Octahedron Iron Oxide Nanocrystals Prohibited Clostridium difficile Spore Germination and Attenuated Local and Systemic Inflammation

Wei-Ting Lee1, Ya-Na Wu2, Yi-Hsuan Chen3, Shang-Rung Wu2, Tsai-Miao Shih2, Tsung-Ju Li1, Li-Xing Yang1, Chen-Sheng Yeh4, Pei-Jane Tsai1,3,5 & Dar-Bin Shieh1,2,6,7

Clinical management of Clostridium difficile infection is still far from satisfactory as bacterial spores are resistant to many chemical agents and physical treatments. Certain types of nanoparticles have been demonstrated to exhibit anti-microbial efficacy even in multi-drug resistance bacteria. However, most of these studies failed to show biocompatibility to the mammalian host cells and no study has revealed in vivo efficacy in C. difficile infection animal models. The spores treated with 500 µg/mL Fe3δO4 nanoparticles for 20 minutes, 64% of the spores were inhibited from transforming into vegetative cells, which was close to the results of the sodium hypochlorite-treated positive control. By cryo-electron micro-tomography, we demonstrated that Fe3δO4 nanoparticles bind on spore surfaces and reduce the dipicolinic acid (DPA) released by the spores. In a C. difficile infection animal model, the inflammatory level triple decreased in mice with colonic C. difficile spores treated with Fe3δO4 nanoparticles. Histopathological analysis showed a decreased intense neutrophil accumulation in the colon tissue of the Fe3δO4 nanoparticle-treated mice. Fe3δO4 nanoparticles, which had no influence on gut microbiota and apparent side effects in vivo, were efficacious inhibitors of C. difficile spore germination by attacking its surface and might become clinically feasible for prophylaxis and therapy.

Nanomaterials have attracted significant interest in medicine. Certain microorganism-reactive nanomaterials have been used as alternative bactericides1, namely, silver, zinc oxide, and titanium oxide nanoparticles, all of which have remarkable antibacterial properties2,3. The antibacterial mechanisms of nanoparticles may be attributable to their generation of reactive oxygen species, disruption of cell membranes, ability to bind thiol groups, and their release of toxic ions4. Spore-formation enables bacteria to survive nutritional deprivation and harsh environments. They can resist ultraviolet radiation, desiccation, high temperatures, extreme freezing, and chemical disinfectants5. Spores can reactivate themselves to the vegetative state when the environment becomes favorable. Therefore, Clostridium species, spore-forming pathogens, usually challenge clinical disease management and prevention. Clostridium difficile, a pathogen associated with healthcare-facility-related (nosocomial) infections, is a major cause of antibiotic-treatment-related diarrhea, pseudomembranous colitis, abdominal pain, fever, and death6. The normal flora in the gut can inhibit the growth of C. difficile and therefore protect patients from...
developing *C. difficile* infection. *C. difficile* infection usually occurs in patients on a long-term regimen of antibiotics, and it is often initiated by the spores acquired from healthcare workers. Once a patient develops *C. difficile* infection, there are only a few antibiotics available to control it. Moreover, the failure rate of first-line antibiotics and the *C. difficile* infection relapse rate are both dramatically high. Consequently, about two decades ago, the attributable post-diagnosis mortality rate was 6.9% at 30 days and 16.7% at 1 year. The spores of *C. difficile* are the major cause of *C. difficile* infection. Compared with oxygen-sensitive vegetative bacteria, *C. difficile* spores survive for up to several months in room air and in low-pH gastric contents. As the spores enter the human digestive tract, they germinate after they have been exposed to bile salts and their derivatives, and then they are colonized in the colon. The virulence of *C. difficile* depends upon the gene expression of tcdA-encoded toxin A, an enterotoxin, and tcdB-encoded toxin B, a cytotoxin. Both cause intestinal inflammation and neutrophil infiltration in the infected foci.

The incidence of *C. difficile* infection has significantly increased in the past 15 years. *C. difficile* infection has become a major cause of nosocomial-associated infection in the world. Antibiotic-resistant *C. difficile* is not only potentially fatal, but it also causes healthcare-associated economic burdens. The available present antibiotics are targeted to vegetative bacterium, however, the infective form is the spore. Current *C. difficile* infection clinical management is still far from satisfactory because the spores are resistant to many chemical agents and physical treatments, which makes effective management of the spores an important problem. Therefore, anti-germination approach could lead to the prevention of *C. difficile* infection. Some newly designed cholate derivatives show promise against *C. difficile* infection; however, they are still under pre-clinical study. Sodium hypochlorite, a standard disinfectant, has outstanding antimicrobial activity but undesirable side effects: it is corrosive and irritates tissue. To control spore germination and *C. difficile* infection, it is important to develop an efficacious and biocompatible spore-control strategy.

There are various well-known antibacterial nanomaterials, e.g., silver (Ag) and zinc oxide (ZnO) nanoparticles, and zero-valent iron nanoparticles, which are prominently bactericidal against *Escherichia coli*. The nanotechnology even has been reported that it could overcome the problem of multi-drug resistant bacteria. However, most current antibacterial nanomaterials are primarily multifunctional generic biocidal agents against vegetative cells; their sporicidal activity at high concentrations has been explored in only a few studies. Many studies demonstrated the excellent antibacterial ability of nanoparticles, but their impacts to the gut microbiota were not clear. The healthy gut microbiota prevent the host from colonic pathogen infection. Therefore, an ideal antibacterial nanomaterial should exhibit the specific pathogen targeting capability without causing disruption of healthy gut microbiota.

The toxicity of various bactericidal nanoparticles to mammalian cells has become an increasing concern. Iron-containing particles are generally recognized as highly biocompatible nanomaterials. A few specific iron oxide nanoparticles have recently been reported to be antibacterial; they were synthesized and showed therapeutic potential for cancer therapy. In this study, we evaluated whether such single-crystal nonstoichiometric Fe₃O₄ magnetite nanoparticles could inhibit *C. difficile* spore germination in vivo. The potential mechanisms were explored from both material science and molecular biology perspectives.

**Results**

**Testing the sporicidal activity of Fe₃O₄ nanoparticles against *C. difficile***. We used a previously described method to explore the inhibitory efficacy of Fe₃O₄ nanoparticles to spores. After they had been incubated with sodium hypochlorite for 20 minutes, inactivated spores were used as positive controls, as described elsewhere. After they had been incubated with nanomaterials of various concentrations (5, 50, and 500 µg/mL) for 20 minutes, the treated spores were then stimulated using 10 mM taurocholate to induce germination.

Spores treated with 50 and 500 µg/mL of Fe₃O₄ nanoparticles showed significantly (***P < 0.001) inhibited germination (Fig. 1), but spores treated with 5 µg/mL did not. There were no significant differences between the sodium hypochlorite-treated positive controls and the Fe₃O₄ nanoparticle-treated
Intriguingly, there was no significant difference in germination inhibition between spores treated with 14-nm and 22-nm Fe$_3$O$_4$ nanoparticles (Fig. 2A). In Supplementary Table 1, the germination kinetic results show that the Michaelis-Menten constant ($K_m$) for taurocholate-treated *C. difficile* CCUG 37780 spores increased from 4.34 to 8.43 M in mice treated with 50 μg/mL of Fe$_3$O$_4$ nanoparticles. The kinetic analysis suggested that Fe$_3$O$_4$ nanoparticles have an inhibiting constant ($K_i$) of approximately 62 μg/mL. To confirm the sporicidal property of 500 μg/mL of Fe$_3$O$_4$, the number of Fe$_3$O$_4$- and 3%-sodium hypochlorite-treated colony-forming unit (CFU) spores was counted. Compared with the negative control set, the number of CFUs of Fe$_3$O$_4$-treated and 3%-sodium hypochlorite-treated spores was ~60% lower (Fig. 2B). Furthermore, the inhibition rates of CFUs in Fe$_3$O$_4$-treated and in 3%-sodium hypochlorite-treated spores were not significantly different.

Phase-contrast microscopy showed that in the positive controls, to which taurocholate was added for 15 minutes, but not in the negative controls, the contrast of the spores transitioned from bright to dark (Fig. 2C). The contrasts of the spores treated with 500 μg/mL Fe$_3$O$_4$ nanoparticles were not significantly different. The luminescence signal that accompanies DPA release was significantly higher during taurocholate-induced spore germination (Fig. 2D). However, the luminescence signal was less intense and less abundant in the Fe$_3$O$_4$-treated group than in the taurocholate-treated group. No luminescence signal was detected in the group of spores not treated with taurocholate because the spores did not release DPA.

Inhibitory effect of Fe$_3$O$_4$ nanoparticles on virulent strains of *C. difficile*. *C. difficile* strain CCUG 37780 does not have the *tcdA* and *tcdB* genes. To determine what doses of Fe$_3$O$_4$ nanoparticles affect the spore germination of *C. difficile*, CCUG 19126 and BAA-1805 *tcdA*- and *tcdB*-positive strains were treated for 20 minutes with 5, 50, and 500 μg/mL doses of Fe$_3$O$_4$ nanoparticles, and then treated with 10 mM taurocholate. The germination of *C. difficile* CCUG 19126 spores and of *C. difficile* BAA-1805 spores was not significantly different.
between those treated with 500 μg/ml of Fe$_{3}$O$_{4}$ nanoparticles and those treated with sodium hypochlorite, but CCUG 19126 was relevantly resistant to treatment with 50 μg/mL of Fe$_{3}$O$_{4}$ nanoparticles (Fig. 3A).

To evaluate the bactericidal effect of Fe$_{3}$O$_{4}$ nanoparticles on vegetative cells in different strains, three groups of vegetative cells—CCUG 37780, CCUG 19126, and BAA-1805—were first treated for 20 minutes with various concentrations of Fe$_{3}$O$_{4}$ nanoparticles and then with 10 mM taurocholate to induce spore germination. Spore germination was effectively inhibited in the two Fe$_{3}$O$_{4}$-treated pathogenic strains, as it was in the nonpathogenic CCUG 37780 strain. (**P < 0.001, *P < 0.05; one-way analysis of variance [ANOVA] followed by Tukey’s Multiple Comparison test).

**Figure 3.** Spore germination was significantly inhibited in Fe$_{3}$O$_{4}$-treated pathogenic strains of *C. difficile*. Spores from two pathogenic strains of *C. difficile* (CCUG 19126 and ATCC BAA-1805) were treated for 20 minutes with various concentrations of Fe$_{3}$O$_{4}$ nanoparticles and then with 10 mM taurocholate to induce spore germination. Spore germination was effectively inhibited in the two Fe$_{3}$O$_{4}$-treated pathogenic strains, as it was in the nonpathogenic CCUG 37780 strain. (**P < 0.001, *P < 0.05; one-way analysis of variance [ANOVA] followed by Tukey’s Multiple Comparison test).

**Figure 4.** The viability of *C. difficile* vegetative cells was not significantly affected by Fe$_{3}$O$_{4}$-treatment. Fe$_{3}$O$_{4}$ nanoparticles (500 μg/mL) did not inhibit the viability of vegetative *C. difficile* in all three test strains compared to cells treated with 3% sodium hypochlorite. (***P < 0.001; Student’s t test).
The mechanisms underlying Fe$_{3-δ}$O$_4$-inhibited spore germination. To determine whether any additional underlying mechanisms allow Fe$_{3-δ}$O$_4$ nanoparticles to inhibit spore germination, we first excluded the sporicidal activity of the coating polymer poly-(styrene-alt-maleic acid) (PSMA) on Fe$_{3-δ}$O$_4$ nanoparticles. At a concentration higher than that on PSMA-coated Fe$_{3-δ}$O$_4$ nanoparticles, PSMA did not inhibit spore germination (Supplementary Fig. 1). In addition, to verify whether Fe$_{3-δ}$O$_4$ nanoparticles promoted or hindered the germination-inducing activity of taurocholate, 10 mM taurocholate was incubated for 20 minutes with different concentrations of Fe$_{3-δ}$O$_4$ nanoparticles and then used to treat spores. The taurocholate incubated with Fe$_{3-δ}$O$_4$ nanoparticles still induced C. difficile spore germination (Supplementary Fig. 2).

To identify any additional underlying mechanisms that allow Fe$_{3-δ}$O$_4$ nanoparticles to inhibit spore germination, we took advantage of the magnetic property of Fe$_{3-δ}$O$_4$. Some C. difficile BAA-1805 spores were incubated with Fe$_{3-δ}$O$_4$ nanoparticles bound to the surface of C. difficile (ATCC BAA-1805) spores, which allowed magnetic attraction. (A) Schematic illustration of the magnetic separation of spores with which Fe$_{3-δ}$O$_4$ nanoparticles had bound: these spores were attracted by a magnet, but spores with which the nanoparticles had not bound were retained in the supernatant. (B) The spores were incubated for 20 minutes with Fe$_{3-δ}$O$_4$ nanoparticles, and then tcdB genes were isolated from the supernatant or the magnetically concentrated precipitates were detected after PCR amplification. Lanes 1–3 are spores alone and lanes 4–6 are spores treated with 500 µg/mL of Fe$_{3-δ}$O$_4$ nanoparticles. TcdB amplicons were detected in the supernatant of the control group, but they were found only in the magnetically concentrated fraction after they had been treated with Fe$_{3-δ}$O$_4$ nanoparticles. (C) TEM images of the spores (upper panel: 10,000 ×; lower panel: 30,000 ×). The left image shows that native spores have a smooth coat; accumulated Fe$_{3-δ}$O$_4$ nanoparticles on spore surfaces are evident after incubation. The spores treated with 500 µg/mL were completely covered by the Fe$_{3-δ}$O$_4$ nanoparticles.
with 500μg/mL of Fe₃O₄ nanoparticles for 20 minutes and then separated, using a magnet, into two parts: the nonmagnetic supernatant and the magnetic pellets (Fig. 5A). Each part was collected, and PCR was used to detect the tcdB gene. The tcdB gene was detected only in the supernatant (Fig. 5B) in the spores-alone groups. In the Fe₃O₄-treated spores, the tcdB gene was also detected in the magnetic pellets.

To clarify the interaction between Fe₃O₄ and spores, the Fe₃O₄-treated spores were examined under a TEM. The TEM images showed that spores treated with 50μg/mL of Fe₃O₄ nanoparticles were only spotted with Fe₃O₄ nanoparticles, but that spores treated with 500μg/mL were almost fully surrounded (Fig. 5C). To reconfirm that Fe₃O₄ nanoparticles were simply attached to the surface of the spores instead of diffused into the spores, the spores were examined using cryo-electron tomography. The rotated dynamic video (Supplementary Video 1) revealed that Fe₃O₄ nanoparticles were directly and dose-dependently bound to the surface of C. difficile spores.

Fe₃O₄ nanoparticles as an in vivo sporicidal agent against C. difficile infection. Before the mouse experiments, the cytotoxicity of Fe₃O₄ nanoparticles was evaluated using a methylthiazol tetrazolium (MTT) assay. The colorectal cells were treated with various concentrations of Fe₃O₄ nanoparticles (5 to 500μg/mL) for 24 hours. The viability of C2BBe1 cell line were decreased at the 500μg/mL of Fe₃O₄ nanoparticles. However, the cell viability assay showed no significant cytotoxicity of the Fe₃O₄ nanoparticles in HT-29 cell line compared with the control group (Supplementary Fig. 3). To investigate the feasibility of Fe₃O₄ nanoparticles as an in vivo sporicidal agent, we used a C. difficile infection mouse model. The inflammatory signal peaked 3 days after C. difficile infection had been induced (Supplementary Fig. 4). Therefore, in the mouse experiment, we examined the C. difficile infection pathology 3 days after the mice had been treated with spores. Mice treated with Fe₃O₄ nanoparticles plus spores lost significantly less body weight than the mice treated with spores alone (P = 0.0119, Student’s t test) (Fig. 6A). The cecum weight of the Fe₃O₄-treated mice was significantly higher than that of the mice treated with spores alone (P = 0.0024, Student’s t test) (Fig. 6B), which showed minor inflammation under an in vivo imager (P = 0.0406, Student’s t test) (Fig. 6C). The histopathological images also showed that neutrophil infiltration was lower in the mice treated with Fe₃O₄ nanoparticles plus spores (Fig. 6D) than in the control mice treated with spores alone. A real-time reverse transcription quantitative polymerase chain reaction (RT-qPCR) was used to determine the expression of the proinflammatory genes TNF-α, IFN-γ, and IL-1β, compared with the spores-alone groups (Fig. 6E). The level of inflammation caused by the spores was significantly lower (P = 0.0088 for TNF-α, P = 0.0276 for IFN-γ, and P = 0.0097 for IL-1β, Student’s t test) at the molecular level in the mice treated with Fe₃O₄ nanoparticles plus spores than in the mice treated with spores alone. To verify whether treatment with 500μg/mL Fe₃O₄ nanoparticles alone induces any adverse side effects in vivo, we fed the mice with nanoparticles alone and monitored the biochemistry markers (Supplementary Fig. 5). Based on the ratio of Firmicutes to Bacteroidetes, we found this nanoparticle does not alter the gut microbiota (Supplementary Fig. 6A), suggesting that this particle did not have the toxicity to the mice.

Discussion

Based on our review of the literature, this is the first study which shows Fe-based nanoparticles that strongly inhibited the germination of C. difficile spores despite the growth inhibitory effect of other type of iron oxide nanoparticles to other bacteria vegetative cells was recently reported.

Iron oxide nanoparticles have existed in the earth’s stratum and some organisms. Due to the iron oxide nanoparticles are widely used in medical applications. The iron oxide nanoparticles not only can be conjugated to other dyes as detection reagents for different imaging systems. The iron oxide nanoparticles could be used as hyperthermia-based cancer therapeutic reagent when combined to radiofrequency. The iron oxide nanoparticles could even be modified with different materials to become the multi-functional nanoparticles, e.g., targeting, drug releasing, and imaging abilities. We performed the new function of iron oxide nanoparticles that could be a C. difficile spore germination inhibition reagent in the study. The 22-nm-Fe₃O₄ nanoparticles presented as unique octahedron single crystals, and the ratio of iron to oxide was 2.966:4, similar to our previously reported 6-nm Fe₃O₄ nanoparticles. The differences between Fe₃O₄ and FeO include size, surface chemistry, and the presence of minor non-oxidized Fe voids in the Fe₃O₄ particles. There was no significant difference between the inhibitory ability of the 22-nm and the 14-nm Fe₃O₄ doses. The amount of DPA released from the spores was significantly lower in the Fe₃O₄-treated group than in the positive controls. The transition of spores from bright to dark can be observed with a phase-contrast microscope after the DPA has been released. In the present study, because Fe₃O₄ nanoparticles inhibited spore germination, there was no transition of spores into phase-dark. However, DPA could be detected, but not in as great abundance as with the taurocholate-treated spores, after the spores had been exposed to Fe₃O₄ nanoparticles. Thus, it is conceivable that there is an underlying mechanism that enables Fe₃O₄ nanoparticles both to induce DPA release and to inhibit spore germination. We hypothesized that taurocholate can be caught by Fe₃O₄ nanoparticles and block the interaction between taurocholate and CspC. CspC is a bile salt germination receptor in C. difficile and transmits the signal to the downstream signaling molecule. However, we later discovered that taurocholate does not bind with Fe₃O₄ nanoparticles, and that taurocholate supernatant without Fe₃O₄ nanoparticles was able to trigger C. difficile spore germination. Therefore, we now hypothesize that Fe₃O₄ nanoparticles inhibit bacterial growth without directly interacting with taurocholate. We also hypothesize that Fe₃O₄ nanoparticles do not penetrate spores based on our cryo-TEM Supplementary Video 1. The details of the molecular mechanisms of Fe₃O₄ nanoparticles require additional study.

Zero-valent iron (ZVI) has recently been reported to be involved in the selective inhibition of cancerous cells while sparing noncancerous cells because it depletes mitochondrial membrane potential. Because mitochondria share distinct structural and biochemical similarities as well as evolutionary links to bacteria, we hypothesize that the nonstoichiometric non-oxidized Fe of Fe₃O₄ also inhibits spore germination. In fact, one study reported...
Figure 6. *C. difficile* infection-induced colitis was attenuated in mice treated with Fe₃O₄ nanoparticles. Purified CCUG 19126 spores were incubated for 20 minutes with Fe₃O₄ nanoparticles, and then the mice were injected with them. Three days later, the mice were killed. (A) The mean bodyweight-loss of the Fe₃O₄ nanoparticle-treated mice was significantly lower than that of the spores-alone control mice. (B) The mean weight-loss of the cecums of the Fe₃O₄ nanoparticle-treated mice was also significantly lower than that of spores-alone control mice. (C) The inflammation of the isolated colon from 11FNL/FVB/NJ transgenic mice with NF-κB-signal-activated bioluminescence was detected. The left image shows a spores-alone control colon and the right image a Fe₃O₄-nanoparticle-treated colon. *C. difficile* infection-induced inflammation was significantly lower in the Fe₃O₄ nanoparticle-treated colon (right panel). (D) A histopathological examination of the isolated colon tissue shows prominent infiltration of neutrophils (black arrows in left 400× panel) in the control group, but attenuated inflammation in the Fe₃O₄-nanoparticle-treated group (right 400× panel). Neutrophils were morphometrically counted in 10 randomly selected fields in the tissue sections. The number
of neutrophils in the control group was 16 ± 5 per 170 m × 130 μm field; the number in the Fe3O4 nanoparticles-treated group was 2 ± 1 per 170 m × 130 μm field. (E) A reverse transcription-quantitative polymerase chain reaction (RT-QPCR) shows that the total RNA extracted from the Fe3O4 nanoparticle-treated spores expressed significantly lower levels of proinflammatory genes than did the spores in the control group. (*P < 0.05, **P < 0.01; Student's t test) (n = 6).

a prominent bactericidal effect of ZVI-based nanoparticles on E. coli because they effectively caused oxidative damage to bacterial membranes. Our study showed that Fe3O4 treatment significantly inhibited the germination of C. difficile spores by damaging spore membranes but not the vegetative cells and therefore did not damage the balance of intestinal normal flora. Thus the specific activity to spore germination must go through other yet to be discovered mechanisms.

It is well known that the expression of the tcdA and tcdB genes is closely associated with the virulence of C. difficile infection. We found that Fe3O4 nanoparticles inhibited germination in both the tcdA(−)tcdB(−) strain and the more virulent tcdA(+)tcdB(+) strain. In an in vivo C. difficile infection mouse experiment, we found that Fe3O4 significantly reduced the inflammation indices in the groups treated with spores +Fe3O4, and that it effectively inhibited C. difficile spore germination, thus controlling in vivo the virulence of C. difficile. Intriguingly, treatment with 500 μg/mL of Fe3O4 nanoparticles did not kill vegetative cells.

Because antibiotics are overserved everywhere in the world, multidrug-resistant microbes have emerged as a major public health threat46. Nanomedicines have inspired various new strategies to control microbes without allowing them to develop drug resistance46. Bacterial spores are even more resistant than are the bacteria themselves to harsh environments and antibiotics, but nanoparticle-spore interaction and potential clinical applications are rarely discussed. The germination of Bacillus subtilis spores is greatly compromised by treatment with polyionic polymers and then silica nanoparticles47. Such treatment, however, does not affect spore viability. We also found that excess PSMA alone did not significantly inhibit the germination of spores; therefore, PSMA must not be important for inhibiting spore germination. In the present study, 500 μg/mL of Fe3O4 nanoparticles were sufficient to significantly, and almost as efficaciously as 3% sodium hypochlorite, inhibit spore germination, which indicated the potential of Fe3O4 nanoparticles for clinical development as a novel and effective treatment for managing and perhaps even preventing C. difficile infection.

Many studies showed their materials existed excellent antibacterial activity, but sometimes the impact of the materials on the normal flora and biocompatibility were rarely discussed. The guts normal flora plays a vital role in animals, because it attend the metabolism and pathogenic colonization prevention in the host48. Therefore, we should have to understand the potential adverse effect of Fe3O4 nanoparticles-treated group was 2 ± 1 per 170 m × 130 μm field. The two main phyla of Firmicutes and Bacteroidetes are the common bacteria in the large intestine48,49. Our results showed that Fe3O4 nanoparticles did not damage the normal flora destruction after administering to human and other animals. We also found excellent biocompatibility between Fe3O4 nanoparticles and a variety of colorectal cells. We previously reported40,41 that Fe3O4 nanoparticles did not damage their vegetative cells. Another study42 reported that a high concentration (3 mg/mL) of Fe3O4 nanoparticles for 72 hours, RT-qPCR was used to analyze the results of normal flora population changes. The two main phyla of bacteria in the gut are Firmicutes and Bacteroidetes45. We found that the Fe3O4 nanoparticles did not change the ratio of Firmicutes to Bacteroidetes in the mice (Supplementary Fig. 6A). Bacteroides fragilis and Enterococcus faecalis are the common bacteria in the large intestine46,48,49. Our results showed that Fe3O4 nanoparticles did not cause damage to the two bacteria (Supplementary Fig. 6B). According to our results we hypothesize that Fe3O4 nanoparticles do not cause imbalance of gut microbiota. The Fe3O4 nanoparticles will not be an issue in the normal flora destruction after administering to human and other animals. We also found excellent biocompatibility between Fe3O4 nanoparticles and a variety of colorectal cells. We previously reported40,41 that Fe3O4 nanoparticles used with MRI scans and cancer therapy were safe. Although the safe dosages are different in these three studies, we have shown that Fe3O4 nanoparticles are biocompatible and safe in vivo.

Metronidazole and vancomycin are antibiotics frequently used to treat patients with C. difficile infection, but the some clinical isolates are metronidazole-resistant46. Antibiotics unselectively damage pathogens as well as normal flora and might trigger a highly contagious state of super-shedding C. difficile spores, thus increasing the risk of infection47. Spores can survive for many years in harsh environments and wait to be ingested by animals48. The imbalance of normal flora and spore germination might be the major cause of a C. difficile infection relapse. Moreover, bowel inflammation has long been recognized as an important index for C. difficile infection severity and prognosis. Therefore, we quantitatively assessed C. difficile infection by directly injecting mice with spores, which is most relevant to the pathogenesis of clinical C. difficile infection. Transgenic mice with NF-κB-dependent luciferase reporter enabled us to visualize bowel inflammation using the IVIS imaging system. Because of their unique ability to selectively kill spores without damaging normal flora, mammalian cells, or the mouse gastrointestinal tract, Fe3O4 nanoparticles offer a new strategy for controlling infectious clinical diseases. Fe3O4 nanoparticle treatment not only inhibited the germination of different clinical strains of C. difficile spores in vitro, but also reduced the virulence of spores that induced C. difficile infection in vivo. We also report the first relatively low-dose but efficacious Fe-based nanoparticle inhibition of spore germination, one which uses a novel mechanism that does not interfere with the growth of healthy vegetative flora, which is an important clinical consideration in C. difficile infection management. We hypothesize that coating the surfaces of medical devices with Fe3O4 nanoparticles will reduce the threat of spores in hospitals. Additional research on improving the design of Fe3O4 nanoparticles should help nanoparticle-based therapeutics become safer and more effective, and should contribute to the development of new therapeutics for C. difficile infection and other emerging infectious diseases.
Methods

Ethics statement and animals. All NF-κB-dependent reporter mice (FVB/N) or naïve genetic background) were obtained from the National Laboratory Animal Center in Taiwan and housed in a pathogen-free barrier facility. All animal experiments were approved and in accordance with the relevant guidelines and regulations required by the Institutional Animal Care and Use Committee of National Cheng Kung University (NCKU-IACUC-102-296).

Bacterial culture and spore purification. *C. difficile* CCUG 37780 (tcdA+, tcdB+) and CCUG 19126 (tcdA−, tcdB+) were purchased from the Culture Collections of the University of Göteborg (Göteborg, Sweden), and BAA-1805 (tcdA−, tcdB+) from American Type Culture Collection (Manassas, VA). All strains were incubated in brain-heart infusion-supplemented medium (BHIS; BD Difco, Franklin Lakes, NJ), with 0.5% yeast extract (BD Difco) and 0.1% L-cysteine (Amresco, Solon, OH), at 37 °C under anaerobic conditions. The spores were prepared and purified as previously described with a slight modification. Briefly, *C. difficile* in BHIS medium was diluted in fresh BHIS medium to an optical density (OD) (600 nm) of 0.2. The diluted bacterial suspension (900 μL) was added to a 6-well dish with BHIS agar, and then the dish was incubated at 37 °C in an anaerobic jar (Thermo Fisher, Oxoid Ltd., Basingstoke, England) for 4 days. The whole cells were washed 5 times with ice-cold sterile water, the bacteria were resuspended with 3 mL of ice-cold sterile MQ water. The suspension was spread on top of a 10-mL 50% (wt/vol) sucrose solution (J.T. Baker Chemical, Phillipsburg, PA) in a centrifuge tube, and then centrifuged at 3500 g for 20 minutes to separate spores from vegetative cells. The purified spores at the bottom of the centrifuge tube were washed 5 times with ice-cold sterile water to remove sucrose, and then stored at 4 °C.

Preparation of the nanoparticles. Fe3O4 nanoparticles (22 nm and 14 nm) were synthesized using thermal decomposition, as previously described. Briefly, 1.42 g of iron acetylacetonate was mixed with 0.57 mL of oleic acid and 20 mL of trietylamine (all 3: Sigma-Aldrich, St. Louis, MO). The solution was refluxed at 325 °C in an argon environment for 30 minutes. After the solution had cooled down to room temperature, the precipitates were collected with a magnet and washed 3 times with toluene. The Fe3O4 nanoparticles were collected with a magnet and then transferred to chloroform solutions (Merck, Whitehouse Station, NJ) containing 0.4 mg/mL of PSMA (Sigma-Aldrich) and were left there for 4 hours at 55 °C. The Fe3O4 nanoparticles were collected, washed 3 times with MQ water, and then stored at 4 °C.

Optical density based spore germination test. Spores were germinated as previously described. Before the germination experiments, the spore suspension was incubated at 60°C for 30 minutes. The heat-treated spores were then moved to ice. The *C. difficile* spores (concentration: OD600, 0.5) were co-incubated with various nanoparticles or at different concentrations (5 to 500 μg/mL) in BHIS in a 96-well plate for 20 minutes. The concentration of Fe3O4 nanoparticles was measured based on the total particle weight per 1 mL of solution. Fe3O4 (22-nm) nanoparticles were used here. Spores treated with 3% bleach (Wako, Osaka, Japan) were the positive control to inhibit germination, and spores treated with BHIS only were the negative controls. After the spore + nanoparticle co-incubation, the spores were treated with 10 mM taurocholate (Sigma-Aldrich) to induce germination. The OD600 of treated spores was kinetically determined at 1 minute intervals using a spectrophotometer (TECAN, Grödig, Austria) at room temperature for 12 minutes. The OD at different time points was used to plot the spore germination curve.

Phase-contrast microscopy observation of spore germination. The images of spore germination were recorded using phase-contrast optical microscopy described elsewhere. The purified spores were incubated at 60°C for 30 minutes. The spores were co-incubated for 20 minutes with 500 μg/mL of Fe3O4 nanoparticles or with buffer alone. Taurocholate was then added to the samples to a final concentration of 10 mM in BHIS. Five microliters of samples were then dropped onto the 0.7% agarose surface on the glass slide. After 15 minutes, phase-contrast images of the spores were captured (Nikon Eclipse 80i; Tokyo, Japan). Six random fields of view were obtained for each experimental group.

Dipicolinic acid (DPA) release assay. The DPA release was estimated by modified method from a previous study. Briefly, the spores were heated as described and washed with distilled water then resuspended in spore germination buffer (10 mM Tris [pH 7.5], 150 mM NaCl, and 100 mM glycine). The spores were treated with MQ water or 500 μg/mL of Fe3O4 nanoparticles for 20 minutes in a black 96-well plate before taurocholate (10 mM) and terbium (III) chloride (TbCl3) (100 μM) were added. The total DPA of the spores was extracted by boiling the samples for 30 minutes. The spores treated with TmCl3 only were negative controls. The DPA-Tb signal was monitored immediately in real-time using a microplate reader (FlexStation 3 Multi-Mode; Molecular Devices, CA) with excitation/emission at 270 nm and 545 nm, and a cutoff at 530 nm.

Spore viability analysis. After the spore germination curves had been recorded, each aliquot was removed from the 96-well plate and plated on BHIS agar dishes and serially diluted. The plate-counts (CFU/mL) at each dilution were done after 48 hours of incubation in an anaerobic environment. The inhibition rates were calculated using the following formula:

\[ 1 - \left( \frac{\text{treatment CFU}}{\text{control CFU}} \right) \times 100\% \]
Nanoparticle-spore binding analysis.  

C. difficile BAA-1805 spores at a concentration of OD_560=0.5 were co-incubated with Fe_3O_4 nanoparticles (500 μg/mL) in BHIS in a 96-well plate for 20 minutes. The spore samples without Fe_3O_4 treatment were controls. The samples were placed next to a magnet for 5 minutes and then all supernatant was removed to other tubes. The magnet-attached parts and supernatant were washed 3 times with 1× phosphate-buffered saline. All samples were resolved in distilled deionized water, and the tcdB DNA in samples was detected using a polymerase chain reaction (PCR). Spore and nanoparticle images were captured using a transmission electron microscope (TEM) (JEM-1400; JEOL, Tokyo, Japan).

In vivo analysis of C. difficile infection.  
To directly monitor the colonic inflammation, we infected mice with C. difficile spores in a previously generated NF-κB-dependent reporter mouse model containing the luciferase transgene under the transcriptional control of NF-κB (NF-κB-RE-luciferase)²⁵. Before the mice were fed the spores with NF-κB reporter, they were given an antibiotic cocktail (0.4 mg/mL of kanamycin, 0.035 mg/mL of gentamicin, and 0.057 mg/mL of colistin) in their drinking water for 48 hours and a refreshed cocktail every 24 hours for 2 days. The mice were gavaged with a 200-μL proton pump inhibitor (PPI) (2 mg/mL) every 12 hours for 2 days before the C. difficile spore infection. C. difficile CCUG 19126 spores (2×10⁶ CFUs) were co-incubated with or without 500 μg/mL of Fe_3O_4 nanoparticles for 20 minutes before they were gavaged with 100 L of sample solution. While the spores and Fe_3O_4 nanoparticles were incubating, all the mice were gavaged with a 50-μL PPI (2 mg/mL) and then intraperitoneally injected with clindamycin (4 mg/kg). The antibiotic cocktail water was replaced with normal water after the C. difficile infection. All mice were monitored for C. difficile infection symptoms, e.g., diarrhea, weight loss, hunched posture, and death. Seventy-two hours post-infection, the mice were intraperitoneally infected with 150 mg/kg of luciferin (PerkinElmer, Waltham, MA) to show NF-κB activation-mediated luminescence. The mice were anesthetized with isoflurane and oxygen, and then images were collected for 5 minutes using an imaging system (Xenogen IVIS® Spectrum; Advanced Molecular Vision, Grantham, Lincolnshire, UK). Data were analyzed (Xenogen Living Image®), and luciferase activity was presented in photons/sec/cm²/stereadian (p/s/cm²/sr). After the IVIS images had been obtained, a reagent (TRI-Sigma–Aldrich) was used to extract RNA from colon tissue samples. The levels of inflammatory gene expression were estimated using a real-time PCR assay (StepOnePlus; Applied Biosystems).

Histopathology examination.  
Histopathological analysis was used to evaluate C. difficile infection-induced mucosal damage and inflammation. Resected colon tissue samples were fixed in 4% formaldehyde buffered with PBS and then embedded in paraffin. Deparaffinized 6-μm-thick sections were stained with hematoxylin and eosin. The results were captured using optical microscopy. Neutrophils were randomly counted in 10 fields for both the spores-alone group and the Fe_3O_4-treated-spores group.

Statistical analysis.  
GraphPad Prism 5.01 was used for all statistical analyses. All experiments were done in triplicate. Data are means ± standard error of the mean (SEM) from three independent experiments. One-way analysis of variance (ANOVA) and then Tukey’s Multiple Comparison test were used for spore germination curve analyses, and Student’s t test was used in the other CFU inhibition test.

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Competing Interests:

Additional Information

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Author Contributions

D.-B. Shiieh and P.-J. Tsai form the concept, designed the study and edited the manuscript. W.-T. Lee did most experiments and draft the manuscript. Y.-N. Wu helped the organization of the manuscript and perform the primary language editing. Y.-H. Chen participated in most animal tests. S.-R. Wu performed all TEM and cryo-EM imaging and interpretation. T.-M. Shih performed part of the spore germination test. T.-J. Li and L.-X. Yang participated in results discussion and part of the animal tests. C.-S. Yeh provided nanoparticles fabrication protocol and guided the synthesis detail.

Additional Information

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