

# In vitro and in vivo evaluation of biologically synthesized silver nanoparticles for topical applications: effect of surface coating and loading into hydrogels

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**Abstract:** In the present study, silver nanoparticles (AgNPs) were synthesized via biological reduction of silver nitrate using extract of the fungus *Fusarium verticillioides* (green chemistry principle). The synthesized nanoparticles were spherical and homogenous in size. AgNPs were coated with polyethylene glycol (PEG) 6000, sodium dodecyl sulfate (SDS), and  $\beta$ -cyclodextrin ( $\beta$ -CD). The averaged diameters of AgNPs were  $19.2 \pm 3.6$ ,  $13 \pm 4$ ,  $14 \pm 4.4$ , and  $15.7 \pm 4.8$  nm, for PEG-, SDS-, and  $\beta$ -CD-coated and uncoated AgNPs, respectively. PEG-coated AgNPs showed greater stability as indicated by a decreased sedimentation rate of particles in their water dispersions. The antibacterial activities of different AgNPs dispersions were investigated against Gram-positive bacteria (methicillin-sensitive and methicillin-resistant *Staphylococcus aureus*) and Gram-negative bacteria (*Escherichia coli*) by determination of the minimum inhibitory concentrations (MICs) and minimum bactericidal concentrations (MBCs). MIC and MBC values were in the range of 0.93–7.5 and 3.75–15  $\mu\text{g/mL}$ , respectively, which were superior to the reported values in literature. AgNPs-loaded hydrogels were prepared from the coated-AgNPs dispersions using several gelling agents (sodium carboxymethyl cellulose [Na CMC], sodium alginate, hydroxypropylmethyl cellulose, Pluronic F-127, and chitosan). The prepared formulations were evaluated for their viscosity, spreadability, in vitro drug release, and antibacterial activity, and the combined effect of the type of surface coating and the polymers utilized to form the gel was studied. The in vivo wound-healing activity and antibacterial efficacy of Na CMC hydrogel loaded with PEG-coated AgNPs in comparison to the commercially available silver sulfadiazine cream (Dermazin<sup>®</sup>) were evaluated. Superior antibacterial activity and wound-healing capability, with normal skin appearance and hair growth, were demonstrated for the hydrogel formulations, as compared to the silver sulfadiazine cream. Histological examination of the treated skin was performed using light microscopy, whereas the location of AgNPs in the skin epidermal layers was visualized using transmission electron microscopy.

**Keywords:** silver nanoparticles, green synthesis, coating agents, hydrogel, wound healing, antibacterial activity

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## Introduction

Multidrug resistance is commonly developed against several pathogenic bacteria consequent to the use of several antibiotics.<sup>1-4</sup> Recovery from multidrug-resistant infections is a problematic issue<sup>5,6</sup> and requires multiple treatments with broad-spectrum antibiotics, which is usually associated with toxicity and high cost.<sup>6</sup> Methicillin-resistant *Staphylococcus aureus* (MRSA) is the main cause of worldwide nosocomial

infections and usually associated with increasing mortality and morbidity rates.<sup>7</sup> In general, *S. aureus* causes superficial lesions in human skin, localized abscesses, septicemia, pneumonia, and bacteremia.<sup>8</sup>

Historically, silver was one of the frequently used antibacterial agents before the introduction of antibiotics.<sup>9</sup> Silver compounds, silver ions, and silver nanoparticles (AgNPs) have been validated extensively for their antibacterial, antifungal, and antiviral activities.<sup>2,10–13</sup> Silver ions in silver-containing creams (such as silver sulfadiazine) are reported to exhibit high toxicity when applied to denuded skin due to the increased absorption of silver ions.<sup>2</sup> In addition, they have low stability due to inactivation by complexation with available counter ions and subsequent precipitation.<sup>3</sup> Among several tested metallic nanoparticles, AgNPs were found to be the most effective antimicrobial agent against bacteria, viruses, and other eukaryotic microorganisms.<sup>14</sup> AgNPs can be used as an effective broad-spectrum antibacterial agent against Gram-negative and Gram-positive bacteria including antibiotic-resistant bacteria.<sup>15</sup> Silver attacks multiple targets in the microorganisms, and therefore decreases their ability to develop resistance.<sup>13,16,17</sup> AgNPs exhibit higher toxicity to microorganisms than mammalian cells.<sup>18</sup> The broad-spectrum antibacterial effectiveness of AgNPs is due to their small size that enhances the accumulation in the microbial membrane and thus results in disruption and loss of integrity of the microbial membrane.<sup>8,19–21</sup>

AgNPs are fabricated by different methods including chemical, physical, and green methods.<sup>1,3,22,23</sup> Chemical and physical methods are usually expensive, with low production rates, and are not ecofriendly due to the toxicity of the used chemical-reducing agents and capping agents.<sup>4</sup> On the other hand, green chemistry method is preferred because the synthesized products are obtained from natural extracts.<sup>24</sup> Green chemistry involves different biological methods for the synthesis of AgNPs which employ plants extract or microorganisms, such as fungi and bacteria.<sup>1,20,24,25</sup>

To prevent aggregation of particles, different coating materials, such as carboxylic acids, polymers, polysaccharides, and surfactants, have been exploited to increase stability of the particles.<sup>26</sup> Polyethylene glycols (PEGs) are stable, hydrophilic, and nonirritant to the skin,<sup>27</sup> and have been used as an organic capping agent to stabilize AgNPs and control further growth of the particles via trapping of seeds and steric repulsion.<sup>28</sup>

Topical application of AgNPs for treatment of skin infections could be more effective if the particles are suspended in a hydrogel matrix, as compared to free particles suspended in water. Hydrogels are easily applied

and removed with improved patient compliance. Various gelling agents were reported in literature for the preparation of antimicrobial hydrogels due to their antibacterial activity per se, which might potentiate the efficacy of the loaded antimicrobial agents. Several authors reported the preparation of topically applied formulations intended for the delivery of AgNPs.<sup>21,24,29,30</sup>

The present work aimed at biological synthesis of AgNPs in a simple, ecofriendly, and cost-effective manner using fungus *Fusarium verticillioides* ASU1 (KT587649). A comparative detailed study based on the effect of three factors on AgNPs activity was provided. First, effect of different coating agents, including PEG 6000, sodium dodecyl sulfate (SDS), and  $\beta$ -cyclodextrin ( $\beta$ -CD), on AgNPs stability and antimicrobial activity was investigated. Second, various gelling agents, namely sodium carboxymethyl cellulose (Na CMC), sodium alginate (Na alginate), hydroxypropylmethyl cellulose (HPMC), Pluronic F-127, and chitosan, were used in the preparation of AgNPs-loaded hydrogels to aid in the topical application of AgNPs. The effect of coating type and gelling agent on the viscosity, spreadability, in vitro release profile, and antibacterial activity of the medicated hydrogels was studied. Third, effect of type of bacteria on the activity of AgNPs was evaluated. Gram-negative (*Escherichia coli*) and Gram-positive (MRSA and methicillin-sensitive *S. aureus* [MSSA]) bacteria were used to evaluate the antimicrobial activities of the various formulations. Efficacy of a selected 0.1% AgNPs-loaded hydrogel was studied in vivo versus the commercially available 1% silver sulfadiazine cream (Dermazin<sup>®</sup>), and wound-healing and antimicrobial activities in rats and in MRSA-infected mouse model were compared.

## Materials and methods

### Materials

Silver nitrate ( $\text{AgNO}_3$ ),  $\beta$ -CD, Pluronic F-127, and HPMC were purchased from Sigma Aldrich (St Louis, MO, USA). Na CMC of molecular weight 30,000 Da, propylene glycol (PG), and PEG 6000 were purchased from Adwic, EL-Nasr Pharmaceutical Chemicals (Cairo, Egypt). SDS was purchased from Novartis (Basel, Switzerland). Chitosan of molecular weight 100,000–300,000 Da was purchased from Acros Organics (Geel, Belgium). Na alginate was purchased from Judex Laboratory Reagent (Sudbury, UK).

### Biological synthesis of AgNPs

#### Microorganism and media

The strain *F. verticillioides* ASU1 (KT587649), isolated from *Zea mays* seeds, was used in this study. The strain was

maintained on potato dextrose agar medium at  $28^{\circ}\text{C}\pm 2^{\circ}\text{C}$  and stored at  $4^{\circ}\text{C}$  for subsequent study.

### Biosynthesis of AgNPs

For the synthesis of AgNPs, the biomass of fungus *F. verticillioides* ASU1 (KT587649) was prepared by growing the fungus in a potato dextrose broth medium. Flasks were inoculated, incubated on an orbital shaker at  $28^{\circ}\text{C}\pm 2^{\circ}\text{C}$ , and agitated at 150 rpm for 72 h. After complete incubation, the biomass was harvested by filtering through filter paper followed by repeated washing with distilled water to remove any medium components from the biomass. About 15 g (wet weight) was brought in contact with 100 mL of sterilized double-distilled water for 24 h at  $28^{\circ}\text{C}\pm 2^{\circ}\text{C}$  in a 250 mL Erlenmeyer flask and agitated again at 150 rpm. The cell filtrate was obtained after the incubation by passing it through Whatman® filter paper Number 1. The filtrate was treated with aqueous 1 mM  $\text{AgNO}_3$  solution in an Erlenmeyer flask and incubated at room temperature. Cell-free filtrate without  $\text{AgNO}_3$  solution was also run as a control.<sup>31</sup> AgNPs dispersion was then centrifuged at 14,000 rpm for 20 min. The supernatant was discarded, and the pellets were redispersed in distilled water, and washed three times with sterilized distilled water to remove the free entities and unbound biological molecules from AgNPs.

## Characterization of the synthesized AgNPs

### Atomic absorption spectroscopy (AAS)

The concentration of AgNPs in the suspensions was measured using the Buck Scientific atomic absorption spectrophotometer (Model 210 VGP; Norwalk, CT, USA) equipped with a silver hollow cathode lamp and air-acetylene flame. Standard silver solutions were also prepared and measured.

### X-ray diffraction (XRD)

Crystalline structure of the biologically synthesized AgNPs powder was identified using a Philips X-ray diffractometer model PW 1710 (Philips, Amsterdam, the Netherlands) with  $\text{CuK}_{\alpha}$  radiation ( $\lambda=1.5405 \text{ \AA}$ ), at an operating applied voltage of 40 kV and 30 mA current, equipped with a graphite monochromator and automatic divergence slit. Measurements were swapped from  $2\theta=35^{\circ}$  to  $2\theta=80^{\circ}$  with a scanning speed of  $0.06^{\circ}/\text{min}$ .

### UV-visible spectrophotometry

The bioreduction of  $\text{AgNO}_3$  solution and formation of AgNPs by fungal extract was confirmed by scanning the resulted dispersion at 300–900 nm wavelength range using

a double-beam spectrophotometer (Thermo Scientific Evolution™ 300 UV-Vis Spectrophotometer; Thermo Fisher Scientific, Waltham, MA, USA). A strong absorption of electromagnetic waves was exhibited by metal nanoparticles in the visible range due to the surface plasmon resonance (SPR) effect.

### Transmission electron microscopy (TEM)

A drop of the biologically synthesized AgNPs dispersion ( $100 \mu\text{g}/\text{mL}$ ) was placed on a carbon-coated copper grid and dried at room temperature. TEM micrograph of the sample was taken using the JEOL TEM (Model 100 CX II; Tokyo, Japan) at the Electron Microscopy Unit, Assiut University, Egypt. Size and shape of AgNPs were studied.

## Coating of AgNPs

Effect of coating agents on AgNPs sedimentation rate and antibacterial activity was studied using different coating agents, namely PEG 6000 (hydrophilic polymer), SDS (anionic surfactant), and  $\beta$ -CD (cyclic oligosaccharide polymer), which were added separately after the biological synthesis of AgNPs. PEG, SDS, and  $\beta$ -CD solutions were prepared by dissolving their powders separately in distilled water. Then, the prepared solutions were added directly to the freshly prepared AgNPs suspension ( $500 \mu\text{g AgNPs}/\text{mL}$ ) to obtain final concentrations of 5 and 10 mM of each coating agent. The final solutions were sonicated for 1 h using an ultrasonicator (Crest Ultrasonics Corp., Trenton, NJ, USA).<sup>28,32,33</sup> The ability of different coating agents, prepared at different concentrations, to maintain the nanoparticles dispersed in solutions and prevent their sedimentation over time was studied. Stability of AgNPs dispersions ( $500 \mu\text{g}/\text{mL}$ ) was examined visually after 1, 2, 3, 6, 9, and 12 months at  $4^{\circ}\text{C}$ . Also, the coated AgNPs dispersions were scanned at 300–900 nm using a UV-visible spectrophotometer, and TEM micrographs were also taken as previously described.

### Dynamic light scattering (DLS) and zeta-potential measurements

The mean hydrodynamic diameters and size distribution (polydispersity indices, PDIs) of the nanoparticles in water as a dispersant were measured using a Zetasizer Nano ZS instrument (Malvern Instruments, Malvern, UK) equipped with a backscattered light detector operating at  $173^{\circ}$ . The CONTIN program was used to extract size distributions from the autocorrelation functions. The zeta-potential values were determined by laser Doppler anemometry using a Malvern Zetasizer Nanoseries ZS.

## Antibacterial activity assay

### Bacterial strains and growth condition

The following microbial strains were used: MSSA and MRSA, which are Gram-positive bacteria; and *E. coli*, which are Gram-negative bacteria. All bacterial strains were clinically isolated at the infection control unit (Assiut University Hospital). Strains were grown in Mueller–Hinton broth or Mueller–Hinton agar medium at 37°C. A single isolated colony of each bacterium was picked from the agar plate culture and suspended in 0.9% (w/v) sterile aqueous saline solution to prepare bacterial suspensions of 0.5 McFarland standard and diluted to give a final concentration of 10<sup>8</sup> colony-forming units (CFU)/mL.

### Minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC)

In order to investigate the antibacterial activity of AgNPs and the role of coating agents, MIC of AgNPs dispersions against clinically isolated bacteria was determined using microtiter broth dilution method according to the guidelines of the Clinical and Laboratory Standards Institute.<sup>34</sup> Sterile 96-well rounded-bottom plates were used. AgNPs dispersions were sonicated before the experiment to ensure homogenous distribution of the particles. Mueller–Hinton broth (100 µL) was dispensed into all wells of the microtiter plate. Each tested dispersion was added to the first well, and serial twofold dilutions were made. Bacterial dispersions (5 µL of 10<sup>6</sup> CFU/mL) were added into all wells except for the sterility control blank. Plates were incubated at 37°C for 24 h and evaluated for the MIC by visual examination of the culture turbidity. In order to determine the MBC of the tested AgNPs dispersions, 5 µL from the dilution that showed the MIC (no visible bacterial growth) and four of the more concentrated dilutions were plated, cultured at 37°C for 24 h without AgNPs supplementation, and counted to determine viable CFU/mL. The MBC is the lowest concentration that showed the same predetermined reduction in CFU/mL when compared to the MIC dilution.

### Agar-well diffusion method

Another method was also used to evaluate the antibacterial activity of the AgNPs. Bacterial suspensions (100 µL) were mixed with Sabouraud agar (20 mL) in sterile petri dishes (9 cm in diameter), and the agar plates were allowed to solidify. After solidification, wells were made in the agar plates using a pore-maker of size 10 mm and filled with the tested AgNPs dispersions (200 µL of 240 µg AgNPs/mL) after sonication to ensure complete dispersion of the particles. Pure solutions of PEG, SDS, and β-CD were also used as positive growth controls. Plates were incubated at 37°C for 24 h, and the diameters

of inhibition zone were measured without subtracting the well diameter from the total inhibition zone diameter using a digital caliber.<sup>35,36</sup> Inhibition zone diameter is an indicator of the antibacterial activity of AgNPs dispersions.

## Formulation of AgNPs-loaded hydrogels

In order to make AgNPs dispersions applicable as topical treatment and to improve patient compliance, hydrogel formulations were prepared using five different polymer-gelling agents. The composition of the formulated hydrogels containing 500 µg AgNPs per gram of hydrogel is shown in Table 1. As illustrated in the table, 25 formulations (F1–F25) were prepared by incorporating five different AgNPs dispersions (uncoated and 5 mM PEG-, 10 mM PEG-, 5 mM SDS-, and 5 mM β-CD-coated AgNPs) into each of five gelling polymers (Na CMC, Na alginate, HPMC, Pluronic F-127, and chitosan). Based on previous optimizations, PG (20%, w/w) was added to all the prepared hydrogels to improve consistency and decrease the viscosity of the hydrogel formulations. In addition, PG enhances the hydrogen bonding between water and gelling agents, thus improving the AgNPs dispersion stability in the polymer network.<sup>37,38</sup>

The specified amounts (3%, w/w) of the gelling polymers (Na CMC, Na alginate, and HPMC) were added slowly, each separately, to a previously formed mixture of 20% (w/w) PG and specific volumes of AgNPs stock dispersions. Then, the mixtures were stirred continuously using a magnetic stirrer (Gallenkamp, Loughborough, UK) and allowed to soak overnight for complete polymer solvation. Twenty-five percent (w/w) Pluronic F-127 gels were prepared by the cold method described by Schmolka.<sup>39</sup> The required amount of Pluronic F-127 was dissolved in PG and cold AgNP dispersions. Dispersions were left in a refrigerator overnight. A clear transparent gel was obtained when the dispersion was maintained at room temperature. Three percent (w/w) chitosan gel was prepared by addition of chitosan and 1% (v/v) glacial acetic acid to specific amount of AgNPs dispersion and 20% PG using a magnetic stirrer. The pH of chitosan hydrogel was adjusted to 5.5 by addition of NaOH solution.

## Evaluation of the prepared hydrogel formulations

### Physical appearance

The prepared formulations were inspected visually for their color and homogeneity.

### Viscosity and rheological behavior

The flow rheological properties of the prepared topical formulations (F1–F25) were studied for blank and medicated

**Table 1** Composition of AgNPs hydrogels (500 µg AgNPs/g hydrogel). AgNPs were coated with 5 mM PEG 6000, 10 mM PEG 6000, 5 mM SDS, and 5 mM β-CD, and AgNPs-loaded hydrogels were prepared using various polymers (Na CMC, Na alginate, HPMC, Pluronic F-127, and chitosan)

Formulation	Composition	Coating agent type					PG	
		Gelling agent type	Uncoated AgNPs	5 mM PEG 6000-coated AgNPs	10 mM PEG 6000-coated AgNPs	5 mM SDS-coated AgNPs		5 mM β-CD-coated AgNPs
			✓	✓	✓	✓		✓
F1	3% Na CMC	✓					20%	
F2			✓					
F3				✓				
F4					✓			
F5						✓		
F6	3% Na alginate	✓						
F7			✓					
F8				✓				
F9					✓			
F10						✓		
F11	3% HPMC	✓						
F12			✓					
F13				✓				
F14					✓			
F15						✓		
F16	25% Pluronic F-127	✓						
F17			✓					
F18				✓				
F19					✓			
F20						✓		
F21	3% Chitosan	✓						
F22			✓					
F23				✓				
F24					✓			
F25						✓		

**Abbreviations:** AgNPs, silver nanoparticles; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate; β-CD, β-cyclodextrin; Na CMC, sodium carboxymethyl cellulose; Na alginate, sodium alginate; HPMC, hydroxypropylmethyl cellulose; PG, propylene glycol.

formulations (20 g) at 25°C using a Brookfield DV-III ULTRA programmable rheometer, model RV, T-bar spindle numbers 95 (Brookfield Engineering Laboratories, Inc., Middleboro, MA, USA). Effect of AgNPs incorporation on the rheological characteristics of the hydrogel formulations was studied. Also, the effect of coating agents and gelling agents on viscosity of the hydrogel formulations was investigated. Measurements were performed over the range of speed settings from 5 to 50 rpm with 1 min interval between each two successive speeds to generate a complete flow curve. The rheograms were generated by plotting viscosity readings in centipoises versus spindle speed (rpm).

### Spreadability

Spreadability of the prepared formulations (F1–F25) was determined as reported previously.<sup>40</sup> Each formulation (0.25 g) was moderately pressed between two glass slides and left for 5 min. Diameters of the spreaded sample circles

were measured in centimeter and taken as comparative values for spreadability. Results obtained were the average of three measurements.

### In vitro release

The in vitro release of AgNPs from the prepared hydrogel formulations (F1–F25) was studied. One-gram sample of each formulation was accurately weighed and placed on nylon fabric bag as a donor compartment (44±15 µm pore size). The bag was immersed into a beaker (receptor compartment) containing 10 mL of double-distilled water as the release medium (pH 6). The system was maintained for 3 h at 37°C±0.5°C in a thermostatically controlled shaking water bath (Gallenkamp) at 50 rpm. Aliquots of 2 mL were withdrawn at intervals of 0.5, 1, 1.5, 2, and 3 h, which were replaced by the same volume of fresh water (kept at the same temperature) to maintain constant volume. Samples were analyzed for AgNPs content spectrophotometrically at

$\lambda_{\max}$  420 nm against a blank sample that was similarly treated. Results were compared to the AgNPs dispersion of same concentration. All experiments were carried out in triplicates, and the average values were calculated. Kinetics of the AgNPs release data from the prepared hydrogels was analyzed, and details are provided in the [Supplementary materials](#).

### Antibacterial activity using agar-well diffusion method

In order to evaluate the release of the loaded AgNPs from the hydrogels (F1–F25) in the agar plates and their antibacterial activity, the prepared medicated formulations (200  $\mu\text{g}$  of 500  $\mu\text{g}$  AgNPs/g) were loaded inside agar wells. Plates were incubated at 37°C for 24 h, and then were examined for the diameters of the bacterial growth inhibition zone as previously described. Blank hydrogel formulations were also tested as positive control.

### In vivo studies

Based on the previous in vitro studies and characterizations, 0.1% 10 mM PEG-coated-AgNPs-loaded Na CMC hydrogel (F3) was selected for the in vivo studies. The in vivo studies involved the evaluation of wound-healing-promoting activity of AgNPs-loaded hydrogel as described previously.<sup>30</sup> Then, in vivo antibacterial activity was evaluated,<sup>41</sup> and the protocols were approved by the Assiut University Animal Ethical Committee. Ethical and legal approvals were obtained prior to the commencement of the study and all the animal experiments were approved by the Institutional Animal Ethical Committee of Faculty of Pharmacy, Assiut University, and it adheres to the Guide for the Care and Use of Laboratory Animals, 8th Edition, National Academies Press, Washington, DC, USA.

### In vivo wound-healing efficacy

The wound-healing activity of the prepared 0.1% 10 mM PEG-coated-AgNPs-loaded Na CMC hydrogel (selected formulation, F3) was investigated. Ten adult female rats aged 8–10 weeks and weighing 200–250 g were used. Animals were purchased from the Animal Facility of the Faculty of Medicine (Assiut University, Egypt). Rats were divided randomly into two groups ( $n=5$ ). Group I served as test group which received 0.1% AgNPs hydrogel (F3), and group II served as a control and received the commercially available 1% silver sulfadiazine cream. Rats were kept in large cages and allowed food and water ad libitum. Rats were anesthetized using intraperitoneal (IP) thiopental injections, and were then shaved on their dorsal surfaces. A circular piece of full-thickness skin ( $1 \times 1 \text{ cm}^2$ )

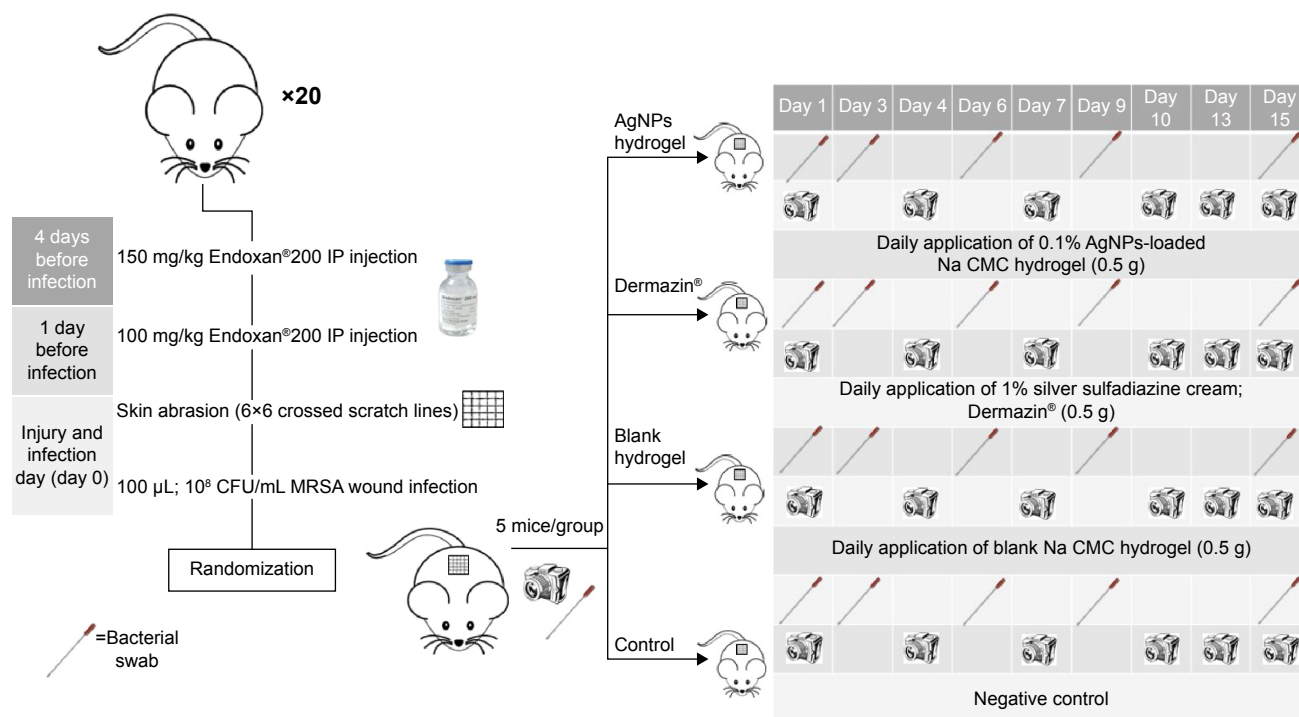
from each rat was excised using scissors, and wound was left undressed to the open environment.<sup>42</sup> Treatment of both groups was started at the day of wound creation and lasted for 10 d. Rate of healing in both groups, represented by percent wound contraction, was assessed and compared. Percent wound contraction was determined according to the following equation:

$$\% \text{ wound contraction} = \frac{\text{Wound area day 0} - \text{Wound area day } n}{\text{Wound area day 0}} \times 100$$

### In vivo antibacterial activity

The antibacterial activity of 0.1% 10 mM PEG-coated-AgNPs-loaded Na CMC hydrogel (selected formulation, F3) was evaluated using a superficial skin infection model<sup>41</sup> using MRSA strain in animals and compared with that of 1% silver sulfadiazine cream, hydrogel blank formulation (3% Na CMC blank hydrogel), and a group of animals with no treatment as a control group.

Twenty adult female mice were used in this study. All mice were 6–8 weeks old and weighing 20–25 g. Mice were purchased from the Animal Facility of the Faculty of Medicine (Assiut University, Egypt). Experimental study design is presented in Figure 1. Mice were housed in large cages (five animals per cage) and allowed food and water ad libitum. Four days and 1 d prior to the infection, mice were injected intraperitoneally with two doses of Endoxan<sup>®</sup>200 (cyclophosphamide). The first dose of cyclophosphamide was 150 mg/kg followed by a second dose of 100 mg/kg. Cyclophosphamide was reported to reduce the mice peripheral blood neutrophils to  $<100/\mu\text{L}$  blood, thus providing a susceptible environment for infection.<sup>41</sup> Mice were anesthetized with IP thiopental injections, and then shaved on their dorsal surfaces. Skin wounds were created using the abrasion method which was applied on the dorsal surfaces of mice using 24-gauge needles. Crossed scratch lines ( $6 \times 6$ ) were created in a  $1 \text{ cm}^2$  area with damaged stratum corneum and upper layer of the epidermis, without affecting the dermis. Five minutes after wounding, an aliquot of MRSA suspension (100  $\mu\text{L}$  of  $10^8$  CFU/mL in phosphate-buffered saline) was inoculated over each defined area containing the crossed scratches with a micropipette tip. Animals were divided into four groups each containing five animals and received the specified treatment 1 d postinfection according to each group (0.5 g/d for 15 d). Group I was treated with the prepared 0.1% AgNPs gel formulation (F3), group II was treated with the commercially available 1% silver sulfadiazine cream, group III was treated with the blank hydrogel formulation (hydrogel



**Figure 1** Graphical chart represents the design of the study on the in vivo antibacterial activity of AgNPs.

**Abbreviations:** AgNPs, silver nanoparticles; IP, intraperitoneal; CFU, colony-forming units; MRSA, methicillin-resistant *Staphylococcus aureus*; Na CMC, sodium carboxymethyl cellulose.

without AgNPs), and group IV was the control (without any application).

Swabs from the surface of the crossed scratches were taken using a sterilized cotton swab at 1 d (just before treatment), and at 3, 6, 9, and 15 d postinfection. MRSA (CFU/mL) were counted after serial dilution of the bacterial swab and cultured on a selective solid agar media (mannitol salt agar), and then incubated for 24 h at 37°C.

### Histopathological examinations

#### Light microscope examination using hematoxylin and eosin stain

To assess morphological alterations during the in vivo antibacterial activity study, the tissue samples from the mice back skin were harvested after euthanizing the mice and were processed routinely for light microscopic examination. Skin sections were evaluated for intercellular and intracellular epidermal edema, dermal edema, and inflammation. For the histological examinations by light microscope, small fresh pieces from the skin were taken from the four groups, fixed in 10% of neutral buffered formalin (pH 7.2), dehydrated in graded series of ethanol, cleared in methyl benzoate, and then embedded in paraffin wax. Thin paraffin sections (5 µm) were prepared and stained with Harris's hematoxylin and eosin.

### Transition electron microscope examination

To visualize the AgNPs diffusion from the hydrogel into the epidermal layers of the skin surface, TEM was used. Selected skin tissues were fixed in 2% glutaraldehyde followed by osmium tetroxide and then dehydrated in ascending concentrations of ethanol. Samples were embedded in Epon 812 according to the protocol of the Electron Microscopy Unit (Assiut University, Egypt). Semi-thin (1 µm) sections were cut on an Ultracut E ultramicrotome using a diamond knife, stained with toluidine blue, and examined by light microscope. Finally, ultrathin sections (60 nm) were cut with an ultramicrotome (Reichert Ultracuts), contrasted with an uranyl acetate and lead citrate, and examined with a transmission electron microscope (JEOL TEM 100 CXII) at 80 kV and imaged.<sup>43</sup>

### Statistical analysis

Statistical analysis was performed using SPSS statistical package.<sup>44</sup> Basic statistical parameters (mean, standard deviation, or standard error) of the presented data were estimated. According to the number of the studied factors, results were subjected to one-, two-, and three-way analysis of variance (ANOVA) to study the simple and main effects between levels of the different factors (coating agents, gelling agents, and/or bacterial type). Test of homogeneity was carried out.

Homogeneity of variance was evident in most cases and assumed in other cases. Tukey's post hoc test was carried out for each factor, and the homogenous subsets were identified at 0.05 level of significance.

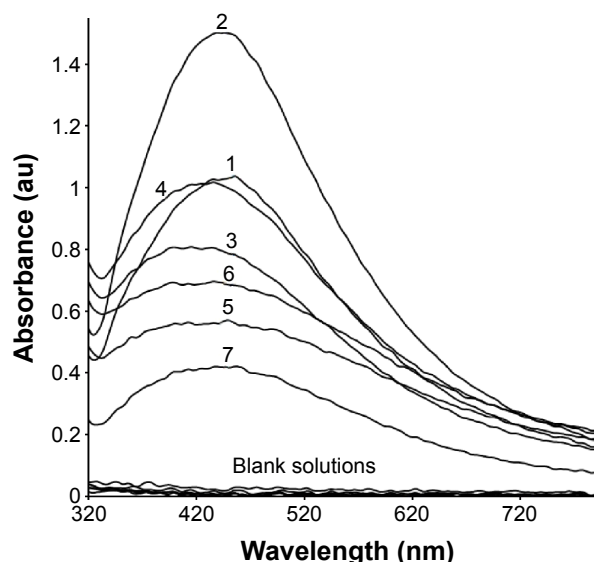
## Results and discussion

### Biosynthesis and characterization of AgNPs

AgNPs were produced using a simple, fast, clean, benign, and ecofriendly alternative method to the complex chemical synthetic methods, where the crude fungal extract acts as reducing agent.<sup>45</sup> Several groups have utilized the crude extract of fungi for myco-synthesis of AgNPs.<sup>31,46,47</sup> Different studies have discussed the fabrication of AgNPs based on fungi-assisted bioreduction processes, as well as their antibacterial activities.<sup>20,48,49</sup> The fungus *Fusarium* sp. can reduce aqueous silver ions extracellularly to generate AgNPs.<sup>46</sup> The process of silver ions reduction occurs through the action of both reductase enzymes and electron shuttle quinones.<sup>47</sup> In addition, biological nanoparticle synthesis often yields a more consistent size distribution pattern, as compared to other methods. In order to ensure the reproducibility of AgNPs yield and characteristics, synthetic conditions, such as duration, pH, temperature, and AgNO<sub>3</sub> starting concentration, were optimized. The optimal conditions for maximum AgNPs production were pH 7, 1 mM concentration of AgNO<sub>3</sub>, and 24 h incubation time.

The production of AgNPs in the solution was confirmed by visual detection of the color transformation from transparent to pale brown. The concentration of the synthesized AgNPs stock dispersion was measured using the AAS. XRD pattern of AgNPs synthesized using the fungus *F. verticillioides* extract is shown in Figure S1. The sharp peaks in the pattern indicate the existence of the silver and the crystalline nature of the biologically synthesized AgNPs. The four distinguished reflections at 38.1°, 44.3°, 64.5°, and 77.5° in the XRD pattern are indexed to the (111), (200), (220), and (311) crystallographic planes of the face-centered cubic structure of silver, respectively (JCPDS file no 00-004-0783).

AgNPs were coated with PEG 6000, SDS, and β-CD to improve the stability of the nanoparticles and reduce the growth of the particles via trapping of seeds and steric repulsion. Silver particles were characterized using different analytical techniques. UV-visible absorption spectra of the prepared AgNPs dispersions (before and after coating) were recorded in the range of 300–800 nm using a double-beam UV-visible spectrophotometer (Figure 2). Spectra of AgNPs dispersions exhibited a single SPR absorption band with a maximum of ca.

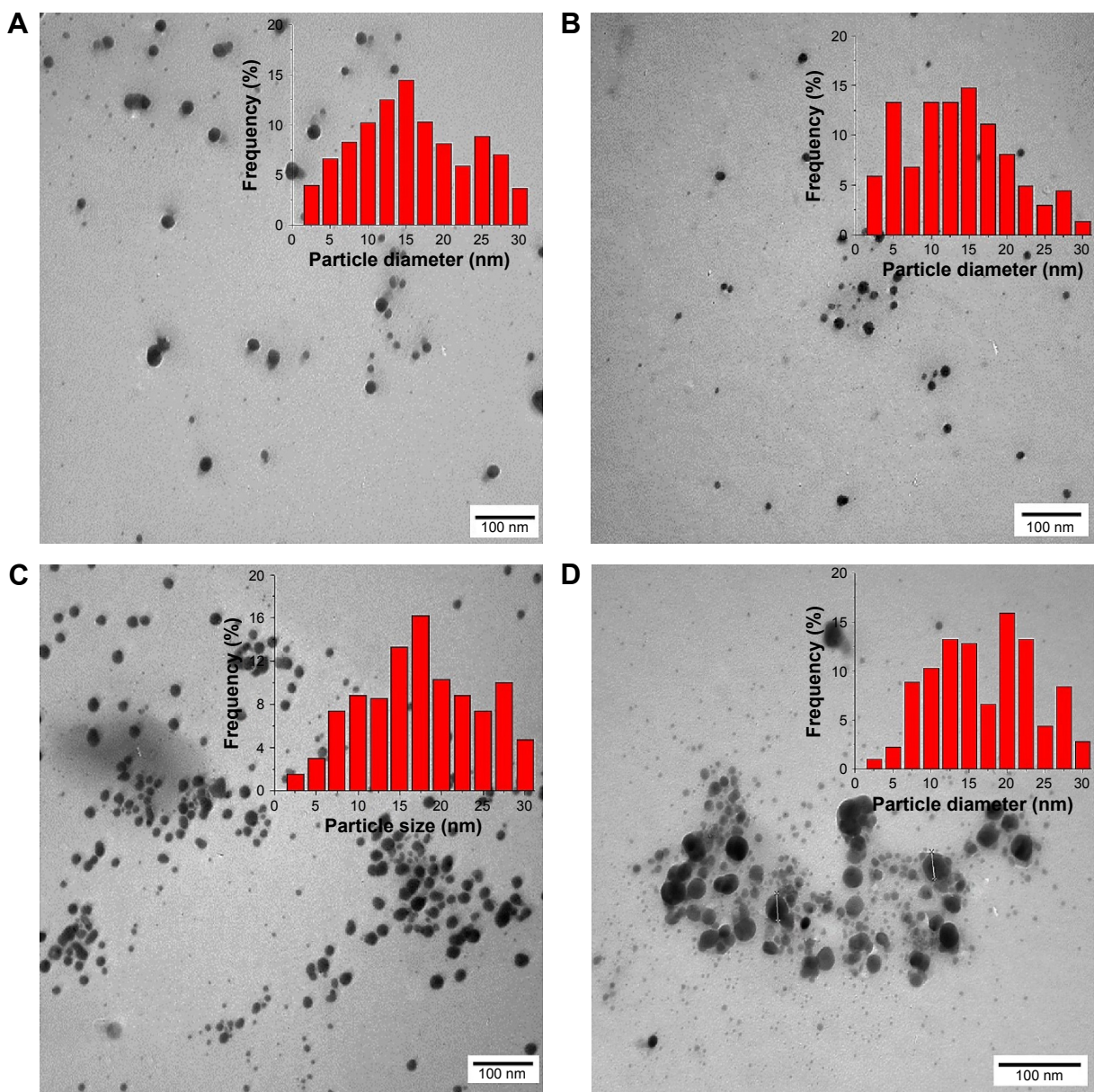


**Figure 2** UV-visible spectrophotometric spectra of AgNPs dispersions: (1) 5 mM PEG-coated AgNPs; (2) 10 mM PEG-coated AgNPs; (3) 5 mM SDS-coated AgNPs; (4) 10 mM SDS-coated AgNPs; (5) 5 mM β-CD-coated AgNPs; (6) 10 mM β-CD-coated AgNPs; and (7) uncoated AgNPs and their solutions without AgNPs (blank solutions).

**Abbreviations:** AgNPs, silver nanoparticles; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate; β-CD, β-cyclodextrin; UV, ultraviolet.

420 nm that indicated the presence of spherical nanoparticles.<sup>16</sup> Pal et al<sup>16</sup> reported that the number of SPR peaks decreases as the symmetry of the nanoparticles increases, and they found that spherical AgNPs show only one peak, whereas disks and triangular nanoplates show two or more peaks. Coating of AgNPs using different materials led to an increase in the peak intensity. Furthermore, this peak intensity increase was directly proportional to the polymer concentration. PEG-coated AgNPs dispersions of 5 and 10 mM concentrations showed a slight red shift to 450 nm. SPR of metal nanoparticles absorption spectra is usually shifted to longer wavelengths when particle size increases.<sup>16</sup> Size of AgNPs was determined using DLS and TEM imaging. The morphology of the particles was studied by TEM. Representative TEM micrographs of 5 mM PEG-, SDS-, and β-CD-coated and uncoated AgNPs are shown in Figure 3. Most of the nanoparticles were homogenous and spherical in shape, and the average particle size was 19.2±3.6, 13±4, 14±4.4, and 15.7±4.8 nm for 5 mM PEG-, SDS-, and β-CD-coated and uncoated AgNPs, respectively (n=50). Histograms (right-hand insets in the figures) indicate the AgNPs size distribution. There were no significant differences ( $P>0.05$ ) between the sizes of AgNPs measured using TEM. Therefore, DLS technique was exploited to measure the hydrodynamic size in water dispersions. Particle size measured using DLS was bigger than that obtained from TEM micrographs (Table 2) because DLS measures the hydrodynamic diameter of the particles.<sup>50</sup> In addition,





**Figure 3** Representative TEM micrographs for the aqueous dried AgNPs (100  $\mu$ g AgNPs/mL): (A) uncoated AgNPs; (B) SDS-coated AgNPs; (C) PEG-coated AgNPs ( $\times 100,000$ ); (D)  $\beta$ -CD-coated AgNPs ( $\times 140,000$ ) with sizes =  $15.7 \pm 4.8$ ,  $13 \pm 4$ ,  $19.2 \pm 3.6$ , and  $14 \pm 4.4$  nm, respectively ( $n=50$ , bar represents 100 nm). Insets indicate histograms of AgNPs size distribution.

**Abbreviations:** TEM, transmission electron microscopy; AgNPs, silver nanoparticles; SDS, sodium dodecyl sulfate; PEG, polyethylene glycol;  $\beta$ -CD,  $\beta$ -cyclodextrin.

**Table 2** Characterizations of the AgNPs dispersions in terms of size, PDI, and zeta-potential. Size was measured by using DLS and TEM

AgNPs dispersion	DLS: average particle size $\pm$ SD (nm)	DLS: average PDI $\pm$ SD	TEM: average particles size $\pm$ SD (nm)	Zeta-potential $\pm$ SD (mV)
5 mM PEG-coated AgNPs	$54.2 \pm 8.1$	$0.3 \pm 0.0$	$19.2 \pm 3.6$	$15.5 \pm 0.9$
10 mM PEG-coated AgNPs	$77.7 \pm 11.0$	$0.3 \pm 0.0$	–	$11.9 \pm 0.6$
5 mM SDS-coated AgNPs	$43.3 \pm 4.4$	$0.5 \pm 0.1$	$13.0 \pm 4.0$	$25.3 \pm 1.3$
10 mM SDS-coated AgNPs	$57.2 \pm 10.0$	$0.4 \pm 0.1$	–	$28.2 \pm 0.9$
5 mM $\beta$ -CD-coated AgNPs	$86.4 \pm 3.2$	$0.4 \pm 0.0$	$14.0 \pm 4.4$	$17.8 \pm 1.1$
10 mM $\beta$ -CD-coated AgNPs	$86.3 \pm 12.6$	$0.3 \pm 0.0$	–	$15.2 \pm 0.7$
Uncoated AgNPs	$48.5 \pm 14.3$	$0.3 \pm 0.0$	$15.7 \pm 4.8$	$20.1 \pm 1.5$

**Note:** <sup>a-c</sup>Refer to significant differences between different groups at the same column.

**Abbreviations:** AgNPs, silver nanoparticles; PDI, polydispersity index; DLS, dynamic light scattering; TEM, transmission electron microscopy; SD, standard deviation; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate;  $\beta$ -CD,  $\beta$ -cyclodextrin.

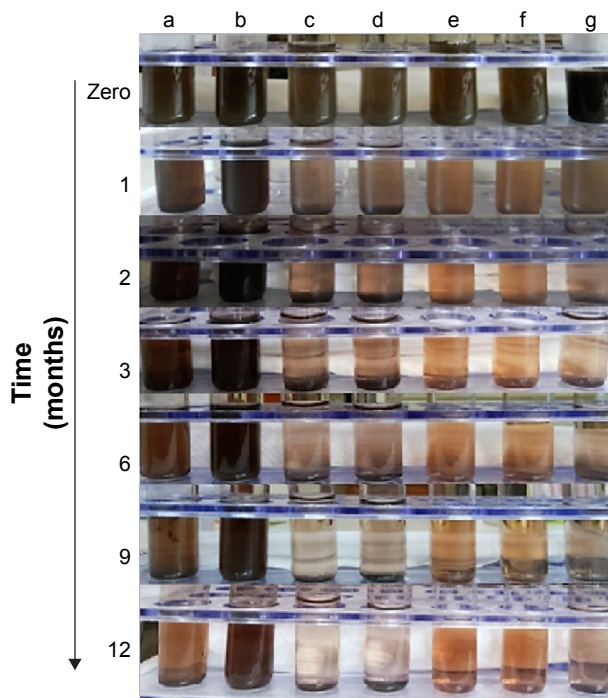
Ma et al<sup>51</sup> have reported that the increased size obtained from DLS over TEM is due to the coating effect that decreased the particle diffusivity, thus resulting in a larger hydrodynamic size. DLS measurements of 5 and 10 mM SDS-coated AgNPs and 5 mM PEG-coated AgNPs were not significantly different ( $P>0.05$ ) from that of uncoated AgNPs; however, increasing the concentration of PEG from 5 to 10 mM increased the hydrodynamic size significantly from 54.2 to 77.7 nm. PDIs of all the AgNPs dispersions were  $<0.5$ , thus indicating a narrow and homogenous particle size distribution (Table 2).<sup>52,53</sup> The surface charges of the synthesized AgNPs were determined by zeta-potential measurements (Table 2). Uncoated AgNPs showed a negative zeta-potential of  $-20.1$  mV which was decreased upon coating with 5 and 10 mM of neutral PEG to  $-15.5$  and  $-11.9$  mV, respectively. Coating of the AgNPs with 5 and 10 mM of the anionic surfactant SDS increased the negative charge significantly ( $P<0.05$ ) to  $-25.3$  and  $-28.2$  mV, respectively. Sundararajan et al<sup>54</sup> have reported that particles that possess zeta-potential values  $>+30$  and  $<-30$  mV are relatively stable. In the current study, AgNPs were stabilized by both electrostatic and steric repulsions.

### Effect of coating agent on the sedimentation rate of AgNPs in water

The rates of sedimentation of the prepared dispersions of AgNPs were evaluated visually at  $4^{\circ}\text{C}$ , and imaged at different time intervals for 1 year, and are shown in Figure 4. It is evident that 10 mM PEG-coated AgNPs formed a stable dispersion without any particles aggregation and sedimentation for ca. 9 months, whereas lower concentration of the same polymer (5 mM) afforded stability for ca. 3 months. On the contrary, sedimentation of uncoated, 5 and 10 mM SDS-coated, and 5 and 10 mM  $\beta$ -CD-coated AgNPs from their colloidal dispersions was significant after the first month. Coating layers are reported to increase the stability of the particles by enhancing the electrostatic (as in the case of anionic SDS) or steric (as in the case of neutral PEG and  $\beta$ -CD) repulsion.<sup>55</sup> AgNPs stabilized with steric repulsion (ie, PEG) are usually more stable than those stabilized via electrostatic repulsion (ie, SDS).<sup>55</sup> Stiufluc et al<sup>56</sup> have demonstrated that coating AgNPs with PEG increased not only the steric distance between nanoparticles but also the hydrophilicity via hydrogen bonds formation with the solvent, thus preventing particles aggregation.

### MIC and MBC

Table 3 summarizes the MIC and MBC values of the different AgNPs dispersions against both Gram-positive



**Figure 4** Effect of coating type on AgNPs ( $500 \mu\text{g AgNPs/mL}$ ) sedimentation rate in water over time: (a) 5 mM PEG-coated AgNPs; (b) 10 mM PEG-coated AgNPs; (c) 5 mM SDS-coated AgNPs; (d) 10 mM SDS-coated AgNPs; (e) 5 mM  $\beta$ -CD-coated AgNPs; (f) 10 mM  $\beta$ -CD-coated AgNPs; and (g) uncoated AgNPs.

**Abbreviations:** AgNPs, silver nanoparticles; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate;  $\beta$ -CD,  $\beta$ -cyclodextrin.

(MRSA and MSSA) and Gram-negative bacteria (*E. coli*). In the current study, MIC and MBC values of the studied AgNPs dispersions against MSSA, MRSA, and *E. coli* bacteria were in the range of  $0.93$ – $7.5$  and  $3.7$ – $15 \mu\text{g/mL}$ , respectively. The high antibacterial activity of the nanoparticles could be due to their small size (in the range of  $13\pm 4$  to  $19.2\pm 3.6$  nm), and subsequently large surface area of the particles. Several authors have reported that the antibacterial activity of AgNPs increases upon reduction of their particles size due to the increase in the surface area which allows greater contact with bacterial cells.<sup>57–59</sup> Ansari et al<sup>57</sup> studied the antibacterial activity of AgNPs ( $5$ – $10$  nm) against clinically isolated MSSA and MRSA, and reported that their MIC and MBC were in the range of  $12.5$ – $50$  and  $12.5$ – $100 \mu\text{g/mL}$ , respectively. In the present work, the antibacterial activity of AgNPs dispersions was higher, which might be due to the difference in AgNPs synthetic method and the sources of the bacterial strains utilized in the two studies.

MIC and MBC values were subjected to two-way ANOVA without replicates to investigate the effects of coating agent and bacterial type in the study (Table 3). PEG, SDS, and  $\beta$ -CD were studied previously as coating agents to

**Table 3** MIC and MBC of AgNPs dispersions against MSSA, MRSA, and *Escherichia coli* determined using microtiter plate method. Statistical data indicate the effect of bacterial type on the MIC/MBC of the prepared AgNPs, either uncoated or coated with various materials, regardless of the coating type

Dispersion (240 µg AgNPs/mL)	MIC (µg/mL)			MBC (µg/mL)		
	MSSA	MRSA	<i>E. coli</i>	MSSA	MRSA	<i>E. coli</i>
Dispersion (240 µg AgNPs/mL)						
5 mM PEG-coated AgNPs	1.87	3.75	7.50	7.50	7.50	15.00
10 mM PEG-coated AgNPs	1.87	3.75	7.50	7.50	7.50	15.00
5 mM SDS-coated AgNPs	0.93	1.87	7.50	3.75	3.75	7.50
10 mM SDS-coated AgNPs	1.87	3.75	3.75	7.50	7.50	3.75
5 mM β-CD-coated AgNPs	1.87	3.75	7.50	3.75	3.75	7.50
10 mM β-CD-coated AgNPs	1.87	3.75	7.50	7.50	3.75	7.50
Uncoated AgNPs	1.87	3.75	7.50	7.50	7.50	15.00
Main effects of bacteria ± SD	1.70 <sup>a</sup> ±0.35	3.80 <sup>b</sup> ±0.35	6.90 <sup>c</sup> ±0.35	6.40 <sup>a</sup> ±0.93	5.80 <sup>b</sup> ±0.93	10.20 <sup>b</sup> ±0.94

**Note:** <sup>a-c</sup>Refer to significant differences between different groups.

**Abbreviations:** MIC, minimum inhibitory concentration; MBC, minimum bactericidal concentration; AgNPs, silver nanoparticles; MSSA, methicillin-sensitive *Staphylococcus aureus*; MRSA, methicillin-resistant *S. aureus*; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate; β-CD, β-cyclodextrin; SD, standard deviation.

enhance the antibacterial activity of AgNPs, as compared to the naked nanoparticles.<sup>10,19</sup> In the present work, the effect of two different concentrations (5 and 10 mM) of PEG, SDS, and β-CD as coating agents on the antibacterial activity of AgNPs was studied. Neither the change in the type of coating nor the concentration of the coating materials had significant effect on the MIC and MBC values of AgNPs ( $P=0.0468$  and  $P=0.061$ , respectively).

Bacterial types included Gram-negative bacteria represented by *E. coli* and Gram-positive bacteria represented by two strains, MRSA and MSSA. MIC and MBC values of different AgNPs dispersions against *E. coli* were significantly higher than those against MRSA and MSSA ( $P<0.0001$  and  $P<0.01$ , respectively). Also, the effect of AgNPs dispersions against sensitive (MSSA) and resistant (MRSA)

Gram-positive bacteria was studied. MIC values of AgNPs dispersions against MRSA showed twofold significant increase than those against MSSA ( $P>0.0001$ ). However, MBC values against both MRSA and MSSA showed no significant difference, which indicates that AgNPs activity was not affected by the different resistance mechanisms exerted by MRSA.<sup>8</sup>

The effect of AgNPs dispersions on the bacterial growth was confirmed using agar-well diffusion method (Table 4). Results indicated that the effects of coating agents and bacterial type on the measured inhibition zones were similar to those obtained from MIC and MBC measurements. MSSA and MRSA were significantly ( $P<0.01$ ) more sensitive to the antibacterial effect of the AgNPs dispersions than *E. coli* which exhibited the lowest

**Table 4** Determination of inhibition zones for different AgNPs dispersions against MSSA, MRSA, and *Escherichia coli* using agar-well diffusion method. Statistical data indicate the main effects of bacterial type on the inhibition zones, regardless of the coating type, and the main effects of coating agents, regardless of the bacterial type

Dispersion (240 µg AgNPs/mL; 200 µL)	Zone of inhibition ± SD (mm) (n=3)			Main effects of coating ± SD (mM)
	MSSA	MRSA	<i>E. coli</i>	
Dispersion (240 µg AgNPs/mL; 200 µL)				
5 mM PEG-coated AgNPs	22.0 <sup>a</sup> ±0.5	21.0 <sup>a</sup> ±0.5	16.0 <sup>b</sup> ±0.0	19.7 <sup>a</sup> ±0.2
10 mM PEG-coated AgNPs	23.0 <sup>a</sup> ±0.5	22.0 <sup>a</sup> ±0.0	16.0 <sup>b</sup> ±0.0	20.2 <sup>a</sup> ±0.2
5 mM SDS-coated AgNPs	25.0 <sup>a</sup> ±0.5	23.0 <sup>a</sup> ±0.0	18.0 <sup>b</sup> ±0.5	21.7 <sup>b</sup> ±0.2
10 mM SDS-coated AgNPs	25.0 <sup>a</sup> ±0.0	24.0 <sup>a</sup> ±0.0	19.0 <sup>b</sup> ±1.5	22.5 <sup>c</sup> ±0.2
5 mM β-CD-coated AgNPs	23.0 <sup>a</sup> ±0.5	21.0 <sup>a</sup> ±0.0	17.0 <sup>b</sup> ±1.0	20.2 <sup>a</sup> ±0.2
10 mM β-CD-coated AgNPs	23.0 <sup>a</sup> ±0.5	22.0 <sup>a</sup> ±0.0	17.0 <sup>b</sup> ±0.0	20.3 <sup>a</sup> ±0.2
Uncoated AgNPs	22.0 <sup>a</sup> ±0.5	21.0 <sup>a</sup> ±0.5	16.0 <sup>b</sup> ±0.0	19.7 <sup>a</sup> ±0.2
Main effects of bacteria ± SD (mM)	22.8 <sup>a</sup> ±0.1	22.1 <sup>a</sup> ±0.1	16.7 <sup>b</sup> ±0.1	

**Notes:** <sup>a-c</sup>Refer to significant differences at the same row (between bacterial types). <sup>A-C</sup>Refer to significant differences at the same column (between coating agents).

**Abbreviations:** AgNPs, silver nanoparticles; MSSA, methicillin-sensitive *Staphylococcus aureus*; MRSA, methicillin-resistant *S. aureus*; SD, standard deviation; PEG, polyethylene glycol; SDS, sodium dodecyl sulfate; β-CD, β-cyclodextrin.

inhibition zone. The sensitivity of both MSSA and MRSA to AgNPs dispersions was similar ( $P > 0.01$ ), except in the case of 5 mM SDS-coated AgNPs and 5 mM  $\beta$ -CD-coated AgNPs. Regarding the coating materials, there were no significant differences in the antibacterial activity of AgNPs dispersions ( $P > 0.01$ ), except for 5 and 10 mM SDS-coated AgNPs. SDS-coated AgNPs exerted a slightly enhanced activity, which might be due to SDS ability to disrupt bacterial cell wall and increase its permeability.<sup>19</sup> Some studies reported a higher sensitivity for *E. coli* than *S. aureus* toward AgNPs.<sup>23,60,61</sup> This was explained by the different characteristics of Gram-positive and Gram-negative bacteria in the membrane structure particularly in the thickness of the peptidoglycan layer.<sup>59</sup> However, several authors reported that the antibacterial activity of AgNPs dispersions is more pronounced in case of MSSA and MRSA, as compared to *E. coli*.<sup>10,20,62,63</sup> Therefore, the bacterial membrane structure is not the only factor that affects the antibacterial activity of nanoparticles, but also the sources of the bacterial strains.

## Evaluation of AgNPs-loaded hydrogels

The formulated AgNPs hydrogels (F1–F25) were homogeneous with brownish color due to the presence of AgNPs, while the blank hydrogels were transparent. Formulation F24 (5 mM SDS-coated AgNPs loaded in 3% [w/w] chitosan hydrogel) precipitated immediately after preparation, which could be due to interaction between the anionic surfactant SDS and the positively charged chitosan polymer (gelling agent). Hence, this formulation was excluded from further characterizations and studies.

In order to study the flow behavior of the formulations, viscosity values were plotted against spindle speeds (rpm) and examined for the shape of the generated rheograms. It was noticed from the rheograms (Figure S2) that all the prepared AgNPs-loaded hydrogel formulations showed a pseudoplastic behavior (shear thinning behavior). Hydrogels with shear thinning behavior flow constantly under shear stress and recover quickly after stress removal.<sup>64</sup> This behavior is prerequisite for gel formulations intended for topical applications to fulfill the filling and application requirements.<sup>65</sup> Although incorporation of AgNPs within the polymer network of the prepared hydrogels did not affect flow behavior, it decreased viscosity when compared to the blank hydrogels (data not shown). Generally, uncoated AgNPs-loaded hydrogels exhibited the highest viscosity followed by those loaded with PEG-coated AgNPs. AgNPs-loaded hydrogels formulated using chitosan and Pluronic

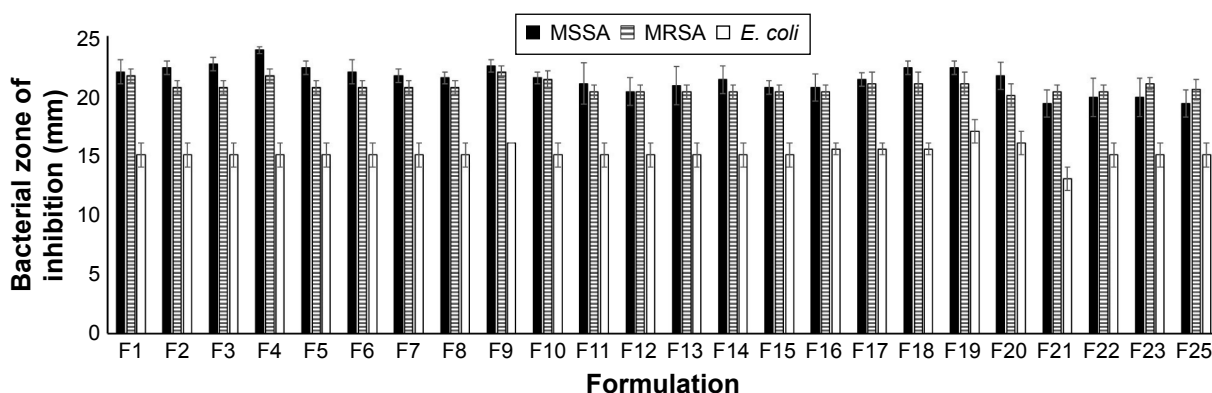
F-127 as gelling agents exhibited the highest viscosity, while those prepared using HPMC showed the lowest viscosity. Similar to the viscosity study, hydrogels with low viscosity showed high spreadability (data not shown).

Release studies showed that hydrogel formulations exhibited AgNPs release (30%–90%) when compared to AgNPs dispersion (100%) after 3 h. All the prepared hydrogels released AgNPs at a concentration ( $\geq 10 \mu\text{g}$  AgNPs after 30 min) that exceeded their MIC against MSSA, MRSA, and *E. coli* (0.93–7.5  $\mu\text{g}/\text{mL}$ ). AgNPs-loaded hydrogels could be used as promising topical antibacterial formulations that allow rapid release of AgNPs, and hence, rapid antibacterial effect at the infection sites before disease progression or gel clearance occurs.

Time-dependent in vitro release of AgNPs over 3 h from all the studied hydrogel formulations is shown in Figure S3. Differences in the release of AgNPs from the prepared hydrogel formulations after 1.5 h were evaluated using two-way ANOVA to study the effect of coating agent and gelling polymer (Table S1). Na CMC, Na alginate, and chitosan hydrogels loaded with 5 mM  $\beta$ -CD-coated AgNPs exhibited the lowest release rate, which might be due to the crosslinking that occurred between cyclodextrin and the polymer matrices which retarded AgNPs release. Bibby et al<sup>66</sup> reported a crosslinking between CD and polymeric formulations which resulted in a decrease in the polymer mesh size, thus decreasing drug release. Izawa et al<sup>67</sup> designed a unique system for controlled release of ondansetron which was composed of  $\beta$ -CD crosslinked with alginate as the main chain polymer. Many authors reported the advantages of using  $\beta$ -CD and chitosan together in controlling drug release from polymeric matrices.<sup>68,69</sup> Regarding other coating agents, the release from hydrogel formulations was in the following descending order: 10 mM PEG-, 5 mM PEG-, and 5 mM SDS-coated-AgNPs-loaded and uncoated AgNPs-loaded formulations. The resulted release patterns of 10 mM PEG-coated-AgNPs-loaded hydrogel might be attributed to its low viscosity, whereas the higher viscosity exhibited by uncoated AgNPs-loaded hydrogels resulted in a slower release pattern. Analysis of the kinetics of the in vitro release of AgNPs demonstrated that the release kinetics of AgNPs from most of the hydrogel formulations was best fitted to Higuchi diffusion model. Further details are provided in the Supplementary materials (Table S2).

## Antibacterial activity of hydrogels

Agar-well diffusion method was used to evaluate the release of AgNPs in the agar plates and their subsequent



**Figure 5** Bacterial inhibition zones of AgNPs-loaded hydrogels against MSSA, MRSA, and *Escherichia coli* using agar-well diffusion method, where F1–F5, F6–F10, F11–F15, F16–F20, and F21–F25 are Na CMC, Na alginate, HPMC, Pluronic F-127, and chitosan hydrogels loaded with AgNPs, respectively.

**Abbreviations:** AgNPs, silver nanoparticles; MSSA, methicillin-sensitive *Staphylococcus aureus*; MRSA, methicillin-resistant *S. aureus*; Na CMC, sodium carboxymethyl cellulose; Na alginate, sodium alginate; HPMC, hydroxypropylmethyl cellulose.

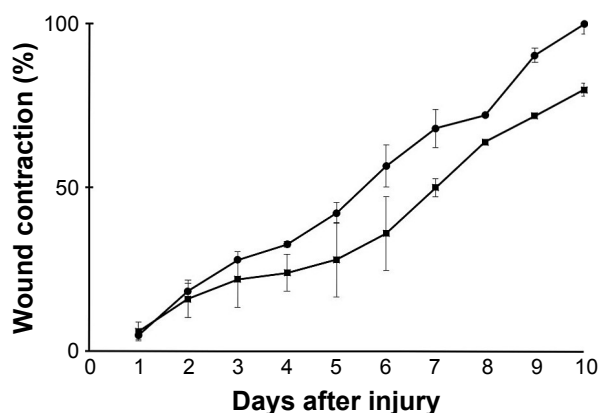
antimicrobial activity and zones of bacterial growth inhibition (Figure 5). All the tested formulations demonstrated antibacterial activities. The bacterial inhibition zone data were analyzed by factorial ANOVA, to study the main effects of gelling polymers, coating agents, and bacterial type. No zone of inhibition was obtained with the control gel formulations (without AgNPs). Gelling agents included five levels (Na CMC, Na alginate, HPMC, Pluronic F-127, and chitosan), and coating agents also had five levels (no-coat, 5 mM PEG 6000, 10 mM PEG 6000, 5 mM SDS, and 5 mM  $\beta$ -CD), while bacteria comprised three levels (MSSA, MRSA, and *E. coli*). Similar to what has been observed in the antibacterial activity study of AgNPs dispersions, AgNPs with different coating agents showed similar ( $P > 0.001$ ) effect on the bacterial growth inhibition zone with all the studied gelling agents except for Na alginate gel and Pluronic F-127 gel, where using 5 mM SDS-coated AgNPs showed an increase in the bacterial growth inhibition zones. Formulations with different gelling agents showed significant differences in the bacterial growth inhibition zones where the zones of inhibition for the Na CMC, Na alginate, and Pluronic F-127 gelling agents were significantly ( $P < 0.001$ ) wider than those of HPMC and chitosan. Regarding the bacterial type, all the formulations showed significant differences in the bacterial growth inhibition zones between *E. coli* and both MSSA and MRSA, where the latter two bacteria demonstrated higher sensitivity to the AgNPs-loaded hydrogels. These results were similar to those obtained from the determination of MIC and MBC, and agar-well diffusion methods for the AgNPs dispersions (vide supra). In conclusion, the most evident effect on the bacterial growth inhibition zones was for the bacterial type, followed by the gelling agent and the coating type.

## In vivo studies

From the previous in vitro studies and characteristics of the prepared formulations, it was noticed that, although SDS-coated AgNPs showed a slight increase in the antibacterial activity in the agar-well diffusion study, 10 mM PEG-coated AgNPs dispersion was preferred. The use of PEG as a coating agent resulted in a stable dispersion without any particles aggregation or sedimentation for at least 9 months. PEG-coated-AgNPs-loaded hydrogel also showed faster release of the loaded AgNPs from the hydrogels. Hydrogels prepared using 10 mM PEG as the coating agent and 3% Na CMC as the gelling agent had the most reasonable viscosity values among the tested AgNPs-loaded hydrogels, in addition to an intermediate spreadability and good consistency. Hence, formulation (F3) that contains 10 mM PEG and 3% (w/w) Na CMC was selected for the further animal studies to investigate the in vivo efficacy of AgNPs. First, the wound-healing efficacy and prevention of initial growth of any bacteria at wound sites were studied. Second, the efficacy against MRSA-infected wound model was investigated.

## Wound healing

Dermal wound healing is a complicated process that involves proper harmony between several skin components to allow for the repair of the damaged tissues and to restore the normal skin functions.<sup>70</sup> The use of a broad-spectrum antibacterial agent (eg, AgNPs) that provides a septic environment might aid in the effective wound repair. Liu et al<sup>71</sup> highlighted the role of AgNPs in wound-healing process and demonstrated that AgNPs could enhance reepithelialization and wound contraction process. Ten adult female rats were used and divided randomly into two groups (n=5). After skin excision, the wound was left undressed to the open environment.



**Figure 6** Percent of wound contraction in rats during the 10 d of treatment. Hydrogel loaded with 0.1% AgNPs (circles) was compared to 1% silver sulfadiazine cream (squares).

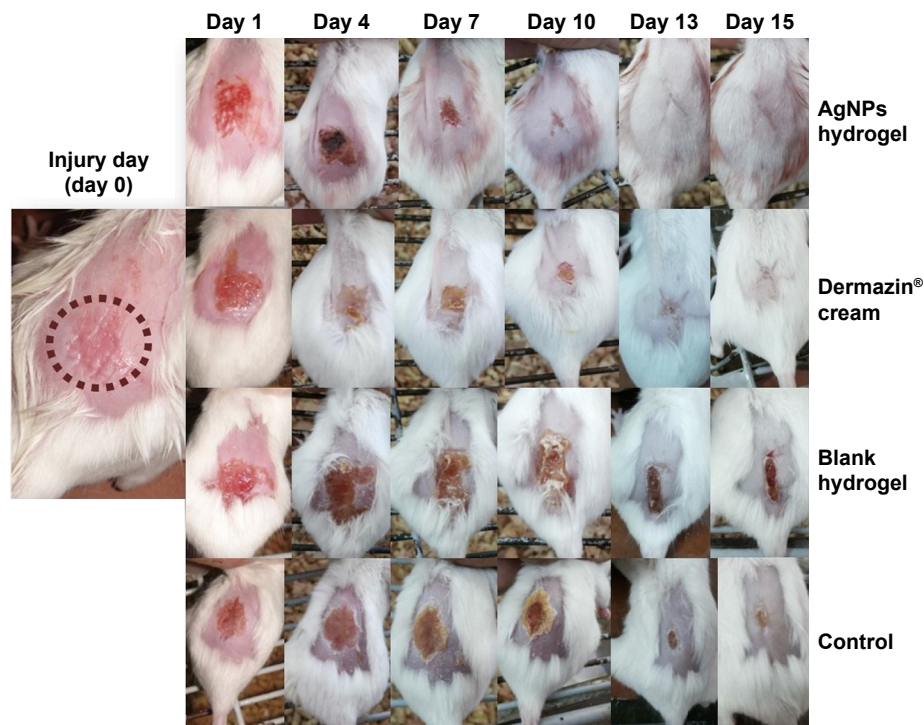
**Abbreviation:** AgNPs, silver nanoparticles.

Group I received 0.1% AgNPs hydrogel (F3), and group II received the commercially available 1% silver sulfadiazine cream. Treatment of both groups was started at the day of wound creation and lasted for 10 d. Group I exhibited faster wound closure as compared to group II that received the commercially available cream, as demonstrated by the higher percentage of wound contraction during the experiment as compared to the values measured for the commercial silver sulfadiazine cream (Figure 6). Similar observations were obtained by Tian et al<sup>42</sup> who investigated the wound-healing efficacy of AgNPs-coated dressing in different diabetic and nondiabetic animal wound models. Rapid wound healing with better scar appearance and normal hair growth in the AgNPs-treated group, as compared to silver sulfadiazine-treated group, was observed. Franková et al<sup>72</sup> have studied the in vitro effect of AgNPs as wound-healing promoters. They have reported that AgNPs decreased the release of some pro-inflammatory cytokines and growth factors from normal human epidermal keratinocytes (NHEKs) and normal human dermal fibroblasts after 24 h at all the studied AgNPs concentrations (0.25, 2.5, and 25  $\mu\text{g}/\text{mL}$ ). Other authors studied the effect of different concentrations of polyvinylpyrrolidone-coated AgNPs on NHEKs and reported that AgNPs decreased the keratinocytes viability, metabolism and proliferation, and their migratory potential.<sup>73</sup> However, they mentioned that the effect might be different in vivo as keratinocytes are surrounded by other types of cells that protect them against harmful stimuli. In our in vivo wound-healing study, 0.1% AgNPs-loaded hydrogel promoted wound healing more than the commercially available 1% silver sulfadiazine.

## In vivo antibacterial activity

Skin cuts and abrasions are susceptible to infections, and it is important to rapidly eradicate the bacteria from the infectious

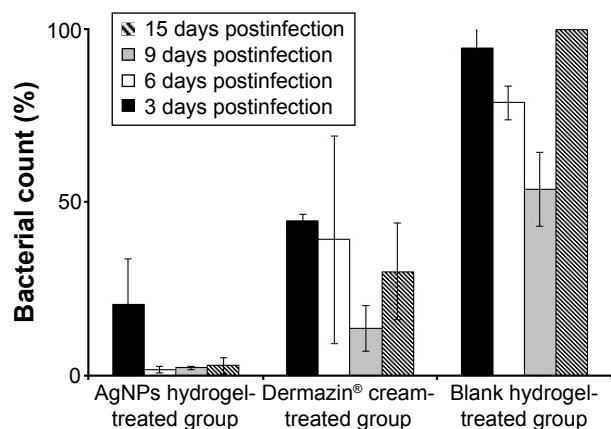
lesions to allow for rapid wound healing and to reduce the morbidity rate.<sup>74</sup> MRSA is the main cause of worldwide nosocomial infections and usually causes superficial lesions in human skin, localized abscesses, septicemia, pneumonia, and bacteremia.<sup>8</sup> The antibacterial activity of the biologically synthesized 0.1% 10 mM PEG-coated-AgNPs-loaded Na CMC hydrogel (F3) against MRSA-infected skin model was tested in mice. Dai et al<sup>41</sup> reported that this experimental model produces a chronic and stable wound infection which might develop a severe morbidity or mortality in mice. All the studied groups were observed, and representative mice skin abrasion wounds were imaged daily at different time intervals (Figure 7). The group treated with AgNPs-loaded hydrogel (F3) showed more efficient healing of the infected abrasion as compared to the group treated with the silver sulfadiazine cream. No healing was observed in the infected wounds of the untreated and blank hydrogel-treated mice groups, in addition to oozing of pus at the infected sites. Wound swabs from each group were taken at different time intervals and cultured on a selective agar media to measure the growth of the microorganisms. The average bacterial count in the mice groups before beginning of the treatment (1 d) was ca.  $5 \times 10^5$  CFU/mL. The percentages of bacterial count (as compared to the control untreated group) in the mice skin infected abrasions (n=5) at different time intervals after daily application of the topical treatments were calculated and are presented in Figure 8. The mice group treated with 0.1% AgNPs hydrogel (F3) exhibited a significantly ( $P < 0.05$ ) faster and superior bacterial growth reduction when compared to the other groups during the 15 d of the treatment. AgNPs hydrogel-treated skin exhibited 20.5% of the bacterial count in the control untreated skin versus 44.7% and 94.5% for the silver sulfadiazine cream- and blank hydrogel-treated groups, respectively, in the first bacterial swab analysis after beginning of the treatment (3 d). In the last bacterial swab analysis (15 d), AgNPs hydrogel-treated skin exhibited 2.7% of the bacterial count in the control untreated skin, whereas silver sulfadiazine cream- and the blank hydrogel-treated groups exhibited 30% and 100% of the bacterial count, respectively, as compared to the previous bacterial swab analysis (9 d). The observed increase of bacterial count measured at 15 d compared to 9 d in the silver sulfadiazine cream-treated mice group (Figure 8) was not significant ( $P > 0.05$ ). In addition, the bacterial count in the blank hydrogel-treated mice group decreased insignificantly over time ( $P > 0.05$ ) when compared to the 0.1% AgNPs hydrogel-treated group and 1% silver sulfadiazine cream-treated group. This might be due to the normal recovery of their immune systems over time after initial suppression using cyclophosphamide.



**Figure 7** Successive images of representative mice skin abrasion wounds infected with MRSA at different time intervals. Two groups were treated with 0.1% AgNPs hydrogel and 1% silver sulfadiazine cream. The two other groups were the blank hydrogel-treated group and control untreated mice.

**Abbreviations:** MRSA, methicillin-resistant *Staphylococcus aureus*; AgNPs, silver nanoparticles.

Mortality and morbidity of animals were observed during the experiment in the different studied groups (Table 5). No weight loss and suppuration or death were observed in mice treated with the AgNPs hydrogel with normal hair growth, whereas the silver sulfadiazine cream-treated group showed 20% weight loss and 20% mortality. Similar results were obtained by Hendi<sup>75</sup> who reported that AgNPs-treated group showed the most resemblance to normal skin with thin epidermis and nearly normal hair follicles.



**Figure 8** Bacterial count in the infected abrasion wounds of different mice groups at different time intervals expressed as percentages of the bacterial counts in the control untreated mice group. The studied groups are 0.1% AgNPs hydrogel-treated group, 1% silver sulfadiazine cream-treated group, and blank hydrogel-treated group.

**Abbreviation:** AgNPs, silver nanoparticles.

Blank hydrogel-treated mice and untreated mice showed 50% weight loss and 60%–80% suppuration at the end of the treatment course. Proliferation of MRSA at the injury sites in the latter two mice groups resulted in the observed inflammation, pus formation, and disrupted wound-healing process.

### Histopathological studies

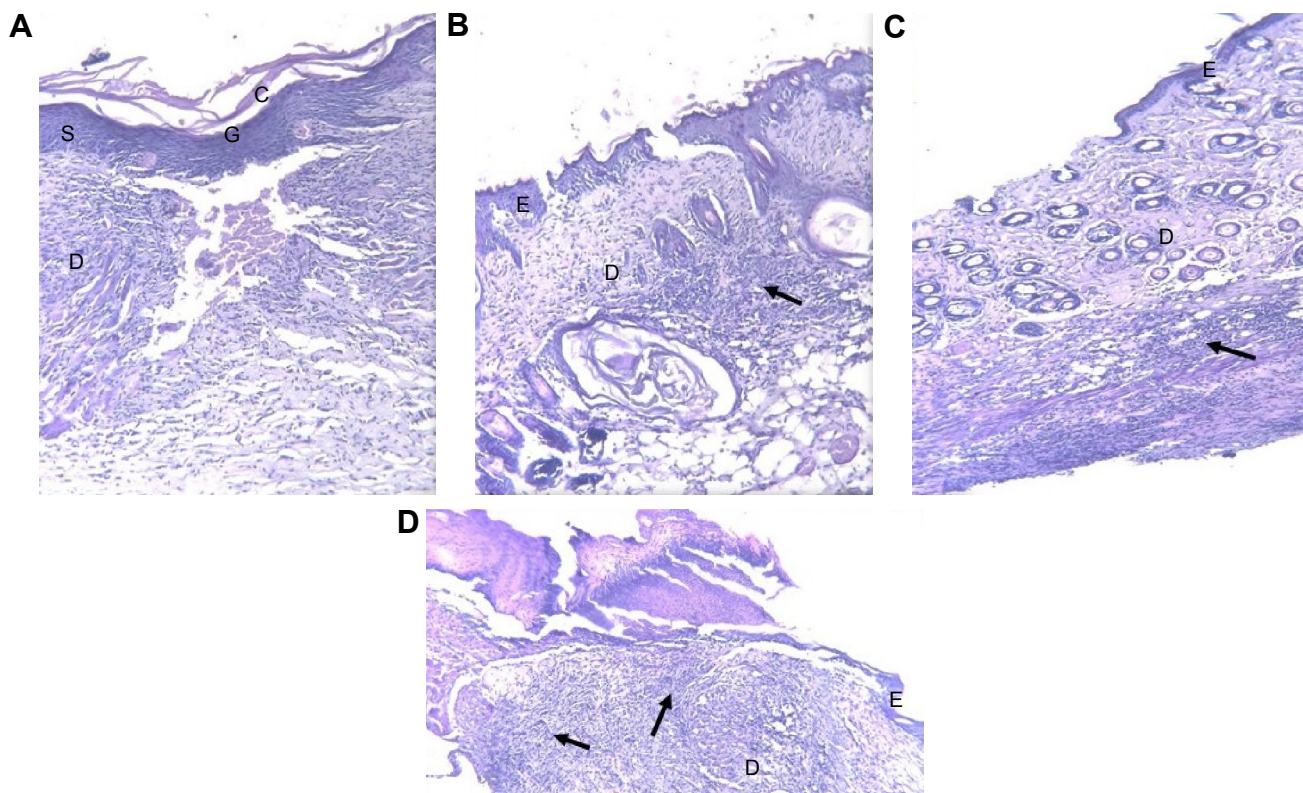
Light microscopic examination of the skin samples after staining with hematoxylin and eosin showed great histological differences between the four groups. Skin samples from the 0.1% AgNPs hydrogel-treated group showed a well-stratified

**Table 5** Mortality and morbidity observed at the end of the study in the MRSA-infected mice groups

Observation	Percentage of animals (n=5)			
	AgNPs hydrogel	silver sulfadiazine cream	Blank hydrogel	Untreated mice
Morbidity*				
Suppuration	0%	0%	80%	60%
Weight loss	0%	20%	50%	50%
Hair growth	Normal	Normal	Retarded	Retarded
Argyria**	0%	0%	0%	0%
Mortality***	0%	20%	0%	0%

**Notes:** \*Morbidity refers to the relative incidence of the disease in the animal. \*\*Argyria refers to gray staining of the skin due to deposition of Ag. \*\*\*Mortality refers to death of the animals.

**Abbreviations:** MRSA, methicillin-resistant *Staphylococcus aureus*; AgNPs, silver nanoparticles.



**Figure 9** Light microscopy images of representative skin samples (stained with hematoxylin and eosin) ( $\times 400$ ) after 15 d of treatment: (A) 0.1% AgNPs hydrogel-treated group; (B) 1% silver sulfadiazine cream-treated group; (C) blank hydrogel-treated group; and (D) control group (untreated mice). Arrows refer to lymphocytes. C, G, S, D, and E represent stratum corneum, stratum granulosum, stratum spinosum, dermis, and epidermis, respectively.

**Abbreviation:** AgNPs, silver nanoparticles.

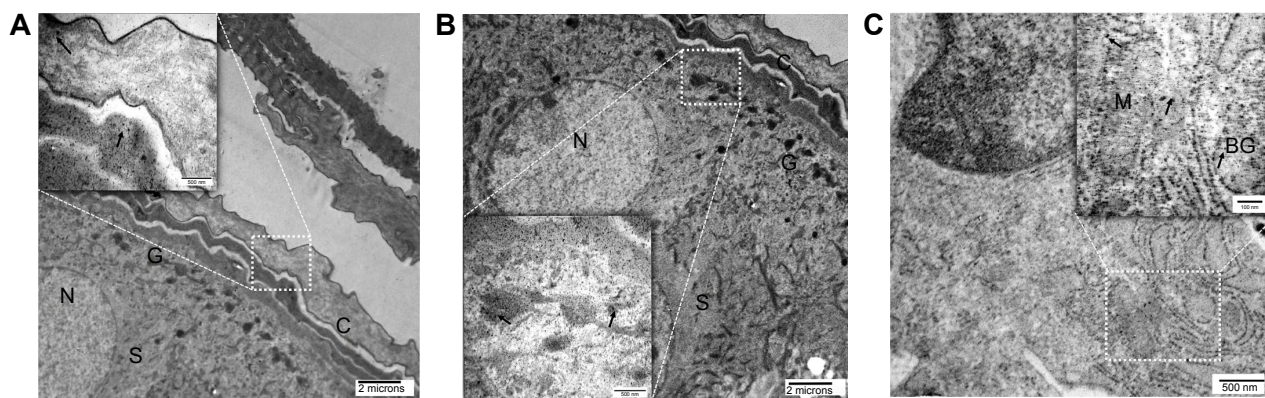
epidermis, complete restoration of epidermal layers, and complete basal, spinosum, granular, and cornfield layers (Figure 9A). Mild damage in the epidermal layers and incomplete healing were observed in the 1% silver sulfadiazine cream-treated skin (Figure 9B). In addition, slight infiltration of inflammatory cells (lymphocytes) was observed, which indicates a prolonged inflammatory response at the wound sites.<sup>76</sup> In the representative skin samples from the blank hydrogel-treated group and control group, unhealed skin was clearly observed, where epidermis was severely damaged and no reformation was observed (Figure 9C and D). A varying degree of infiltration of inflammatory cells with accumulation of lymphocytes was observed, and damage was severe in both cases. In addition, severe focal dermal inflammation with intercellular edema was observed.

Skin samples from mice group that was treated daily with 0.1% AgNPs-loaded 3% Na CMC hydrogel for 15 d were examined under TEM. The uptake, distribution, and localization of AgNPs in the epidermal layers are demonstrated in the micrographs (Figure 10A and B), while localization of the AgNPs inside the Langerhans cell (antigen-presenting immune cells) is indicated in the stratum spinosum layer (Figure 10C).

## Conclusion

In the current work, fungi-assisted biological reduction of  $\text{AgNO}_3$  solution resulted in spherical-shaped AgNPs with small and homogenous size, which were coated with several materials prior to formulation into hydrogels suitable for topical applications. Coating of the nanoparticles with PEG reduced the sedimentation rate of particles in their aqueous dispersions and greatly enhanced the shelf-life stability over 9 months. AgNPs dispersions were effective against both Gram-positive (MRSA and MSSA) and Gram-negative (*E. coli*) bacteria with preferential inhibitory effect against Gram-positive bacteria. Homogenous hydrogels were prepared using various gelling agents, where Na CMC hydrogel loaded with 10 mM PEG-coated AgNPs showed superior characteristics based on the viscosity, spreadability, and release studies, and therefore, was selected for further in vivo studies. Na CMC hydrogel loaded with 0.1% 10 mM PEG-coated AgNPs showed superior in vivo antibacterial activity and wound healing with good skin appearance and normal hair growth when tested on MRSA-infected wound mice model as compared to the commercially available 1% silver sulfadiazine cream. Complete restoration of the epidermal layers in the mice skin treated with the AgNPs





**Figure 10** Transmission electron micrographs: (A) epidermal layers with magnification of stratum corneum; (B) epidermal layers with magnification of stratum granulosum ( $\times 4,800$ , inset:  $\times 29,000$ ), (C) Langerhans cell in stratum spinosum ( $\times 19,000$ , inset:  $\times 72,000$ ). Inset arrows indicate localized AgNPs. C, G, S, N, BG, and M represent stratum corneum, stratum granulosum, stratum spinosum, nucleus, Birbeck granules, and mitochondria, respectively.

**Abbreviation:** AgNPs, silver nanoparticles.

hydrogel was demonstrated, and localization of the nanoparticles in the different epidermal layers was detected in the TEM micrographs of the treated skin. Lower toxicity and higher tolerability were observed in the mice treated with the AgNPs-loaded hydrogels as compared to the silver sulfadiazine cream-treated group as indicated in the animal experiments and histological studies.

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## Disclosure

The authors report no conflicts of interest in this work.

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