Highly sensitive detection of protein biomarkers via nuclear magnetic resonance biosensor with magnetically engineered nano ferrite particles

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Abstract: Magnetic-based biosensors are attractive for on-site detection of biomarkers due to the low magnetic susceptibility of biological samples. Here, we report a highly sensitive magnetic-based biosensing system that is composed of a miniaturized nuclear magnetic resonance (NMR) device and magnetically engineered nanoferrite particles (NFPs). The sensing performance, also identified as the transverse relaxation ($R_2$) rate, of the NMR device is directly related to the magnetic properties of the NFPs. Therefore, we developed magnetically engineered NFPs (MnMg-NFP) and used them as NMR agents to exhibit a significantly improved $R_2$ rate. The magnetization of the MnMg-NFPs was increased by controlling the Mn and Mg cation concentration and distribution during the synthesis process. This modification of the Mn and Mg cation directly contributed to improving the $R_2$ rate. The miniaturized NMR system, combined with the magnetically engineered MnMg-NFPs, successfully detected a small amount of infectious influenza A H1N1 nucleoprotein with high sensitivity and stability.

Keywords: biosensor, NMR, nanoferrite particles, transverse relaxation, magnetization, on-site detection

Introduction

On-site detection for rapid initial examination regarding possible virus infection is critical in preventing serious contagions. Accordingly, there has been an increase in the development of sensitive biosensor systems that include electrical, optical, and magnetic-based mechanisms.1–6 In particular, the magnetic-based sensing system is an attractive method for biosensor applications. This is possible because intrinsically biological media have low magnetic susceptibility. Consequently, the magnetic sandwich assay can achieve a high detection signal with low interferences even in a complex biological background.7

Nuclear magnetic resonance (NMR) is a useful magnetic-based sensing technology that enables rapid, stable, and highly sensitive biomarker detection by utilizing magnetic resonance technology to detect biomarkers labeled with magnetic particles.8,9 The transverse relaxation ($R_2$) rate of the water protons in the vicinity of the magnetic particles and the target biomarker is modified upon reacting with the magnetic particles, and this change can be detected by the NMR. The biomarkers labeled with magnetic particles exhibit a faster decay rate (larger $R_2$) of NMR signals than that of non-magnetic particle-labeled biomarkers.10–12 The NMR-based detection technology requires minimal sample purification steps and consequently reduces sample loss. In addition, because the NMR signal is generated from the whole volume, the binding kinetics for the NMR signal
is much quicker than that of surface reaction-based sensors. This allows the NMR assay the advantage of obtaining a rapid sensing signal at a faster pace. Naturally, the NMR signal is directly relevant to the magnetic particles. Because magnetic particles with a higher \( R_2 \) rate can improve the detection sensitivity in NMR-based sensing, developing a magnetic NMR agent that exhibits a higher \( R_1 \) rate is imperative in achieving maximal sensitivity in the NMR sensing.

In this study, we developed a high-performance magnetic particle to improve the NMR sensitivity and then demonstrated its feasibility as an NMR agent by conducting a detection test for the nucleoproteins of the influenza A H1N1 virus. In order to achieve this goal, we magnetically engineered Fe-nanoferrite particles (Fe-NFPs) to enhance the \( R_2 \) rate. The Fe ions in tetrahedral A (T\(_4\)) sites or octahedral B (O\(_6\)) sites were substituted with Mn ions (Mn-NFPs) and Mn/Mg ions (MnMg-NFPs) for increased magnetization; the \( R_2 \) rate of the NFPs is proportional to the magnetization (\( R_2 = \frac{d^3 M^2}{d} \); \( d \) represents the diameter of the nanoparticle).\(^{13}\)

### Materials and methods

#### Synthesis of nanoferrites and surface modification

All the NFPs were synthesized using a high temperature thermal decomposition method.\(^{14,15}\) The metal precursors (Fe (III) acetylacetonate \([\text{C}_9\text{H}_8\text{O}_2\text{Fe}]\) (>99.9%), Mn (II) acetate tetrahydrate \([\text{C}_6\text{H}_5\text{COO}]_2\text{MnH}_{12}\text{O}_4\) (99.99%) and Mg acetate tetrahydrate \([\text{C}_6\text{H}_5\text{O}_2\text{MgH}_{12}\text{O}\) (99.999%)) and other materials such as the surfactants (oleic acid \([\text{C}_{18}\text{H}_{34}\text{O}]\) (90%), oleylamine \([\text{C}_{16}\text{H}_{33}\text{N}]\) (70%), reducing agents (1,2-hexadecanediol \([\text{C}_4\text{H}_{18}\text{O}]\) (90%)), and solvent (benzyl ether \([\text{C}_{14}\text{H}_{18}\text{O}]\) (99%)) were mixed and heated at 296°C.\(^{11}\) The Mn and Mg cation concentration and distribution in Mn-NFPs and MnMg-NFPs were chemically controlled by adjusting the amount of the Mn and Mg precursors and the amount of the reducing agent during the synthesis process. Then, all the synthesized NFPs were rendered water soluble by coating the surface of particles with polyethylene glycol (PEG).

Water-soluble NFPs were obtained by forming a lipid layer (PEGylated lipid) with 1-myristoyl-2-hydroxy-sn-glycero-3-phosphocholine/1,2-dipalmitoyl-sn-glycero-3-phosphoethanolamine-N-(lauroyl)-polyethylene glycol 2000 (MHPC/DSPE-PEG2K). The NFPs were conjugated with influenza A H1N1 nucleoprotein antibodies. The reaction of the primary amines on the antibody with PEG-coated NFPs (NFPs@PEG) was catalyzed by 1-ethyl-3-[3-dimethylaminopropyl]carbodiimide hydrochloride and sulfo-N-hydroxysuccinimide. More detailed protocols regarding the surface modification of NFPs and antibody conjugation with NFPs are indicated in our previous reports.\(^{16-18}\)

### Characterization of nanoferrites

The size and size distribution of the synthesized NFPs were measured by transmission electron microscopy. The hydrodynamic diameter (\(d_h\)) and polydispersity index (PDI) of the NFPs@PEG were measured using a dynamic light scattering system. The magnetic hysteresis characteristics, including the major and minor hysteresis loops for both the non-coated (powder state) and PEG-coated (fluid state) NFPs, were investigated using a vibrating sample magnetometer. The \( R_2 \) rate was measured by employing a miniaturized NMR system.\(^{19}\) The NMR system consists of three components, microcoils, on-board NMR electronics, and a small permanent magnet. The permanent magnet can be used to generate polarizing magnetic field, \(B_0\)=0.1–0.5 T. The concentrations of NFPs for the measurement of \( R_2 \) rate were varied from \(10^{-2}\) to \(10^{-11}\) g/mL.

### Relaxation measurement

The biosensing performances, including sensitivity and limit-of-detection (LOD) of the NFPs, were evaluated by detecting the nucleoproteins of influenza A H1N1 virus using the miniaturized NMR system combined with the magnetic sandwich method. The H1N1 nucleoprotein and antibody to the H1N1 nucleoproteins were purchased from Sino Biological Inc. A total of 15 µL of the H1N1 nucleoprotein mixture solution was first mixed with polystyrene beads (diameter: 3 µm) conjugated with the capture antibody and incubated for 30 min at room temperature. Uncaptured H1N1 nucleoproteins were removed by washing the beads with the wash buffer. The NFPs conjugated with detection antibody were then added, and the mixture was incubated again for 20 min. The unbound NFPs were removed by washing and filtering the mixture with the wash buffer, and finally, 15 µL of phosphate-buffered saline (PBS) was added to the mixture for NMR measurement (Scheme 1). We carried out all \( R_2 \) measurements at the polarizing magnetic field of \(B_0\)=0.5 T. All the measurements were done in triplicate at room temperature.

### Results and discussion

As the coating and dispersion status of the particles inside the fluid affects the magnetic properties, we investigated the PEG
MnMg-NFPs showed a higher magnetization than that of the Fe-NFPs. Considering that the preferential site of Mn$^{2+}$ cations is the O$_A$ site of Fe-NFPs, the improved magnetization of Mn-NFPs is physically thought to be due to the possible substitution of Fe$^{3+}$ cations with 4 $\mu_B$ (Bohr magneton) in the O$_A$ sites by Mn$^{2+}$ cations with 5 $\mu_B$ during the synthesis process (Figure 2C).

In the case of the MnMg-NFPs, where the preference site of Mg$^{2+}$ cations is the T$_A$ site, the Mg$^{2+}$ cations ($\mu_B = 0$) replace the Fe$^{3+}$ cations in the T$_A$ site of Mn-NFPs, and as a result, the magnetic moment of the MnMg-NFPs increases (6.2 $\mu_B$, total net $\mu_B = 6 \mu_B [O_A \; \text{site}] - 5 \mu_B [T_A \; \text{site}]$). The fluid state NFPs@PEG (Figure 2B) also retains the superparamagnetic phase and stable DC magnetic hysteresis loops. This is due to the superior PEG coating status, which allows for a more uniform anti-body conjugation to the NFPs@PEG without spontaneous magnetic aggregation.

The $R_2$ characteristic of each NFPs@PEG was measured using the miniaturized NMR system in a magnetic field $B_0$ (0.5 T) to evaluate the feasibility as an NMR agent for the detection of H1N1 nucleoproteins. In order to measure the $R_2$ rate, Fe-NFPs@PEG, Mn-NFPs@PEG, and MnMg-NFPs@PEG were dispersed in DI water. The concentration of the NFPs@PEG dispersed in DI water varied from 5×10$^{-2}$ g/mL to 5×10$^{-10}$ g/mL. Figure 3 shows the $R_2$ values for the three NFPs@PEGs at various concentrations. The observed $R_2$ values increased as the concentration increased, and the MnMg-NFPs@PEG exhibited the highest $R_2$ value among the different NFPs@PEGs.

Considering the fact that the $R_2$ is proportional to magnetization ($R_2 = 3 \pi^2 M^2$), the highest $R_2$ value of the MnMg-NFPs@PEG is thought to be due to the highly improved magnetization caused by engineering the cation concentration and distribution in the NFPs. This result demonstrates that the MnMg-NFPs@PEG is high enough to be considered as an NMR agent.

In order to demonstrate the applicability of the engineered NFPs as an NMR agent, we used the MnMg-NFPs@PEG and the Fe-NFPs@PEG to detect H1N1 nucleoproteins in a miniaturized NMR system. Figure 4A shows the miniaturized NMR system and representative NMR signal (T2 time of MnMg-NFPs@PEG). Figure 4B exhibits the H1N1 nucleoprotein detection ability ($\Delta R_2 = R_2, \text{P-beads + H1N1 + NFPs} - R_2, \text{P-beads}$) of the MnMg-NFPs@PEG and Fe-NFPs@PEG. Both NFPs@PEGs successfully detected the amount of H1N1 nucleoprotein with
Figure 1 (A) Sizes of synthesized NFPs measured by TEM, (B) size distribution, and (C) hydrodynamic diameter and PDI of the PEG-coated NFPs characterized by a DLS system.

Abbreviations: NFP, nanoferrite particle; TEM, transmission electron microscopy; PDI, polydispersity index; PEG, polyethylene glycol; DLS, dynamic light scattering.
a low error range of <4.5%. The H1N1 nucleoproteins bound to a polystyrene bead can be a cause of the aggregation of NFPs and the corresponding increase in the $R_2$ rate.

As can be clearly seen in the result, the MnMg-NFPs@PEG showed a significantly higher $\Delta R_2$, a wider dynamic range, and a lower LOD than those of Fe-NFPs@PEG. The MnMg-NFPs@PEG showed a $\Delta R_2$ of 0.501 s$^{-1}$ (g/mL)$^{-1}$ with an LOD of $1\times10^{-9}$ g/mL, while the Fe-NFPs exhibited a $\Delta R_2$ of 0.329 s$^{-1}$ (g/mL)$^{-1}$ with an LOD of $1\times10^{-7}$ g/mL. All the results strongly demonstrate that the magnetically engineered MnMg-NFPs have the potential to be a promising NMR agent for the detection of viral biomarkers with high sensitivity and stability.

**Conclusion**

We developed the magnetically engineered MnMg-NFPs for application as an NMR agent to detect infectious viruses. The Mn and Mg cation concentration and distribution in the MnMg-NFPs were chemically controlled during the Fe-NFP synthesis to improve the magnetization of spinel-structured NFPs. As a result, the MnMg-NFPs showed the highest magnetization value among the prepared NFPs, as well as a correspondingly improved $R_2$ rate. The potential of the MnMg-NFPs as an NMR agent was demonstrated by conducting the H1N1 nucleoprotein detection test. The MnMg-NFPs successfully detected a tiny amount of H1N1 nucleoproteins in PBS with a stable and high $R_2$ rate (high sensitivity).
**Figure 4** (A) Miniaturized NMR system and representative measured NMR signal (T2 time). (B) Detection results of influenza A H1N1 nucleoprotein using the miniaturized NMR system with MnMg-NFPs@PEG and Fe-NFPs@PEG. The MnMg-NFPs@PEG and Fe-NFPs@PEG are conjugated with the detection antibodies and used as an NMR agent. The polystyrene bead is conjugated with the capture antibodies.

**Abbreviations:** NMR, nuclear magnetic resonance; NFP, nanoferrite particle; Peg, polyethylene glycol.

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**Disclosure**

The authors report no conflicts of interest in this work.

**References**


