fluorescence optical bio-imaging). Main organs (heart, liver, spleen, lung, and kidney) were harvested and their fluorescence images were acquired to compare MNC biodistribution. For histological evaluation of potential organ toxicity of the MNC, the mice were sacrificed 24 h and 48 h after injection. The heart, liver, spleen, lung, and kidney were harvested and fixed in 4% formaldehyde. Untreated mice were also used as the control ( sham) for comparison. Samples were stained with hematoxylin and eosin (H&E) according to the literature. Individual images were acquired by optical microscope (DMI3000B; Leica, Germany).

Statistical Analysis
The results are indicated as the mean ± standard deviation (SD) unless otherwise noted. The differences among samples were compared by one-way analysis of variance (ANOVA) with Tukey’s post hoc comparison at a significance level of 0.05.

Results and Discussion
Synthesis of Magnetic Nanocomposites
To synthesize MNCs, oleic acid-capped hydrophobic IONPs were first synthesized by the co-precipitation method. TEM images showed that the prepared IONPs had sphere-like shapes with an average diameter of 9.3 ± 2.2 nm (Figure 1A). DLS measurements indicated that the average diameter of the particles was approximately 32.8 nm in chloroform (Figure 1B). Also, X-ray diffraction (XRD) pattern of the IONPs indicated the typical diffraction peaks for oleic-acid coated Fe₃O₄ nanoparticles (Figure S1). The synthesized IONPs showed superparamagnetic behavior and a saturation magnetization of 56 emu/gFe at 300 K (Figure 1C). Application of AMF to IONPs caused a dose-dependent temperature increase (Figure 1D). The SLP value of these oleic acid-capped IONPs was determined to be 37.2–41.6 W/gFe in the concentration range of 0.5 ~ 4 mgFe/mL in the media (1-octanol) (Figure S2).

MNCs containing different amounts of IONPs were prepared using oleic acid-coated IONPs and PLA in an oil/water single emulsion, in which the IONPs were expected to be firmly immobilized within the solid PLA polymer matrix (Scheme 1). The initial weight ratios of the IONPs and PLA were varied to 1:6, 2:6, 3:6, and 4:6, which resulted in the production of various MNCs with different local IONP densities and thus different interparticle interactions. MNC1, MNC2, MNC3, and MNC4 had Fe contents of 9, 13, 17, and 22 wt% (gFe per g nanocomposites), respectively (Table 1). The average diameter of all MNCs was approximately 160 nm when measured by TEM, and their hydrodynamic diameter was approximately 170 nm, as measured by DLS (Figure 2A and B). The zeta potential of MNCs was about −15 mV, which was similar to PLA NPs (Figure S3). Consequently, we successfully obtained various MNCs with different IONP contents but similar overall sizes and distributions. Hence, we could further study the effects of IONP density and interparticle interactions within the solid PLA matrix on the magnetic and hyperthermia properties. Notably, this synthetic process can be further useful for effectively embedding hydrophobic molecules together with IONPs in a polymer matrix in a reproducible manner (Figure 2C). The Nomenclatures, compositions, and dimension of various MNCs are listed in Table 1.

Magnetic Properties of Magnetic Nanomaterials
The field-dependent magnetization of the various samples was measured using a VSM (Figure 3). Both IONPs and MNCs exhibited superparamagnetic-like behavior (Figure 3A). The saturation magnetization (Mₛ) values of all the MNCs were higher than those of the nonencapsulated IONPs. The Mₛ values of the bare IONPs, MNC1, MNC2, MNC3, and MNC4 were 56.6, 72.8, 77.5, 75.8, and 65.1, emu/gFe at 300 K, respectively (Figure 3B). As the IONP content in the MNCs increased (to Fe 13%), Mₛ increased. Above this content, the Mₛ values of the MNCs
decreased. As results, MNC2 had the highest $M_s$ value. The magnetic behavior of IONPs and their complexes can be explained as a combination of the intrinsic magnetic properties of single IONPs (eg, susceptibility) and their interparticle interactions (dipole–dipole interactions). For IONPs with short particle distances, the magnetic dipolar interactions can affect the effective anisotropy and relaxation behaviors of individual particles. Such changes can enhance or reduce the strength of magnetization. For example, IONP clusters can show increased net local magnetization and enhanced magnetic relaxivity. The dipolar coupling between closely packed IONPs can alter the anisotropic energy of the complex. For our MNCs, the interparticle interactions (IONPs embedded in PLA) appear to positively contribute to the magnetization at moderate particle concentrations. Conversely, in MNCs containing large amounts of IONPs (MNC3 and MNC4), excessive interparticle interactions appeared to reduce the magnetic properties, likely due to the reduction of the global magnetic response of the IONPs. We compared the hysteresis loss areas, which are known to be significantly correlated with interparticle interactions and hyperthermia efficiency. MNC1 and MNC2 samples had two times larger hysteresis loss areas than the other.

Table 1 The Nomenclatures, Compositions, and Dimension of Various MNCs

| Sample Name | Input Weight Ratio (IONP:PLA) | Fe Content in MNC (wt%) | Size by TEM (nm) | Size by DLS (nm) | PDI |
|-------------|-------------------------------|-------------------------|------------------|------------------|-----|
| MNC 1       | 1:6                           | 9                       | 157.2            | 165.3            | 0.088|
| MNC 2       | 2:6                           | 13                      | 160.4            | 167.8            | 0.082|
| MNC 3       | 3:6                           | 17                      | 156.7            | 170              | 0.051|
| MNC 4       | 4:6                           | 22                      | 161.9            | 167              | 0.068|

Figure 1 Characterization of the synthesized oleic acid-coated IONPs. (A) A TEM image of the IONPs. (B) A size distribution of the IONPs measured by DLS. (C) Magnetization curve of the IONPs. (D) Temperature increases of the sample solutions containing various amounts of IONPs under the AMF ($H = 3.25$ kA/m and $f = 191$ kHz).
samples. The hysteresis loss areas decreased as the IONP content increased further in MNCs (Figure 3C).

**Magnetic Hyperthermia**

The hyperthermia properties of IONPs and MNCs were studied using an AMF \( (H = 3.25 \text{ kA/m} \text{ and } f = 191 \text{ kHz}) \). The product of the field parameters \( (H \times f = 0.62 \times 10^9 \text{ Am}^{-1} \text{s}^{-1}) \) was approximately 8 times less than the biological safety limit \( (H \times f = 5 \times 10^9 \text{ Am}^{-1} \text{s}^{-1}) \). The heating performance of hydrophobic IONPs and various MNCs was measured in 1-octanol at an identical IONP concentration \( (1 \text{ mg Fe/mL}) \) and compared in terms of SLP. The SLP values of all MNCs were approximately two times larger than those of the nonembedded IONPs (Figures 4A
and S4). The results suggest that the SLP could be improved by embedding IONPs in a PLA polymer matrix, in which individual IONPs were close to others. Among the MNCs, MNC2 had the highest SLP value, which was similar to the results of the saturation magnetization and hysteresis area (Figure 3). For both IONPs and MNC2, heat generation efficiencies were not substantially affected by their concentrations between 0.5 and 4 mgFe/mL (Figure 4B), which implies that no substantial agglomeration of either IONPs or MNC2 occurred during the AMF application and SLP measurement. The SLP of the MNCs was further compared with that of commercial IONPs (ie, Resovist®) in an aqueous medium (DIW) (Figure 4C). The SLP values of MNC1 and MNC2 were higher than those of Resovist®. Particularly, the SLP value of MNC2 was approximately 20% higher than that of Resovist® (Figure S5), implying the suitable performance of our MNCs for hyperthermia application.

The improved heat dissipation of MNCs is likely attributable to several possible factors. First, the high magnetic susceptibility (magnetization strength) of the MNCs might result in a high heating efficiency, based on the linear response theory. In the case of our MNC system, the IONPs are immobilized in the polymer matrix; hence, the Néel relaxation would play a major role in heat dissipation of the MNCs. In addition, as the IONP content increases, dipolar interactions would become substantial. Because the IONPs in the composites can behave collectively, such interparticle interactions may increase or decrease the anisotropy energy and heat dissipation depending on the nanoparticle composition, size, and homogeneity. Interparticle dipolar interactions generate a dipolar magnetic field and can increase the effective magnetic anisotropy, resulting in an increase in heat generation. Hysteresis loss can be a substantial heat generation mechanism for highly interactive magnetic nanomaterials. We observed that the hysteresis loss areas were higher for MNCs than for bare IONPs. In addition, at specific IONP contents (MNC1 and MNC2) such hysteresis loss was distinct (Figure 3D). It is also possible that the aggregation of individual IONPs in the PLA matrix resulted in a multidomain-like property and, thus, improved the hysteresis loss energy of the MNCs. However, excessive interparticle interactions might lead to decreases in magnetic susceptibility and hysteresis loss, resulting in a decrease in SLP. Dipolar interactions can cause magnetic nanoparticles stable against field orientation and increase the relaxation time. Several studies have reported the nonmonotonous heating performance of magnetic nanoparticles with different concentrations, indicating the optimal particle concentration for interparticle interactions and hyperthermic heat generation.

Effects of Magnetic Field Intensity and Frequency

We further investigated the effects of AMF conditions (amplitude (H) and frequency (f)) on heat generation by MNCs. Generally, the application of AMF with a higher H or f results in a higher SLP, and their influence on heat generation is different depending on the characteristics of the magnetic nanoparticles. We first measured the SLP of the samples by varying the amplitudes (H = 1.6 ~ 3.2 kA/m) with constant frequency (f = 191 kHz) (Figure 5A). All nanocomposites showed linear increases in SLP as the
amplitude increased. A two-fold increase in the amplitude resulted in increases in the SLP values by 1.8-, 2.1-, 2.0-, and 1.9-fold for MNC1, MNC2, MNC3, and MNC4, respectively. In addition, in the amplitude–SLP plots, MNC2 showed the highest slope among the samples. Accordingly, MNC2 was further analyzed to study the effects of different $H$ and $f$ ranges on SLP. Specifically, we applied a wide and high range of AMF amplitude (15–45 kA/m) using a lab-made AMF instrument (Figure 5B). MNC2 exhibited a linear relationship between SLP and an applied amplitude, with a drastic increase in SLP as the amplitude increased. For example, SLP values of MNC2 were 32.2, 81.1, and 204.7 (W/gFe) under AMF application of 15, 25, and 45 kA/m, respectively. In addition, MNC2 exhibited a linear relationship between the SLP and AMF frequency (Figure 5C). The SLP values were 23.6, 36.5, and 46.1 (W/gFe) with AMF of 160, 270, and 382 kHz, respectively. An increase in the frequency (2.4 times) resulted in a 2-fold increase in SLP. The results revealed that the influence of the amplitude on the heat generation of MNC2 was substantially greater than that of frequency. We could obtain the highest SLP value (204.7 W/gFe) from MNC2 with $H = 45$ kA/m and $f = 97$ kHz ($H \times f = 4.4 \times 10^9$ Am$^{-1}$s$^{-1}$), which is below the biosafety limit.

**In vitro Colloidal Stability**

The stability of magnetic nanoparticles in the physiological environment is essential for their robust and reliable biomedical applications because many magnetic nanoparticles, including IONPs, undergo severe agglomeration in practical applications, such as in vivo conditions. For stability tests, MNC2, which showed the highest heat generation efficiency among the samples, was selected and compared with Resovist®. After incubation in aqueous solutions, both samples appeared to remain stable without severe aggregation or color changes (Figure 6A). The hydrodynamic size of MNC2 remained unchanged after incubation in both PBS and DMEM. In contrast, the size of Resovist® gradually increased in DMEM, almost doubling after 48 h in DMEM (Figure 6B). In addition, the SLP value of MNC2 did not significantly change after 48 h incubation in DMEM, whereas the SLP value of Resovist® decreased by 14% (Figure 6C). These results indicate that MNC2 is stable in physiological environments. Such excellent stability of MNC2 may be attributed to the stable PLA matrix, which physically prevents direct interactions of the individual IONPs and their spontaneous aggregation. In addition, PLA presents a mild surface charge (zeta potential is -0.4 to -1 in physiological conditions), which offers stability in aqueous solutions. Conversely, the aggregation of Resovist® led to a change in interparticle interactions and performance deterioration.

**Cytocompatibility Tests**

The cytocompatibility of MNC2 and Resovist® was examined by in vitro culture of NIH-3T3 fibroblasts (Figure 7). The cell viability cultured with MNC2 remained high (≥96%)—up to 2 mgFe/mL concentration—corresponding to 15.4 mg MNC2 per mL medium, after 1 d in culture. In contrast, the viability of Resovist® gradually decreased with an increase in its concentration in the culture medium. For example, the cell viability in Resovist® sample was >90% up to 0.5 mgFe/mL; however, at a higher concentration, the cell viability significantly decreased.
For example, the cell viability of Resovist® at 2 mg Fe/mL was 77%. Resovist® is a carboxyldextran-coated IONP with hydroxyl groups on the surface. Different nanoparticles exhibit different cellular uptake, distribution, and reactivity (eg, reactive oxygen species formation) depending on their size, shape, and surface characteristics, which can affect toxicity. We speculate that the relative toxicity of Resovist® might be attributed to its large surface area and instability, resulting in particle aggregation. As stated earlier, the materials used in this study (ie, IONPs and PLA) are all biodegradable and biocompatible; hence, MNCs are believed to be minimally toxic (Figure S6). In addition, the excellent stability of MNCs may relieve a concern with the potential toxicity associated with particle instability (eg, aggregation).

**In vivo Biodistribution and Toxicity Study**

The Cy7-loaded MNC was intravenously injected via tail vein to evaluate biodistribution. In vivo biodistribution of MNC was detected at the time points of 0, 4, 8, 12, 24, and 48 h after injection. According to in vivo biodistribution images (Figure 8A), the injected MNC appeared to first circulate in the whole body through bloodstream and trap in specific organs of the body. In addition, we observed overall decreases in fluorescence signals 4 h after the injection (Figure 8A). As shown in Figure 8B, noticeable signals were observed from liver, spleen, and heart tissues post injection while minimal intensity was detected in the lung at both 24 and 48 h after injection. In general, the major fraction of the intravenously injected IONPs is first filtered by the liver and spleen, and reaches other organs. Based on the fluorescence intensity, the
relative amount of the MNC accumulated in the individual organs were compared. Fluorescence intensities of the most organs except spleen slightly decreased after 48 h of MNC injection, implying the MNC might slowly degrade or be excreted (Figure 8C). Mononuclear phagocytic systems (MPS), such as liver, might play roles in degrading IONPs in the blood. For example, Kupffer cells in the liver are known to actively degrade the IONPs.

After administration of the MNC, all the mice behaved normally over the entire observation period, indicating that the MNC did not induce apparent toxicity in vivo. Furthermore, several organs were harvested at 24 h and 48 h after the injection of control (sham) and MNC for histological analysis (Figure 9). H&E staining of the organs revealed that the heart, liver, spleen, lung and kidney showed no obvious difference between the untreated control and the MNC treated groups.

Figure 8 Biodistribution of MNC after intravenous injection. (A) In vivo fluorescence images of the mice acquired after 0, 4, 8, 12, 24, and 48 h of MNC injection. (B) Fluorescence images of the main organs after 24 h and 48 h of injection. (C) Fluorescence intensities of the individual organs harvested at 24 h and 48 h after injection of MNC.

Figure 9 Histological images of main organs (heart, kidney, liver, lung, and spleen) after 24 h and 48 h of MNC injection and the untreated control group. Scale bar = 100 μm.
with respect to the characteristic tissue structures. Therefore, MNC was determined to be biocompatible.

**Conclusion**

To produce stable and high-performance magnetic nanomaterials for potential hyperthermia applications, we synthesized magnetic nanocomposites by embedding IONPs in PLA while controlling the IONP content in the composites. The fabricated MNCs showed enhanced magnetization and heating efficiency compared with nonembedded IONPs. Notably, MNC2 containing moderate amounts of IONPs (13 wt% of Fe) showed higher magnetization and heating efficiency (ie, SLP) than the other nanocomposites. Further, these maximized characteristics appear to originate from the favorable interparticle interactions. In addition, MNC2 exhibited excellent colloidal stability in aqueous solutions, as evidenced by negligible changes in hydrodynamic sizes and SLP after the incubation, which led to consistent heating in physiological environments against the clinically approved IONPs, Resovist®. In vitro cell cytotoxicity tests revealed that MNC2 had good cytocompatibility—superior to Resovist®. In vivo study indicated that MNC had great biocompatibility. Altogether, we successfully produce MNCs displaying improved magnetic properties, heating performance, stability, and biocompatibility. These unique properties will make our MNCs have great potential for various hyperthermia applications. In addition, our MNCs were prepared using widely used biocompatible materials (ie, PLA and IONPs) for their potential translation to clinical applications. Nevertheless, future studies, such as in vivo applications for cancer treatment, are required to clearly demonstrate their efficacy and biocompatibility.

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**Disclosure**

Professor Jae Young Lee reports a patent KR2021-0016924 pending. The authors report no other conflicts of interest in this work.

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