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Low molecular weight chitosan-coated silver nanoparticles are effective for the treatment of MRSA-infected wounds

Abstract: Silver nanoparticles (AgNPs) are being widely applied as topical wound materials; however, accumulated deposition of silver in the liver, spleen, and other main organs may lead to organ damage and dysfunction. We report here that low molecular weight chitosan-coated silver nanoparticles (LMWC-AgNPs) are effective against methicillin-resistant Staphylococcus aureus (MRSA), have better biocompatibility, and have lower body absorption characteristics when compared with polyvinylpyrrolidone-coated silver nanoparticles (PVP-AgNPs) and silver nanoparticles without surface stabilizer (uncoated-AgNPs) in a dorsal MRSA wound infection mouse model. LMWC-AgNPs were synthesized by reducing silver nitrate with low molecular weight chitosan as a stabilizer and reducing agent, while PVP-AgNPs were synthesized using polyvinylpyrrolidone as a stabilizer and ethanol as a reducing agent. AgNPs with different surface stabilizers were identified by UV-visible absorption spectrometry, and particle size was determined by transmission electron microscopy. UV-visible absorption spectra of LMWC-AgNPs, PVP-AgNPs and uncoated-AgNPs were similar and their sizes were in the range of 10–30 nm. In vitro experiments showed that the three types of AgNPs had similar MRSA-killing effects, with obvious effect at 4 μg/mL and 100% effect at 8 μg/mL. Bacteriostatic annulus experiments also showed that all the three types of AgNPs had similar antibacterial inhibitory effect at 10 μg/mL. Cell counting kit-8 assay and Hoechst/propidium iodide (PI) staining showed that LMWC-AgNPs were significantly less toxic to human fibroblasts than PVP-AgNPs and uncoated-AgNPs. Treatment of mice with MRSA wound infection demonstrated that the three types of AgNPs effectively controlled MRSA wound infection and promoted wound healing. After continuous application for 14 days, LMWC-AgNPs-treated mice showed significantly reduced liver dysfunction as demonstrated by the reduced alanine aminotransferase and aspartate aminotransferase levels and liver deposition of silver, in comparison to mice treated with uncoated-AgNPs or PVP-AgNPs. Our results demonstrated that LMWC-AgNPs had good anti-MRSA effects, while harboring a better biocompatibility and lowering the body’s absorption characteristics.

Keywords: silver nanoparticles, LMWC-AgNPs, MRSA, PVP-AgNPs, PVP, biocompatibility

Introduction

Silver nanoparticles (AgNPs) are increasingly being used as various types of external trauma materials due to their effective antimicrobial and good release properties. However, the toxicology of AgNPs remains largely unknown. Moreover, AgNPs can be absorbed into the bloodstream via different routes of administration, leading to deposition of silver in many organs, including the liver and spleen, and potential organ damage after a certain amount of accumulation. Therefore, in the actual
application process, protecting their antibacterial properties while reducing their absorption and toxicity in the body has become a very important challenge.

Previous studies have shown that different surface stabilizers have distinct impacts on AgNP cytotoxicity. Because of its good biocompatibility and antibacterial properties, chitosan is often used as the active ingredient of topical wound materials. It has been reported that chitosan is a good stabilizer for AgNPs and promotes wound healing by AgNPs. Moreover, chitosan-coated AgNPs demonstrated high efficiency in killing common gram-positive and -negative bacteria, and fungi. Compared with a higher degree of polymerization of chitosan, low molecular weight chitosan (LMWC) has better water solubility and biological activity, suggesting that LMWC may be an ideal stabilizer of AgNPs.

Methicillin-resistant Staphylococcus aureus (MRSA) infection is the most common clinical drug-resistant infection in the burn ward, with MRSA wound infection rate accounting for more than 60%; moreover, the number of different MRSA strains isolated sharply increases and there is a growing number of multidrug-resistant MRSA. Thus, control of MRSA wound infection has become a global health problem. A previous study has shown that AgNPs have a great anti-bactericidal activity against MRSA. In order to obtain more safe and effective AgNP products for the clinical management of MRSA infection, in this study, we used LMWC as a reducing agent and a stabilizer to synthesize AgNPs, and compared its effects on the antibacterial activity, biocompatibility, and body absorption of AgNPs with different surface stabilizers of the same particle size.

Materials and methods
Synthesis of LMWC
Medical grade chitosan with 95% degree of deacetylation was provided by Oriental Medical Institute (Qidong, Jiangsu, People’s Republic of China). Three grams of chitosan was dissolved in 100 mL 3% acetic acid solution, followed by addition of 60 mL 30% H2O2 and supplemented with 300 mL deionized water. After incubation at 50°C in a water bath for 2 hours, the reaction was cooled in ice water, adjusted to pH 7.0 with 2 M NaOH, and filtered. The LMWC was precipitated with 900–1,500 mL of 100% ethanol and dried by vacuum. The molecular weight and traits of synthesized LMWC were determined by high-performance liquid chromatography (Waters 1525, UK) and Fourier transform infrared spectroscopy (FTIR, Nicolet 6700; Thermo Fisher Scientific, Waltham, MA, USA), respectively.

Synthesis of low molecular weight chitosan-coated silver nanoparticles
Four grams of LMWC was mixed with 0.1 g AgNO3 in 50 mL deionized water at 50°C under stirring at 200 rpm overnight in a hood. On the next day, acetone was added slowly with stirring until the solution became cloudy, followed by centrifugation at 2,000 rpm for 30 minutes. The supernatant was transferred to a new tube, dried by vacuum, and dissolved in 2–3 mL deionized water. The concentration of the resulting low molecular weight chitosan-coated silver nanoparticles (LMWC-AgNPs) was determined by UV-Vis (UV3600; Shimadzu Corporation, Kyoto, Japan) and ICP-MS (Agilent Technologies, Richardson, TX, USA). The particle size of LMWC-AgNPs was determined by transmission electron microscopy (TEM) (JEM-100S; JEOL, Tokyo, Japan).

Synthesis of polyvinylpyrrolidone-coated silver nanoparticles
One gram of polyvinylpyrrolidone (MW58000; Aladdin, People’s Republic of China) was completely dissolved in 25 mL of absolute ethanol at 80°C for 1 hour, followed by addition of 0.2 g AgNO3. The reaction continued under stirring (400 rpm) until the brown color became stable (about 1–2 hours). Then, a total volume of 5–10 fold of acetone was added slowly with stirring until the solution became cloudy, followed by centrifugation at 2,000 rpm for 30 minutes. The supernatant was transferred to a new tube, dried by vacuum, and dissolved in 2–3 mL of deionized water. The concentration of the resulting polyvinylpyrrolidone-coated silver nanoparticles (PVP-AgNPs) was determined by UV-Vis and ICP-MS. The particle size of PVP-AgNPs was determined by TEM.

Cell proliferation assay
Silver nanoparticles without surface stabilizer (uncoated-AgNPs) were purchased from Shanghai Nanotechnology Ltd (Shanghai, People’s Republic of China). Human fibroblast HS27 cells were from Shanghai Bioleaf Biotech (Shanghai, People’s Republic of China) and cultured in Dulbecco’s Modified Eagle’s Medium supplemented with 10% fetal bovine serum, 100 U/ml penicillin, and 100 μg/mL streptomycin under a humidified 5% CO2 atmosphere at 37°C in an incubator. Cells were seeded in triplicates in 96-well plates (1×104 cells in 100 μL medium per well). After 24 hours incubation, the cells were added to 10 μL of different AgNPs. After 24 hours, the cell counting kit-8 (CCK-8) assay was performed according to the manufacturer’s recommendations. The absorbency was measured with Infinite M200 Pro
multifunctional microplate reader (Tecan, Männedorf, Switzerland) at a wavelength of 450 nm. All assays were performed in triplicates.

Detection of cell apoptosis assay by Hoechst/propidium iodide staining
Human fibroblast HS27 cells were cultured to 80% confluence and treated with 30 μg/mL of three types of AgNPs for 24 hours. After washing three times with phosphate-buffered saline, the cells were added to 10 μL Hoechst 33258 (5 μg/mL) and incubated at 37°C for 10 minutes, followed by adding 5 μL PI (15 μg/mL) in the dark at room temperature for 10 minutes. The cells were observed under a fluorescence microscope (Ti-S; Nikon, Minato, Japan).

In vitro bacterial killing
A bioluminescent MRSA strain derived from ATCC 33591 was a gift from Michael Hamblin’s lab (Wellman Center for Photomedicine, MGH, Boston, MA, USA). In total, 0.5 mL of logarithmic growth phase MRSA (1.5×10^8 cfu/mL) suspension was mixed with 0.5 mL organic interfering substances (3% bovine serum albumin) and incubated in a 20°C±1°C water bath for 5 minutes. The bacterial solution was added with 4.0 mL of different concentrations of three types of AgNPs and incubated for 30 minutes. Also, 0.5 mL of the mixture was mixed with 4.5 mL of sterile saline and incubated for 10 minutes at room temperature. In total, 100 μL of the sample solution was coated on a flat plate, incubated overnight at 35°C, and the number of bacterial colonies was calculated. For filter paper assay (bacterial killing measurement), 20 μL of 10 μg/mL AgNP solution was dropped onto a 6 mm sterile filter paper, dried at 37°C, and placed onto a dish densely coated with logarithmic growth phase MRSA (1.5×10^8 cfu/mL) suspension with a sterile cotton swab. The dish containing MRSA and AgNP-filter paper was placed at 35°C 5% CO₂ incubator and inhibition zone size was observed 48 hours later.

Animal experiments
The Animal Committee of Ninth People’s Hospital Affiliated to Shanghai Jiao Tong University, School of Medicine approved all the experimental protocols and animal handling procedures. All experimental procedures and postoperative animal care were conducted in accordance with the National Institute of Health’s Guidelines for the Care and Use of Laboratory Animals. Balb/c mice were purchased from SLAC Laboratory Animal Technology (Shanghai, People’s Republic of China). All mice were maintained on a 12-hour light/dark cycle in a room at 22°C–25°C and allowed free access to food and water before surgery. Balb/c mice with 1.5×1.5 cm back full-thickness skin wounds were inoculated with 100 μL of 9×10^8 cfu/mL MRSA bacteria, and infection of mice was assessed 3 days later with Fx CCD luminescent camera (Vilber Lourmat, Collégien, France). The mice were randomly divided into groups. The infected wounds were dressed with 50 μg/g-bw of different AgNPs, bandaged with double gauze bandage, and fixed with 3 M film. In the following 2 consecutive days, Fx CCD luminescent camera was used to observe wound infection and healing. The dressing contained AgNPs was changed every other day. On day 14, fresh blood was taken from mouse eyeballs and centrifuged at 1,500 rpm for 10 minutes. The supernatant sera were subjected to liver function analysis with DXC800 Beckman automatic biochemical analyzer (Beckman Coulter, USA). At the same time, a portion of liver tissue and healed skin tissue on the wounded area was removed, dissolved in 69% concentrated nitric acid solution, and subjected to silver content determination by ICP-MS in order to assess the body’s absorption of AgNPs.

Statistical analyses
Graphs were constructed using GraphPad Prism (GraphPad Software, San Diego, CA, USA). All data are presented as mean ± standard error of mean. Statistical significance was determined by unpaired two-tailed t-tests or two-way analysis of variance. P<0.05 was considered statistically significant.

Results
The characteristics of LMWC-AgNPs, PVP-AgNPs, and uncoated-AgNPs
FTIR tests showed that the depolymerization during the process of production of LMWC did not change the molecular structure of chitosan, with only molecular chain rupture, as demonstrated by similar FTIR spectra characteristic of LMWC and chitosan (Figure 1A). The molecular weight of LMWC was between 2,000 and 2,480 g/mol, mainly in the vicinity of 2,300 g/mol (Figure 1B). As the molecular weight of chitosan monomer is 161.2 g/mol, these results indicated that our synthesized LMWC was a chain of 12–15 chitosan sugar monomers. To compare the cytotoxicity of LMWC with chitosan, human fibroblast HS27 cells were treated with different concentrations of LMWC or chitosan (200–1,000 μg/mL) for 24 hours, and cell viability was assessed by CCK-8 assay. As shown in Figure 1C, LMWC was apparently less toxic to HS27 cells than chitosan. UV-Vis analysis demonstrated that absorption curves of
LMWC-AgNPs, PVP-AgNPs, and uncoated-AgNPs were similar, with maximum absorption peaks at 414, 408, and 421 nm, respectively (Figure 2). The particle size of LMWC-AgNPs (Figure 3A), PVP-AgNPs (Figure 3C), and uncoated-AgNPs (Figure 3E) was detected by TEM. Further, ImageJ software analysis of the size and distribution of AgNPs of LMWC-AgNPs (Figure 3B), PVP-AgNPs (Figure 3D), and uncoated-AgNPs (Figure 3F) showed that their sizes were in the range of 10–30 nm without significant difference.

**Figure 1** Characterization of molecular weight distribution of LMWC.

**Notes:** (A) FTIR spectra of chitosan (full line) and LMWC (short dash line) showed similar absorption. (B) FTIR analysis of molecular weight distribution of LMWC. (C) H527 cells were treated with the indicated concentrations of LMWC or chitosan for 24 hours. Cell viability was assessed by CCK-8 assay.

**Abbreviations:** FTIR, Fourier transform infrared spectroscopy; LMWC, low-molecular weight chitosan; CCK-8, cell counting kit-8; MW, molecular weight.

**Figure 2** UV-Vis absorption spectra of uncoated-AgNPs (black full line), PVP-AgNPs (gray full line), and LMWC-AgNPs (black dash line).

**Abbreviations:** Uncoated-AgNPs, silver nanoparticles without surface stabilizer; PVP-AgNPs, polyvinylpyrrolidone-coated silver nanoparticles; LMWC-AgNPs, low molecular weight chitosan-coated silver nanoparticles.

LMWC-AgNPs, PVP-AgNPs, and uncoated-AgNPs have similar bactericidal effects

The anti-bactericidal effects of the three types of AgNPs were similar, with an IC50 of approximately 4 μg/mL for all the
three types. At 8 μg/mL, the anti-bactericidal effect reached 100%, and there was no growth of bacteria following incubation with 10 μg/mL of any of the three types of AgNPs for 30 minutes (Figure 4A and B). Similar bactericidal effects of the three types of AgNPs were observed by bacteriostatic ring method (Figure 4C). In vivo bactericidal results showed that, on the third day, the amounts of bacteria in wounds on the back of mice were similar in the three types of AgNPs treatments, which were demonstrated by the similar values of MRSA chemiluminescent in the three groups (Figure 4D).

**LMWc-agNPs are less toxic to human fibroblasts than PVP-AgNPs and uncoated-AgNPs**

To test the cytotoxicity of three types of AgNPs on human fibroblasts, HS27 cells were treated with different concentrations of the three types of AgNPs for 24 hours and cell viability was assessed by CCK-8 assay. As shown in Figure 5A, the cell viability of HS27 cells treated with 10 and 20 μg/mL of uncoated-AgNPs or PVP-AgNPs was 98.83%±7.82%, 86.47%±16.92% or 99.84%±7.82%, 89.47%±13.92%, respectively. Uncoated-AgNPs and PVP-AgNPs at 30 μg/mL were obviously toxic to cells with cell viability decreased to 62.75%±14.75% and 68.64%±11.03%, respectively. At 50 μg/mL of either uncoated-AgNPs or PVP-AgNPs, the majority of cells were dead and floating with cell viability of only 16.08%±4.05% and 13.12%±3.75%, respectively. However, the cell viability of cells treated with 10, 20, 30, 40, and 50 μg/mL of LMWC-AgNPs was 109.84%±7.82%, 107.46%±13.92%, 96.75%±11.65%, 85.08%±8.55%, and 66.08%±13.05%, respectively. There were no statistical differences in cell viability among the three types of AgNP treatments with lower concentrations (below 20 μg/mL). Detection of apoptosis by Hoechst/PI staining of cells treated with 30 μg/mL AgNPs for 24 hours showed similar apoptosis rates for uncoated-AgNPs and PVP-AgNPs, while lower apoptosis rate for LMWC-AgNPs when compared with uncoated-AgNPs and PVP-AgNPs (Figure 5B). These results indicated that LMWC-AgNPs were less toxic to human fibroblasts cells compared to uncoated-AgNPs and PVP-AgNPs.

**LMWC-AgNPs, PVP-AgNPs, and uncoated-AgNPs effectively control MRSA wound infection and promote wound healing without any apparent difference**

Topical wound healing rate showed that, in comparison to the untreated control group, each group treated with AgNPs displayed accelerated wound healing on day 3 (Figure 6A and B). On day 14, most wounds healed in all the groups treated with AgNPs (Figure 6A and B). Thus, the three types of AgNPs effectively controlled MRSA wound infection and promoted wound healing without apparent difference.
Figure 4 The anti-bactericidal effects of three types of AgNPs. (A) A bioluminescent MRSA strain was treated with the indicated concentrations of the three types of AgNPs for 48 hours. The viability of MRSA was calculated by RFU. (B) A set of representative photos of three independent experiments of (A) with 10 μg/mL of AgNPs. (C) Analysis of bactericidal effects of the three types of AgNPs by bacteriostatic ring method. (D) Mice wounds were inoculated with MRSA and dressed with different AgNPs. From day 3 to day 5, the amounts of bacteria in wounds on the backs of mice were determined by Fx CCD chemiluminescence assay. Higher intensity of the red color indicates more severe infection. Scale bar means 1 cm.

Abbreviations: AgNPs, silver nanoparticles; MRSA, methicillin-resistant Staphylococcus aureus; RFU, relative fluorescence units; Uncoated-AgNPs, silver nanoparticles without surface stabilizer; PVP-AgNPs, polyvinylpyrrolidone-coated silver nanoparticles; LMWC-AgNPs, low molecular weight chitosan-coated silver nanoparticles; D, day.
Figure 5 The cytotoxicity of the three types of AgNPs on human fibroblast HS27 cells. (A) HS27 cells were treated with the indicated concentrations of the three types of AgNPs for 24 hours. Cell viability was assessed by CCK-8 assay. (B) HS27 cells were treated with 30 μg/mL of the three types of AgNPs for 24 hours. Cell apoptosis was assessed by Hoechst/PI staining. Scale bar means 200 μm.

Abbreviations: AgNPs, silver nanoparticles; CCK-8, cell counting kit-8; PI, propidium iodide; Uncoated-AgNPs, silver nanoparticles without surface stabilizer; PVP-AgNPs, polyvinylpyrrolidone-coated silver nanoparticles; LMWC-AgNPs, low molecular weight chitosan-coated silver nanoparticles.

Figure 6 The effects of LMWC-AgNPs, PVP-AgNPs, and uncoated-AgNPs on the control of MRSA wound infection and wound healing in mice. (A) Fx CCD luminescent camera images of wound infection and healing following dressing with different AgNPs for the indicated days. Scale bar means 1 cm. (B) Quantitation of the topical wound healing rate. (C) After application of AgNPs for 11 days, the ALT and AST levels in the blood of mice were detected for liver function. After applying AgNPs six times over 11 days, the ALT levels in the blood of mice were detected for liver function. *P<0.05; **P<0.01. (D) Mouse liver tissues and healed skin tissue at wounded area were removed after continuous application of AgNPs, on day 14, and subjected to silver content determination by ICP-MS. *P<0.05 in liver; #P<0.05 in skin.

Abbreviations: LMWC-AgNPs, low molecular weight chitosan-coated silver nanoparticles; PVP-AgNPs, polyvinylpyrrolidone-coated silver nanoparticles; uncoated-AgNPs, silver nanoparticles without surface stabilizer; MRSA, methicillin-resistant Staphylococcus aureus; AgNPs, silver nanoparticles; ALT, alanine aminotransferase; AST, aspartate aminotransferase; D, day; ICP-MS, inductively coupled plasma mass spectrometry.
LMWC-AgNPs-treated mice have significantly reduced liver deposition of silver in comparison to mice treated with uncoated-AgNPs or PVP-AgNPs

After six times of application of AgNPs for 11 days (D3/D5/D7/D9/D11/D13), the blood of mice was analysed for liver function by assessing alanine aminotransferase (ALT) and aspartate aminotransferase (AST) levels. The ALT values of mice in the uncoated-AgNP and PVP-AgNP groups were 47.33±4.95 and 48.30±6.18 U/L, respectively, while the ALT value of mice in the LMWC-AgNP group was 38.00±8.03 U/L, which was significantly lower than the uncoated-AgNP and PVP-AgNP groups (P<0.05). Similar results were obtained for AST values (Figure 6C). In order to assess the body’s absorption of AgNPs, mouse liver tissues and healed skin tissue at the wounded area were removed on day 14 after continuous application of AgNPs on day 14, dissolved in 69% concentrated nitric acid solution, and subjected to silver content determination by ICP-MS. The liver silver content in the mice treated with uncoated-AgNPs and PVP-AgNPs was 2,765.85±853.49 and 2,898.48±918.96 ng/g, respectively, while that in the mice treated with LMWC-AgNPs was 1,827.97±294.16 ng/g, which was significantly lower than the uncoated-AgNP and PVP-AgNP groups (P<0.05, Figure 6D). Further analysis of the silver content of the newly healed skin tissues revealed that the silver content in the newly healed skin tissues of the mice treated with uncoated-AgNPs and PVP-AgNPs was 1,670.79±850.66 and 1,509.81±612.51 ng/g, respectively, while the silver content in the mice treated with LMWC-AgNPs was 837.35±371.36 ng/g, which was significantly lower than the uncoated-AgNP and PVP-AgNP groups (P<0.05, Figure 6D).

Discussion

Due to their effective antibacterial property, AgNPs are widely used as various types of medical materials including trauma topical material.4,17,18 A large number of previous studies have shown that, compared with the silver ions, the cytotoxicity of AgNPs has been greatly decreased; however, AgNPs still induce cell damage through induction of apoptosis.19-21 The cytotoxicity of AgNPs could be influenced by particle size, surface-stabilizing agents, pH, and some other factors.1,5,7,22-24 The surface-stabilizing coating is one of the most important factors for protecting cell viability from AgNPs.25-27

Previous studies have shown that chitosan has a very good biocompatibility and antibacterial properties. In addition, AgNPs/chitosan materials could promote wound healing3,17,28-30 In comparison with long-chain high molecular weight chitosan, the short-chain LMWC is better in terms of water solubility and biocompatibility. Besides, it has been confirmed that LMWC enhances the physical and chemical stability of many drugs such as insulin11 and siRNA,32 and works as a suitable carrier for the delivery and controlled release of drugs. In this study, we used hydrogen peroxide method to decompose high molecular weight chitosan and obtained short-chain molecules of chitosan with molecular weight between 2,000 and 2,500 Da. This short-chain molecule had a good solubility in water, thus avoiding the requirement of an acidic solution to dissolve high molecular weight chitosan. Compared to high molecular weight chitosan, this LMWC demonstrated a better biological activity (Figure 1C). The obtained LMWC was used as a stabilizer and a reducing agent for direct synthesis of LMWC-AgNPs with particle diameter size of 10–30 nm. In the current study, the three types of AgNPs were 10–30 nm with the same distribution but different coatings. All the three types of AgNPs caused no obvious cytotoxicity at lower concentrations (10 and 20 μg/mL). When the concentration increased to up to 30 μg/mL, however, treatment with LMWC-AgNPs led to significant less toxicity compared with the other two types of AgNP treatments, in the viability of human fibroblast HS27 cells (Figure 5A). This indicates that the surface coating with LMWC-AgNPs reduces cell toxicity and increases cell viability, although further study is needed for more detailed explanation. In comparison to PVP-AgNPs and uncoated-AgNPs, LMWC-AgNPs had similar effects of killing MRSA effects but were less toxic to human fibroblasts. Mice experiments also showed that the three types of AgNPs had good anti-inflammatory effects and effective protection of wound healing in mice.

Previous studies have shown that, through various means such as oral intake, intraperitoneal injection, and intravenous injection, AgNPs can be distributed and deposited on various organs of the body, with the deposition mainly in the liver and spleen.1,7,9,33,34 In addition, it has been reported that the ALT and AST levels in mice serum rose significantly after 2 days of oral intake of 20 μg/g AgNPs with a size of 35–45 nm.8 Intriguingly, in the current study, we found that after 11 days of application, the liver silver content in the mice treated with LMWC-AgNPs was significantly lower than that in the mice treated with uncoated-AgNPs or PVP-AgNPs. This suggests less LMWC-AgNP absorption into the wound and organs than with uncoated-AgNPs or PVP-AgNPs. Similarly, the silver content of the newly healed skin tissues of the mice treated with LMWC-AgNPs...
was significantly lower than that in the uncoated-AgNP and PVP-AgNP groups. Furthermore, both the ALT and AST levels in mice serum in the LMWC-AgNP-treated group were much less than those in the uncoated-AgNP-treated or PVP-AgNP-treated groups. Our results suggest that LMWC-AgNPs have the advantage of less body absorption while remain the anti-infection properties.

In summary, by application of an LMWC as a stabilizer and a reducing agent, we synthesized a diameter of 10–30 nm LMWC-AgNPs. We reveal for the first time that LMWC-AgNPs the characteristics of anti-MRSA infection, low cytotoxicity, better biocompatibility, and lower body absorption in both in vitro and in vivo experiments, when compared with PVP-AgNP and uncoated-AgNP treatments. Our study suggests that LMWC-AgNPs have less risk of damage to organ functions and are effective and safer AgNPs for medical materials, which is very important for large area burn patients, who need local usage of antibacterial materials on wounds for a long time after injury.

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Disclosure
The authors report no conflicts of interest in this work.

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