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A bio-inspired strategy for the synthesis of zinc oxide nanoparticles (ZnO NPs) using the cell extract of cyanobacterium *Nostoc* sp. EA03: from biological function to toxicity evaluation

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Cyanobacteria, as one of the largest groups of phototrophic bacteria, have a high potential as an excellent source of fine chemicals and bioactive compounds, including lipid-like compounds, amino acid derivatives, proteins, and pigments. This study aimed to synthesize ZnO nanoparticles using the cell extract of the cyanobacterium *Nostoc* sp. EA03 (CEN-ZnO NPs) through a rapid and eco-friendly approach. The biosynthesized nanoparticles, CEN-ZnO NPs, were characterized by UV-Vis spectroscopy, X-ray diffraction (XRD), zeta potential measurement, differential scanning calorimetry (DSC)/thermogravimetric analysis (TGA), FTIR, SEM, TEM, and EDX spectroscopy. The UV-Vis spectrum showed an absorption peak at 370 nm. The star-shaped CEN-ZnO NPs, as observed in the TEM and SEM images, had an average diameter of 50–80 nm. MIC and MBC values for *E. coli*, *P. aeruginosa* and *S. aureus*, were determined to be, respectively, 2000, 2000, and 64 $\mu\text{g ml}^{-1}$, and 2500, 2500 and 128 $\mu\text{g ml}^{-1}$. Further analysis through confocal laser scanning microscopy (CLSM) provided the observable confirmation that the CEN-ZnO NPs stunted the bacterial growth, preventing the formation of exopolysaccharides. The AFM analysis of surface topography of bacterial biofilm samples treated with CEN-ZnO NPs showed a rugged topography in some parts of the biofilm surface, indicating the destruction of biofilms. In contrast, in the untreated control samples, the structured biofilms were flat and prominent. MTT assay indicated that CEN-ZnO NPs had less cytotoxicity on the MRC-5 lung fibroblast cells compared with the cancerous treated A549 cells. As the concentration of the CEN-ZnO NPs increased, the amount of ROS produced in the tested bacterial strains also increased. Analyzing the data obtained from flow cytometry showed that the higher concentrations of CEN-ZnO NPs lead to a reduction in the viability of *P. aeruginosa* PAO1, *E. coli* and *S. aureus*. The biosynthesized ZnO nanoparticles using *Nostoc* cell extracts exhibited different attributes, inspiring enough to be considered for further investigation.

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1. Introduction

Nanoparticles (NPs) are particles with one dimension ranging from 1 to 100 nanometers. They exhibit an enhanced catalytic reactivity, thermal conductivity, non-linear optical performance, and chemical steadiness owing to their large surface area to volume ratio.¹ Nano materials are known as the miracles of modern medicine.² Metallic nanoparticles have different physical and chemical properties due to their wide surface ratio, while their bulk forms may not have these features.³ The synthesis of nanoparticles (NPs) using plant

extracts, bacteria, fungi, algae, *etc.*, recognized as green technology, is the biocompatible, cost-effective and safe approach.⁴ The use of green and eco-friendly approaches for the synthesis of NPs has allowed them to develop for use in food, space, consumer healthcare, and the cosmetic industry. It has also vastly improved our understanding of their role in the control of disease and healthcare-associated infections in recent years.⁵ Various types of inorganic metal oxides have been synthesized and used in recent studies like TiO_2 , CuO, and ZnO. Of all these metal oxides, ZnO NPs is of greatest interest, because their production is not expensive and easy to prepare safely. Zinc oxide (ZnO) is an n-type semi-conducting metal oxide and has attracted significant attention in the past three years due to its wide range of applicability in the field of electronics, optics, and biomedical systems. US FDA has enlisted ZnO as GRAS (generally recognized as safe) metal oxide.^{6,7}

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The presence of bioactive compounds in plants, algae, and microorganisms has made them a good candidate for the synthesis of nanoparticles (NPs).⁸ Cyanobacteria as a group of ancient and abundant microorganisms with prokaryotic cell structures have the capability of carbon dioxide-dependent photosynthesis.⁹ Among microorganisms, cyanobacteria are of great interest in NP synthesis, as they are a potential source of novel metabolites that have great importance from a biotechnological and industrial point of view.⁹ It has shown that they can utilize as a good source with biological features for the synthesis of gold (Au), silver (Ag), platinum (Pt), and palladium (Pd) nanoparticles.¹⁰ The genus *Nostoc* is a valuable source of a wide spectrum of secondary metabolites, such as antioxidant enzymes, phycobiliproteins, vitamins, and phenolic compounds that have many uses as anticancer, anti-HIV, anti-malarial, antifungal, and/or antimicrobial drugs^{11–13}. *Nostoc* cell extract with a rich source of materials such as amino functional group, carboxyl, and hydroxyl, can be used as a metal oxide reducing and capping agents for NPs biosynthesis.¹⁴ Here, the green synthesis of zinc oxide nanoparticles was achieved using the cell extract of an indigenous cyanobacterial strain, *Nostoc* sp. EA03, in the first study of its kind. In this regard, the present study attempted to address the issues listed below: (1) isolation and characterization of a local cyanobacterial strain, (2) preparation of the cell extracts of selected cyanobacterial strain, *Nostoc* sp. EA03, (3) synthesis of ZnO NPs using the cell extract of *Nostoc* sp. EA03, (4) characterization of the CEN-ZnO NPs, (5) assessment of the antibacterial, anti-biofilm and antifungal activities of different concentrations of CEN-ZnO NPs against three important clinical bacterial isolates including *E. coli* 25922, *P. aeruginosa* PAO1, and *S. aureus* 25923, (6) *in vitro* analysis of cytotoxic effect of CEN-ZnO NPs on human lung fibroblast cells MRC-5 and the cancerous human lung alveolar epithelial cells A549, (7) the measurement of reactive oxygen species (ROS) levels in tested bacterial cells after exposure to different concentrations of CEN-ZnO.

2. Materials and methods

2.1. Materials and strains

Zinc acetate dehydrate ($\geq 98\%$ Zn $(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$) and sodium hydroxide ($\geq 98\%$ NaOH) were purchased from Sigma-Aldrich, Australia. The clinical bacterial strains including *E. coli* ATCC 25922, *P. aeruginosa* PAO1, and *S. aureus* ATCC 25923 have received donations from Pasteur Institute of Iran.

2.2. Sampling, isolation, and molecular identification of cyanobacterial strain

To isolate the cyanobacterial strain, sampling was carried out independently from the aquatic environments including lakes, rivers in Alborz province (Iran). The samples were quickly transferred to the laboratory in suitable conditions. Detail regarding isolation and identification of the cyanobacterial strain was performed as described earlier.^{15–18}

2.3. Biomass collection

The selected cyanobacterial strain was grown in laboratory condition in 2000 ml Erlenmeyer flasks containing 400 ml BG11 medium and stirred daily, with continuous illumination (a light intensity of 2000–3000 $1\times$), at a temperature of 28 °C and 65% humidity for 10–12 days to collect the bacterial biomass. The cyanobacterial cells were collected in the late logarithmic phase by centrifugation at 5000 rpm/15 min and then washed two times with deionized water until all the medium debris removed. The collected biomass was used to obtain the cell extracts.

2.4. Preparation of cyanobacterial cell extracts

The biomass was collected and dried in the oven at 45 °C (to reach a constant weight) for 12 hours. In the next step, the dried biomass was crushed by a sterile mortar until it was smooth and uniform. The biomass powder was stored in a vial at 4 °C for further use. To obtain the cell extract, 1 g of dried biomass was dissolved in 50 ml of deionized water using a stirrer at 60 °C for 15 min. The solution was filtered through Whatman no. 41 filter paper and the bacterial cell extract was obtained. The cell extract was kept at 4 °C for the subsequent steps.

2.5. Gas chromatography-mass spectroscopy (GC-MS) analysis

The GC-MS analysis of the cyanobacterial cell extract was performed using a Scion 436-GC Bruker system and Gas chromatograph interfaced to a mass spectrometer (GC-MS) equipped with a BR-5MS (5% diphenyl/95% dimethyl polysiloxane) column, 30 m \times 0.25 mm ID \times 0.25 μm df. The defined conditions for GC-MS detection was carried out as described earlier.¹⁹

2.6. Synthesis of CEN-ZnO NPs and optimization

For the synthesis of CEN-ZnO NPs, 1000 μl of cell extract was added drop by drop to 50 ml zinc acetate dehydrate (Zn $(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$, 0.02 M) through constant stirring using a magnetic stirrer. After 10 min of stirring, 0.1 M NaOH was added drop-wise to the constantly stirred solution until a faded white solution appeared (the color of the reaction mixture changes from green to white). The resultant faded white solution was maintained under vigorous stirring for 2 h followed by centrifugation at 5000 rpm/15 min. The pellets were then washed with Milli-Q water three times and dried in a vacuum oven at 60 °C for 12 h. To optimize the synthesis of CEN-ZnO NPs, a set of experiments was designed at different pH ranges and different concentrations of cell extract, separately. The pH of the reaction mixture was adjusted in the range of 7–9 by adding 0.1 M NaOH, and different concentrations of the cell extract (1000–3000 μl) were used in the reaction mixture. The dried powder material was kept in an airtight microtube at room temperature for subsequent analysis.

2.7. Characterization of the synthesized CEN-ZnO NPs

2.7.1. UV-visible (UV-Vis) spectroscopy analysis. The surface plasmon resonance of the resultant CEN-ZnO NPs were



analyzed using a double-beam T50⁺ UV-Vis spectrophotometer (Germany) with a wavelength range of 200–700 nm.

2.7.2. X-ray powder diffraction (XRD) analysis. The XRD pattern of biosynthesized CEN-ZnO NPs was developed by XRD 3003 PTS, SEIFERT (Germany) in the range of $20^\circ \leq 2\theta \leq 80^\circ$ at 40 kV with CuK α radiation ($k = 1.54 \text{ \AA}$), and the crystalline domain diameter of ZnO NPs was calculated from the XRD peaks by the Debye–Scherrer equation $D = 0.9\lambda/\beta \cos \theta$. Here, D is the crystalline domain diameter of ZnO NPs, λ is the wavelength of the X-ray source used (1.541 \AA), and B is the full width at half maximum of the diffraction peak.

2.7.3. Scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDX) analysis. Scanning electron microscopy (SEM) (TESCAN, VEGA3) was used to image and evaluate the morphology and size of the CEN-ZnO NPs. The CEN-ZnO NPs powder was used for SEM at an accelerating voltage of 20 kV. The scientific instrument Sirius SD model (the EDX spectrometer-equipped SEM) was used to determine the elemental analysis of the CEN-ZnO NPs.

2.7.4. Transmission electron microscopy (TEM) analysis. TEM images have obtained with a Philips CM300 (Netherlands) instrument at an accelerating voltage of 100 kV. The preparation of the samples performed by dropping 20 μl of CEN-ZnO NPs on a copper grid (200 mesh), dried at room temperature.

2.7.5. Fourier transform spectroscopy (FTIR) analysis. The chemical structures of ZnO, CEN-ZnO NPs, and cyanobacterial cell extract have studied by FTIR examination. The ZnO, CEN-ZnO NPs and cell extract was finely ground and dispersed in pellets of KBr (Sigma) before performing the FTIR analysis. The FTIR spectrum was recorded in 3500–500 cm^{-1} region on an FTIR system (PerkinElmer, US). The wave number precision and resolution of the spectrum were 0.01 cm^{-1} and 4, respectively.

2.7.6. Zeta potential. The zeta potential of the biosynthesized NPs was determined using the zeta potential analyzer Zetasizer Nano ZSP (ZSP instrument (Malvern), UK). For this purpose, 1 mg of CEN-ZnO NPs powder was added to 1 ml of deionized water and then placed in an ultrasonic bath for 10 minutes. The zeta potential was attained from the electrophoretic mobility using the Smoluchowski equation.

2.8. Evaluation of antibacterial activity of biosynthesized CEN-ZnO NPs (MIC and MBC values)

Three clinical bacterial strains were cultivated in Blood agar and Mueller–Hinton broth (MHB) and stored at -70°C in 30% glycerol. The MIC values were evaluated by the broth microdilution method as recommended by CLSI.²⁰ The MIC of CEN-ZnO NPs was determined against Gram-positive (*S. aureus* ATCC 25923) and Gram-negative bacteria (*P. aeruginosa* PAO1 and *E. coli* 25922) by 96-well microtiter assay. Briefly, 100 μl of Mueller–Hinton broth with CEN-ZnO NPs at different concentrations (*i.e.*, 4, 8, 16, 32, 64, 128, 256, 512, 1000, and 2500 $\mu\text{g ml}^{-1}$) was added. Then, 5 μl of microbial suspension ($1 \times 10^8 \text{ CFU ml}^{-1}$) was added to each well. The plates were incubated at 37°C for 24 h. Positive and negative controls were included as well. Following incubation, the MIC values were listed for each bacterial strain. The minimum concentration of samples that

showed no visible growth was defined as MIC. Aliquots of 100 μl from all tubes with no visible bacterial growth were seeded on MHA plates without CEN-ZnO NPs. The lowest concentration of samples that killed all the initial bacterial inoculum was defined as MBC.

2.9. Biofilm formation

In this experiment, a flat-bottomed 96-well cell culture microtiter plate was used. Bacterial inoculum concentrations were used to form biofilm by *P. aeruginosa* PAO1, *E. coli* 25922, and *S. aureus* 25923. Using 24 hours bacterial culture, the suspension was adjusted to McFarland standard 0.5 ($1 \times 10^8 \text{ CFU ml}^{-1}$) in TSB supplemented with 0.2% glucose. Hundred microliters of each suspension were inoculated into each well. The test was performed using two replicates for each inoculum concentration. The microplates were incubated at 37°C for 20 h.

2.9.1. Determination of minimum biofilm inhibitory concentrations (MBIC) of CEN-ZnO NPs. The wells were filled with the medium as mentioned above. Two-fold CEN-ZnO NP serial dilutions were made in wells in a way that seven different concentrations were tested. The last row of each microtiter plate was used as controls. Using 24 h *P. aeruginosa* PAO1, *E. coli* 25922, and *S. aureus* 25923 cultures, the suspension was adjusted to McFarland standard 0.5 in TSB supplemented with 0.2% glucose. Two hundred microliters of the suspension was inoculated to each well except for the medium controls. The microplates were incubated at 37°C for 20 h. The percentage of biofilm inhibition was expressed by the following formula: $[1 - (\text{OD}_{570 \text{ nm}} \text{ of samples treated with CEN-ZnO NPs} / \text{OD}_{570 \text{ nm}} \text{ of non-treated control samples}) \times 100]$. Positive and negative controls were included.

2.9.2. Tetrazolium chloride (TTC) assay. The TTC solution was obtained by dissolving 0.025 g of powder TTC in 5 cm^3 of deionized water, vortexed and then passed through a 0.22 micron filter. 50 μl of TTC solution was added to the microtiter wells, and the plate located in a shaker incubator for 30 minutes. After this time, the absorbance of the resultant solution was evaluated in microtiter plate reader at 570 nm.²¹

2.9.3. Crystal violet assay (CV assay). First, we rinsed the microtiter plate thrice with sterile PBS; then, 125 μl of 0.1% crystal violet (Merck, Germany) solution was added to each well. After 10 minutes at room temperature, the wells were discharged, and the microplates were carefully washed under running tap water to remove surplus CV dye. Next, acetic acid 30% was added for 10 minutes. After this time, 125 μl of each well was transferred into the new microplate and read through the microtiter plate spectrometry at a wavelength of 570 nm.²²

2.9.4. Determination of minimum biofilm eradication concentrations (MBEC) of CEN-ZnO NPs. In this research, the effect of the CEN-ZnO NPs on one-, three-, and five-day-old biofilms were examined. As mentioned above, the biofilms were established in microtiter plates (Sigma Aldrich, St. Louis, Missouri, USA). After 24, 48, or 72 hours, the media were removed from the wells, and the plates were washed thoroughly with PBS (pH = 7.4) to remove the unattached cells. After that, two-fold serial dilutions of CEN-ZnO NPs were made in wells in



a way that five different concentrations were tested. In our study, the designated concentrations were from 64 to 2500 $\mu\text{g ml}^{-1}$. The culture medium of bacteria without CEN-ZnO NPs was added to the well as the control. The plates were incubated for 24 hours at 37 °C and stained with crystal violet (CV) and TTC.

2.9.5. The analysis of biofilms by confocal laser scanning microscopy (CLSM). Confocal laser scanning microscopy analysis was performed to investigate the effect of the biosynthesized CEN-ZnO NPs on biofilm formed by *S. aureus* and *P. aeruginosa* PAO1. For confocal imaging of the biofilms, they were grown on glass coverslips and microtitre plates.²³ In summary, six-well microtiter plate seeded with glass coverslips were incubated for 24 h at 37 °C in 5 ml of TSB with 2% glucose. The wells were inoculated with 1000 μl of mid-exponential grown cultures. After 24 h, the CEN-ZnO NPs at their MBEC concentrations were added to the overnight grown bacterial cultures and were incubated at 37 °C for 24 h. The coverslips were removed and gently rinsed with sterile PBS and then stained with 15 μl propidium iodide (PI; Sigma) for 15 min at room temperature to detect red (dead) bacterial cells. After being washed in PBS, the cultures were incubated with 50 μl ml^{-1} of fluorescein isothiocyanate (FITC-Sigma) for 15 min at room temperature to stain the glycocalyx matrix green (live). PI is excited at 520 nm, the emission monitored at 620 nm, and FITC is excited and observed at 495 and 525 nm, respectively.²⁴ The untouched biofilms were examined using a confocal electron microscope Leica TCS SP5 II.

2.10. Assessment of bacterial viability by flow cytometry

The bacterial cells including *P. aeruginosa* PAO1, *E. coli* ATCC 25922, and *S. aureus* ATCC 25923 were selected to determine the cell viability and internalization after exposure to the biosynthesized CEN-ZnO NPs by flow cytometry. Briefly, for cell viability assessment, afresh grown suspensions of the bacterial strains (1×10^8 CFU ml^{-1}) were treated with increasing concentrations of CEN-ZnO NPs (500, 1500, and 2500 $\mu\text{g ml}^{-1}$ for *E. coli* and *P. aeruginosa* and 16, 32, and 64 $\mu\text{g ml}^{-1}$ for *S. aureus*) at 37 °C under shaking condition (110 rpm) in a shaker incubator for 20 h. The bacterial viability was assessed using a flow cytometer BD FACS Calibur (BD Biosciences, San Jose, CA, USA) equipped with an argon laser device (excitation at 488 nm, emission 530 nm), and the flow cytometric graphs drawn with the FLOWJO 10.4.2 software that was made available on <https://www.flowjo.com/solutions/>. For laser calibration purpose and gating selection, unstained and single-stained FDA or PI bacteria were also assessed. The setup procedure for flow cytometry analysis was designed as described elsewhere follows: a fluorescein diacetate (FDA) (Sigma Chemicals F7378) stock solution was prepared by dissolving 10 mg FDA in 10 ml acetone and stored in the dark at −20 °C until further use. A working solution was prepared by mixing 10 μl of stock solution with 2.5 ml of distilled water (this solution is stable for only 20 min and should, therefore, be prepared freshly). The working solution was added to each reaction and kept at room temperature for 15 min in dark condition. Unstained samples were used for light scatter gating, and the control was the sample without CEN-ZnO NP treatment.

2.10.1. Internalization assay. Propidium iodide (PI) staining was used as the dye to check the permeability of the bacteria that were exposed to CEN-ZnO NPs. For this purpose, a fresh culture of the bacterial strains prepared as the bacterial concentration was 1×10^8 CFU ml^{-1} and exposed to the NP concentrations of 500, 1500, and 2500 $\mu\text{g ml}^{-1}$ for *E. coli* and *P. aeruginosa* and 16, 32, and 64 $\mu\text{g ml}^{-1}$ for *S. aureus* for 30 minutes. Subsequently, 2 μl of PI (20 μM) was added to the samples for 15 min in darkness at room temperature. Briefly, 50 μl of the treated cells were added to 950 μl of $1 \times$ PBS, mixed thoroughly and analyzed using flow cytometer BD FACS Calibur (BD biosciences, San Jose, CA, USA).

2.11. Cell culture and cytotoxicity assay (*in vitro*)

The human lung fibroblast cell line MRC-5 and the cancerous human lung alveolar epithelial cell line A549 were kindly provided by the cell bank of Pasture Institute of Iran and used as *in vitro* models of normal and cancerous lung cell lines, respectively. They were grown in RPMI/DMEM culture medium, respectively, containing 10% fetal bovine serum in a 5% CO_2 atmosphere and a relative humidity of 90% at 37 °C. The cultured cells were treated with different concentrations of ZnO NPs (10, 50, 100, and 200 $\mu\text{g ml}^{-1}$). The viability of the cells treated with CEN-ZnO NPs was examined by the MTT test. The percentage of cell viability was assessed using the MTT assay using the following protocol. In brief, the cells (1×10^4) were seeded in 96-well culture plates and allowed to adhere for 24 h under a high-humidity environment with 5% CO_2 and 95% O_2 atmosphere at 37 °C. The cells were then exposed to various concentrations (10–200 $\mu\text{g ml}^{-1}$) of CEN-ZnO NPs for 24 h. After exposure, 10 μl of MTT stock solution (5 mg MTT in 1 ml $1 \times$ PBS) was added and the plate incubated for 4 h at 37 °C. Afterward, the solution was removed, and isopropanol added to the cells to create purple crystal formazan. To better dissolve the MTT sediment, the plate was placed on a shaker for 15 minutes. Then, the concentration of the dissolved material in isopropanol was calculated using an ELISA reader (STAT FAX 2100, USA) at a wavelength of 545 nm. The wells with more cells have higher optical density (OD) than the well with lesser cells. Therefore, the following equation was used to determine the wells with more cells and compared with the control sample.

$$\text{Toxicity}\% = 1 - \frac{\text{mean OD of sample}}{\text{mean OD of control}} \times 100$$

$$\text{Viability}\% = 100 - \text{toxicity}\%$$

2.12. ROS level analysis

The ROS level was assayed by a fluorimetric assay using 2',7'-dichlorofluorescein diacetate (DCFHDA; Sigma). The bacterial cells were treated with different concentrations of CEN-ZnO NPs (500, 1000, 1500, and 2500 $\mu\text{g ml}^{-1}$ for *E. coli* and *P. aeruginosa* and 8, 16, 32, and 64 $\mu\text{g ml}^{-1}$ for *S. aureus*) and incubated with



DCFH-DA (10 mM) for 30–60 min at 37 °C and then centrifuged at 4000 rpm. Subsequently, the cells were washed with PBS. The conversion of DCFH to dichlorofluorescein (DCF) was monitored using a fluorescence spectrophotometer with excitation and emission wavelengths at 480 and 520 nm, respectively.

3. Results

3.1. Identification of the selected cyanobacterial isolate

Light microscopy and SEM revealed the isolation, purity, and morphology of the cyanobacterial strain. The selected cyanobacterial isolate forms filamentous structures and is straight or curved to some extent. Molecular analysis of 16S rRNA gene sequences of the strains disclosed that the selected cyanobacterial isolate belongs to the genus of *Nostoc*. The cyanobacterial strain was identified and tentatively designated as *Nostoc* sp. EA03 and the sequence (1260 bp) deposited in GenBank through the accession number of KY947487.1 (Fig. 1a–c).

3.2. GC-MS analysis of cyanobacterial cell extract

GC-MS chromatogram of methanolic extract of *Nostoc* sp. EA03 along with their retention time (RT) is shown in Table 1. The major components present in the methanol extract of *Nostoc* sp. EA03 along with the molecular formula, molecular weight, and peak area are presented in Table 1 and Fig. 2. The GC-MS chromatogram of methanolic extract of *Nostoc* sp. EA03 cell extract showed the presence of several active principle compounds. Fifteen compounds were identified in the methanolic extract of *Nostoc* sp. EA03 cells. The main compounds were *n*-hexadecanoic acid (39.76%), 9-octadecenoic acid (19.72%), octadecadienoic acid (8.67%), hexadecenoic acid (8.11%), octadecanoic acid (4.11%), benzenedicarboxylic acid (2.25%), heptadecane (2.07%), phytol (1.73%), hexadecane (0.77%), and triphenylphosphine oxide (1.21%). Therefore, fatty acids, palmitic acid, and phytol could be the likely candidates elaborated in the capping ZnO at the nanoscale, besides proteins and other ancillary phytochemicals, which needs to be determined in further studies. The existence of algal metabolite hexadecanoic acid (palmitic acid), as a stable and abundant molecule in the cyanobacterial cell extract, can form a dense coating on the surface of the nanoparticles, enhancing its hydrophobic properties.

3.3. Biosynthesis of CEN-ZnO NPs and optimization

The cell extract of *Nostoc* sp. EA03 was added to an aqueous zinc acetate solution, resulting in a change in color from dark green to white (Fig. 3a–d). The UV-Vis absorption spectra of the biosynthesized CEN-ZnO NPs have shown in Fig. 4a. As illustrated, the surface plasmon resonance (SPR) band centered at 370 nm confirms the formation of ZnO NPs in the solution, whereas no absorption band was determined for zinc acetate and the cell extract at the same region. Similar to the result obtained in our study, Jayaseelan *et al.*²⁵ observed a sharp peak in the 374 nm region as a result of ZnONPs synthesis using *A. hydrophila* biomass. Also, our results were completely consistent with a previous report by Singh *et al.*²⁶ that a similar pattern of peaks

obtained from the synthesized zinc oxide nanoparticles using cyanobacterium *Anabaena* strain L3. The effect of pH and cell extract concentration on the UV absorption pattern of CEN-ZnO NPs was also checked. The results showed different concentration of the characteristic band of ZnO NRs at 370 nm as a function of cell extract concentration without any change in the peak wavelength. The maximum production of CEN-ZnO NPs occurred when the final concentration of the cell extract in the reaction mixture was 1 μ l (Fig. 4b). The same investigation on the effect of pH on the yield of CNE-ZnO NPs also showed that the maximum production of ZnO NPs occurred at pH 9, while other pH values showed lower production of CEN-ZnO NPs, as represented in Fig. 4c. Since the synthesis was carried out at room temperature (lack of high-temperature and alkaline requirements), this study appeared to be a significant method for eco-friendly synthesis of ZnO NPs. Based on the results, the amount of fatty acids, especially palmitic acid, is being present in more than the other metabolites in the cell extract. It is therefore suggested that the palmitic acid play an important role as a reducing agent during the green synthesis of the zinc oxide nanoparticles.

3.4. Characterization of CEN-ZnO NPs

3.4.1. XRD analysis. The structure of the synthesized CEN-ZnO NPs was analyzed by XRD analysis (Fig. 4d). Peaks were observed at 2θ values of 31.59°, 34.22°, 36.08°, 47.40°, 56.40°, 62.62°, and 67.75° which can be indexed to (100), (002), (101), (102), (110), (103), and (112). The XRD pattern was comparable with the reference pattern (JCPDS card no. 89-7102) corresponding to the star-like structure of ZnO NPs.²⁶ The observed intense peaks also showed the high purity and crystalline structure of the synthesized ZnO NPs.

3.4.2. FTIR spectroscopy. The chemical structure ZnO and the biosynthesized CEN-ZnO NPs was also studied by FTIR spectroscopy (Fig. 5a). An absorption band, found at 3382 cm^{-1} , indicates the presence of O–H hydrogen bands. The two prominent peaks of proteins were observed at 1650 cm^{-1} (amide I, C=O stretching) and 1550 cm^{-1} (amide II, C–N stretching and N–H deformation). The peak at 1386.49 cm^{-1} represents the N–O functional group. The high intensity band at \sim 547.38 cm^{-1} is the result of the presence of the oxygen and zinc bands. A peak at 1651 cm^{-1} in the biosynthesized CEN-ZnO NPs, by *Nostoc* sp. EA03, is as a result of amide group that is characteristic of the protein. The peak in the region of 1206 cm^{-1} is due to the presence of the C–O functional group. The analysis of cyanobacterial cell extract showed an absorption peak at 1626.96 cm^{-1} that may correspond to free C=O groups, while in CEN-ZnONPs, the absorption peak in the region of 1513 cm^{-1} resulted from the presence of carboxylate group (COO^-). An absorption peak at the region of 2962.78 cm^{-1} indicates the presence of the –C–H stretches of the alkyl (methyl) group. For all doped and undoped ZnO samples the absorption peaks in the range of 400–700 cm^{-1} could be assigned to the ZnO stretching modes. The presence of absorption peaks in the region of 416.14 and 489.98 cm^{-1} in the CEN-ZnONPs sample represents the ZnO core, while this peak



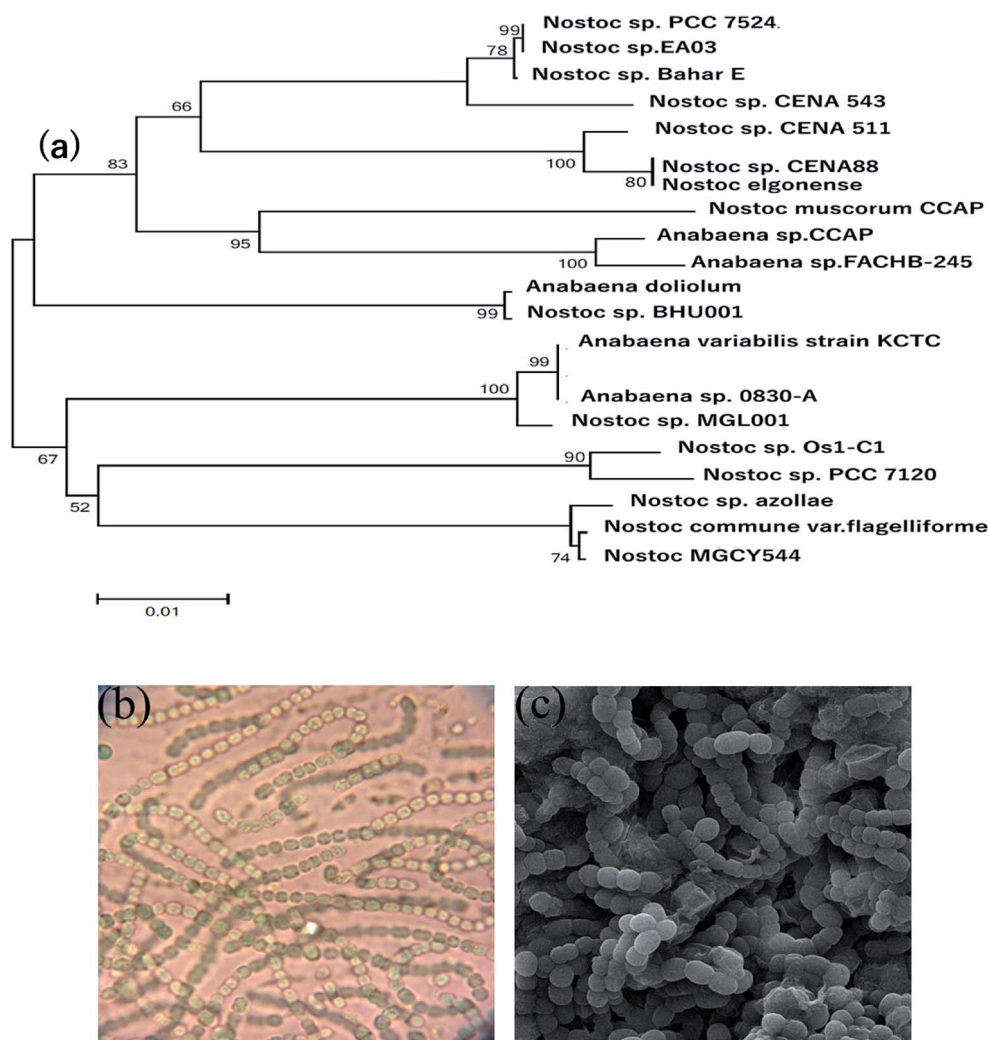


Fig. 1 (a) Phylogenetic tree of the strains *Nostoc* sp. EA03, (b) illustration of the light microscope, (c) scanning electron microscopy (SEM) image of *Nostoc* sp. EA03.

was not observed in cellular extracts and unalloyed ZnO. The absorption peak at 1626 cm^{-1} confirms the presence of C=O groups in the cell extract. The absorption peaks in the region

1352 cm^{-1} and 1044 cm^{-1} in the cell extract, respectively, indicates the presence of C=C and C-O. An absorption peak at 1030 cm^{-1} in CEN-ZnONPs may correspond to the presence of

Table 1 Different compounds of cell extract of the *Nostoc* sp. EA03 and their activities identified by gas chromatography mass spectrometry (GC-MS)

No	RT	Name of the compound (library NIST)	Peak area%	Molecular formula	Compound nature	Activity
1	3.95	Dodecanoyl chloride	1.05	$\text{C}_{12}\text{H}_{23}\text{ClO}$	Saturated fatty acid	No activity reported
2	18.89	<i>n</i> -Hexadecanoic acid	39.76	$\text{C}_{16}\text{H}_{32}\text{O}_2$	Palmitic acid	Antiandrogenic, antioxidant,
3	10.04	Butylated hydroxy toluene	1.81	$\text{C}_{15}\text{H}_{24}\text{O}$	Lipophilic	Antioxidant
4	14.23	Heptadecane	2.07	$\text{C}_{17}\text{H}_{36}$	Alkane	Antioxidant
5	18.46	Hexadecenoic acid	8.11	$\text{C}_{18}\text{H}_{36}\text{O}_2$	Palmitic acid	Antibacterial
6	21.73	Phytol	1.73	$\text{C}_{20}\text{H}_{40}\text{O}$	Diterpene	Antimicrobial, anticancer, anti-inflammatory
7	21.93	9,12-Octadecadienoic acid	8.67	$\text{C}_{20}\text{H}_{38}\text{O}_2$	Oleic acid	Anti-inflammatory, cancer preventive, flavor dermatitogenic
8	22.07	Octadecadienoic acid Oleic acid	19.72	$\text{CH}_3(\text{CH}_2)_{16}\text{COOH}$	Saturated fatty acid	Anticancer
9	27.58	Triphenylphosphine oxide	1.21	$(\text{C}_6\text{H}_5)_3\text{PO}$	Organophosphorus	No activity reported
10	28.50	Benzendicarboxylic acid	2.25	$\text{C}_6\text{H}_4(\text{CO}_2\text{H})_2$	Phthalic acid	Antibacterial, antioxidant



The MIC values of the biosynthesized CEN-ZnO NPs for *E. coli* ATCC 59222, *S. aureus* ATCC 59223, and *P. aeruginosa* PAO1 were 2000, 64, and 2000 $\mu\text{g ml}^{-1}$, respectively, while the MBC values were 2500, 128, and 2500 $\mu\text{g ml}^{-1}$, respectively (Table 2). The maximum MIC and MBC values were 2000 and 2500 $\mu\text{g ml}^{-1}$ for *E. coli* and *P. aeruginosa*, respectively. The biofilm inhibition appeared with increasing concentrations of the biosynthesized CEN-ZnO NPs up to 2500 $\mu\text{g ml}^{-1}$. The rate of biofilm inhibition for the biosynthesized CEN-ZnO NPs concentrations of 128, 256, 500, 1500, and 2500 $\mu\text{g ml}^{-1}$ was 9.49%, 22.15%, 36.30%, 42.67%, and 87.30% in *E. coli*

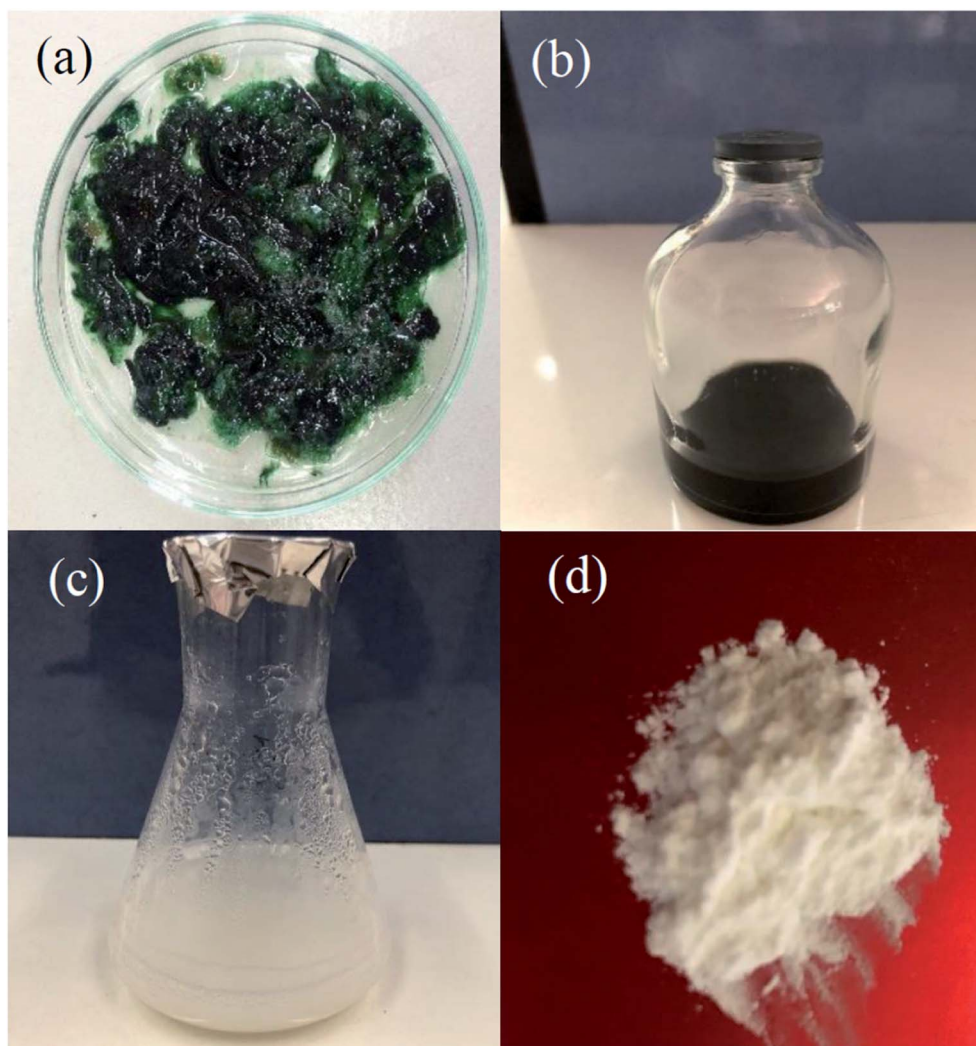


Fig. 3 (a) The biomass of *Nostoc* sp. EA03; (b) purified cell extract; (c) CEN-ZnO NRs; (d) CEN-ZnO powder.

(Fig. 7a), and 51.32%, 53.53%, 54.42%, 55.75%, and 80% in *P. aeruginosa* (Fig. 7b). The rate of biofilm inhibition for *S. aureus* at concentrations of 8, 16, 32, and 64 $\mu\text{g ml}^{-1}$ was 26.35%, 57.36%, 69.11%, and 82.86% (Fig. 7c).

3.6. Eradication of biofilm formation by CEN-ZnO NPs

The biofilm eradication showed that the CEN-ZnO NPs had no significant effect on 3 and 5 days old biofilms, whereas the biosynthesized CEN-ZnO NPs had a significant effect on one day old biofilms. The eradication activity of the NP treatment for *P. aeruginosa* was more pronounced compared to *E. coli* and *S. aureus*. The absorbance value of the biofilm dissolved in 33% (v/v) acetic acid in the microplate at 562 nm was used to evaluate quantitatively the amount of biofilm eradicated from the test material. The results showed that the activity of CEN-ZnO NPs was highest at the concentration of 2 mg ml^{-1} , with an inhibition rate of 88.50% for *P. aeruginosa* (Fig. 7d), and 84.16% for *E. coli* (Fig. 7e). The inhibition rate for *S. aureus*, at a concentration of 64 $\mu\text{g ml}^{-1}$, was 71.53% (Fig. 7f).

3.7. Cell viability

The potential cytotoxic behavior of green synthesized ZnO NPs using *Nostoc* sp. EA03 cell extract against the A549 and MRC-5 cell lines were examined. As shown in Fig. 8a and b, CEN-ZnO NPs do not have any significant effect on the proliferation of A549 and MRC-5 cells at low concentrations; however, when the concentration of CEN-ZnO NPs reached 100 $\mu\text{g ml}^{-1}$, it resulted in a significant decrease in cell viability compared with the control subject. Based on the results, the initial concentration of 50 $\mu\text{g ml}^{-1}$ can be considered as the threshold concentration, at which the toxicity is notably raised ($P \geq 0.05$). However, there were less toxicity, and at a dosage of 50, 100, and 200 $\mu\text{g ml}^{-1}$, the percentage of viable cells was $23\% \pm 06\%$, $13\% \pm 02\%$, and $9\% \pm 44\%$, respectively. The cell extract had no significant effects on the cell viability at 50, 100, and 200 $\mu\text{g ml}^{-1}$ concentrations (Fig. 8a and b). At 50 $\mu\text{g ml}^{-1}$ concentration, CEN-ZnO NPs eliminates 50% of the cancer cells (A549); this concentration kills less than 50% of the healthy cells (MRC-5). The obtained results showed that the CEN-ZnO NPs reduced



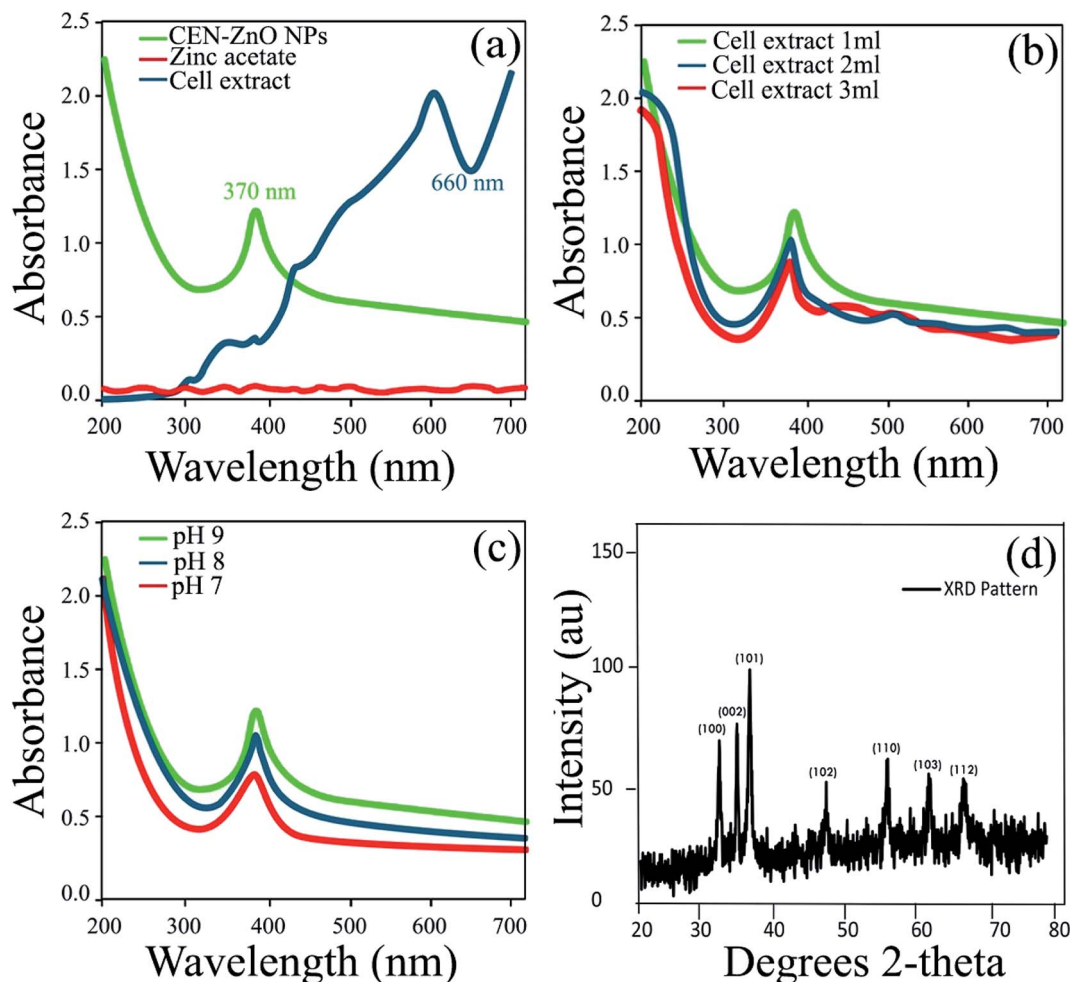


Fig. 4 (a) UV-Vis absorption spectra from 200–700 nm. Green line: CEN-ZnO NPs, blue line: cyanobacterial cell extract, red line: zinc acetate. (b) UV-Vis absorption profiles (200–700 nm) of CEN-ZnO NPs synthesized using different concentrations of cell extract, (c) UV-Vis absorption profiles (200–700 nm) of CEN-ZnO NPs synthesized at different selected pH values, (d) powder X-ray diffraction (XRD) pattern of CEN-ZnO NPs.

the cell viability, and the extent of reduction was concentration/dose-dependent. As determined through the MTT assay, the A549 cell viability observed at 24 h was 85.7%, 57.36%, 23.06%, 13.02%, and 9.44% (Fig. 8a) in contrast, the MRC-5 cell viability observed at 24 h was 94.76%, 77.94%, 58.48%, 9.57%, and 5.44% (Fig. 8b) in response to treatment with 10, 25, 50, 100, and 200 $\mu\text{g ml}^{-1}$ NPs, respectively ($p < 0.05$ for each).

3.8. ROS assay

ROS production upon addition of the biosynthesized CEN-ZnO NPs was determined using the fluorescence dye, DCFH-DA, as an indicator for ROS measurement. In presence of 500–2500 $\mu\text{g ml}^{-1}$ of CEN-ZnO NPs, ROS production was much greater and determined to be 316 ± 4 , 368 ± 8 , 407 ± 8 , and 769 ± 6 in *E. coli*, 410 ± 3 , 550 ± 4 , 616 ± 2 , and 786 ± 9 in *P. aeruginosa*. In presence of 500–2500 $\mu\text{g ml}^{-1}$ of CEN-ZnO NPs, ROS production was much greater and determined to be 316 ± 4 , 368 ± 8 , 407 ± 8 , and 769 ± 6 in *E. coli*, 410 ± 3 , 550 ± 4 , 616 ± 2 , and 786 ± 9 in *P. aeruginosa*. In the case of *S. aureus*, CEN-ZnO NPs at a concentration of 8–64 $\mu\text{g ml}^{-1}$, induced the ROS generation in cells up to 622 ± 1 , 722 ± 66 , $746 \pm$

9, and 768 ± 51 , respectively. These results indicate a significant reduction in the population of viable bacterial cells. The *P. aeruginosa* and *E. coli* cultures showed a higher level of ROS production with increasing CEN-ZnO NPs concentrations, in comparison with that of *S. aureus* (Fig. 8c and d).

3.9. Anti-biofilm activity of CEN-ZnO NPs by confocal laser scanning microscopy

Confocal laser scanning microscopic (CLSM) analysis was performed to evaluate the biofilm formation by bacterial strains *P. aeruginosa* PAO1 (Fig. 9a), and *S. aureus* ATCC 25923 (Fig. 9b). To observe the bacterial biofilms, two dyes, PI and FITC, were used for staining. The microscopic images suggested the formation of biofilm on coverslip surfaces treated with CEN-ZnO NPs. However, most of the cells were dead (red fluorescence), and a small number of cells distinguished by staining with FITC dye (green fluorescence). The bacterial biofilms not treated with CEN-ZnO NPs, most of the bacterial cells surrounded in a network composed of an extracellular matrix (living cells). In the case of *P. aeruginosa* PAO1, the bacterial cells grown in the presence of 2500 $\mu\text{g ml}^{-1}$ of CEN-



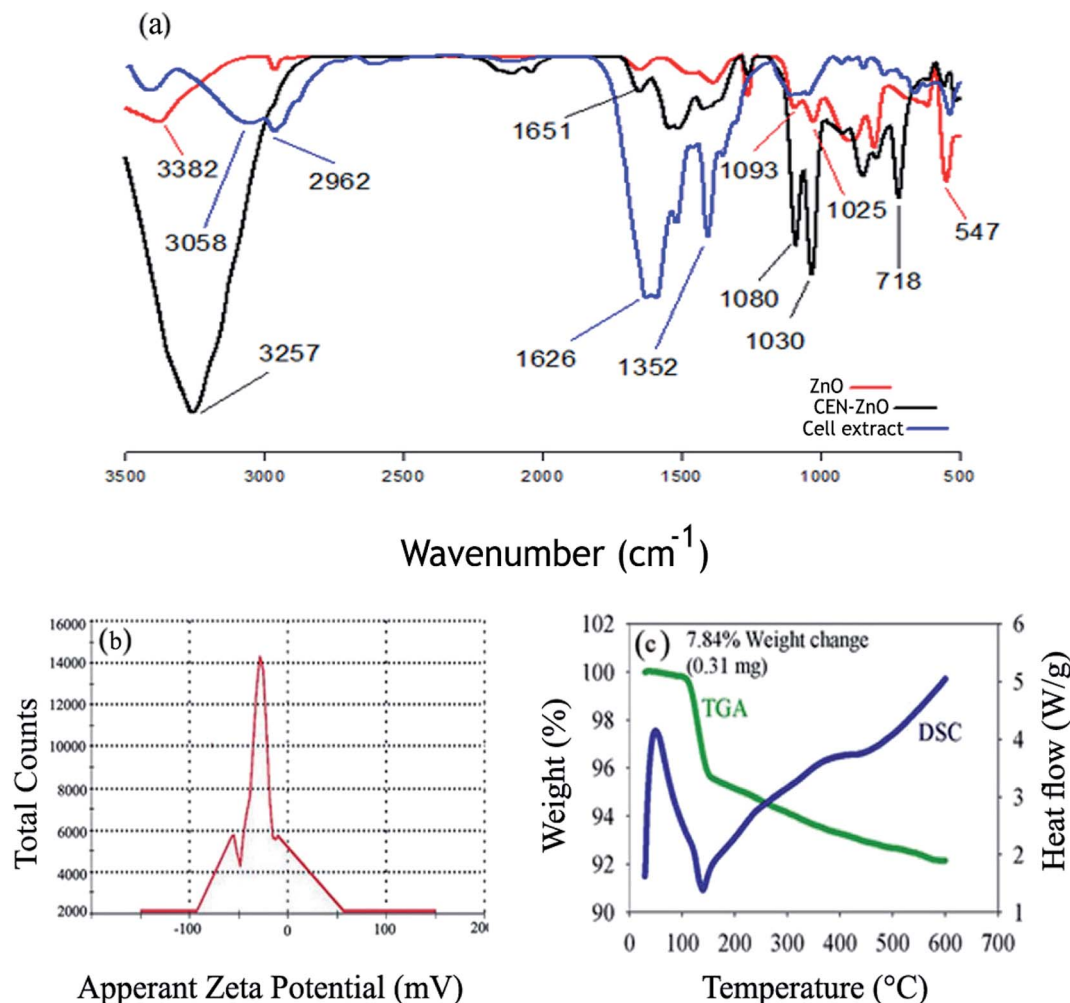


Fig. 5 (a) FTIR profiles of ZnO, CEN-ZnO NPs and cyanobacterial cell extract. (b) Zeta potential distributions for CEN-ZnO NPs. (c) DSC/TGA thermogram profiles of the synthesized CEN-ZnO NPs.

ZnO NPs showed the absence of clumped cells and spread on the surface irregularly. When the CLSM images with red and green fluorescent intensity were placed on top of each other, yellow color appeared. The presence of dark areas among the biofilm might be due to the presence of water channels, heterogeneous matrix production, and all kind of exopolysaccharides inside the biofilms.²⁴ The AFM analysis of surface topography of bacterial biofilm samples treated with CEN-ZnO NPs showed a rugged topography in some parts of the biofilm surface, indicating the destruction of biofilms. AFM 3D topographic images also confirmed the results obtained from the changes in surface topography (Fig. 9c–h). In contrast, in the untreated control samples, the structured biofilms was flat and prominent. Overall, CEN-ZnO NPs cause the destruction of the biofilm, as some parts of the biofilm surface seem to be protruding outward or sunken inwards.

3.10. Flow cytometry analysis

Data were collected and displayed in bi-dimensional FL1-H versus FL2-H cytograms. Fluorescence (FL1-H/FL2-H) was tested from bacterial cells inside gate LL (Fig. 10). As shown, *P.*

aeruginosa PAO1 (Fig. 10a), *E. coli* ATCC 25922 (Fig. 10b), and *S. aureus* ATCC 25923 (Fig. 10c) exhibited a low sensitivity to CEN-ZnO NPs at concentration of 500 µg ml⁻¹ of, and a slow decrease in cell viability (87.01%, 92.04%, and 89.9%, respectively). In fact, less than 10–12% of the bacteria were dead at this concentration. CEN-ZnO NPs at concentrations of 1500 µg ml⁻¹ and 2500 µg ml⁻¹, the cell viability of *P. aeruginosa* PAO1 was reduced by more than 20% and 38%, for *E. coli* by 17.9% and 23.6%, and for *S. aureus* at a concentration of 32 µg ml⁻¹, 64 µg ml⁻¹ by 13% and 16%, respectively. Higher concentrations of CEN-ZnO NPs resulted in an increase in cell debris formation and cell death.

The bacterial cell stained with PI was measured to examine the internalization. The fluorescent emission of this dye was collected in the FL3 channel. The cell internalization of the control group was almost 2.89% for *P. aeruginosa* PAO1 and 6.64% for *S. aureus*, and 3.81% for *E. coli* (Fig. 10). As shown in (Fig. 10d–f), *P. aeruginosa*, *E. coli*, and *S. aureus* showed low sensitivity to CEN-ZnO NPs at concentration of 500 µg ml⁻¹, with a gentle decrease in cell integrity (cell integrity: 3.34%, 9.81%, and 8.11%, respectively). At 1500 µg



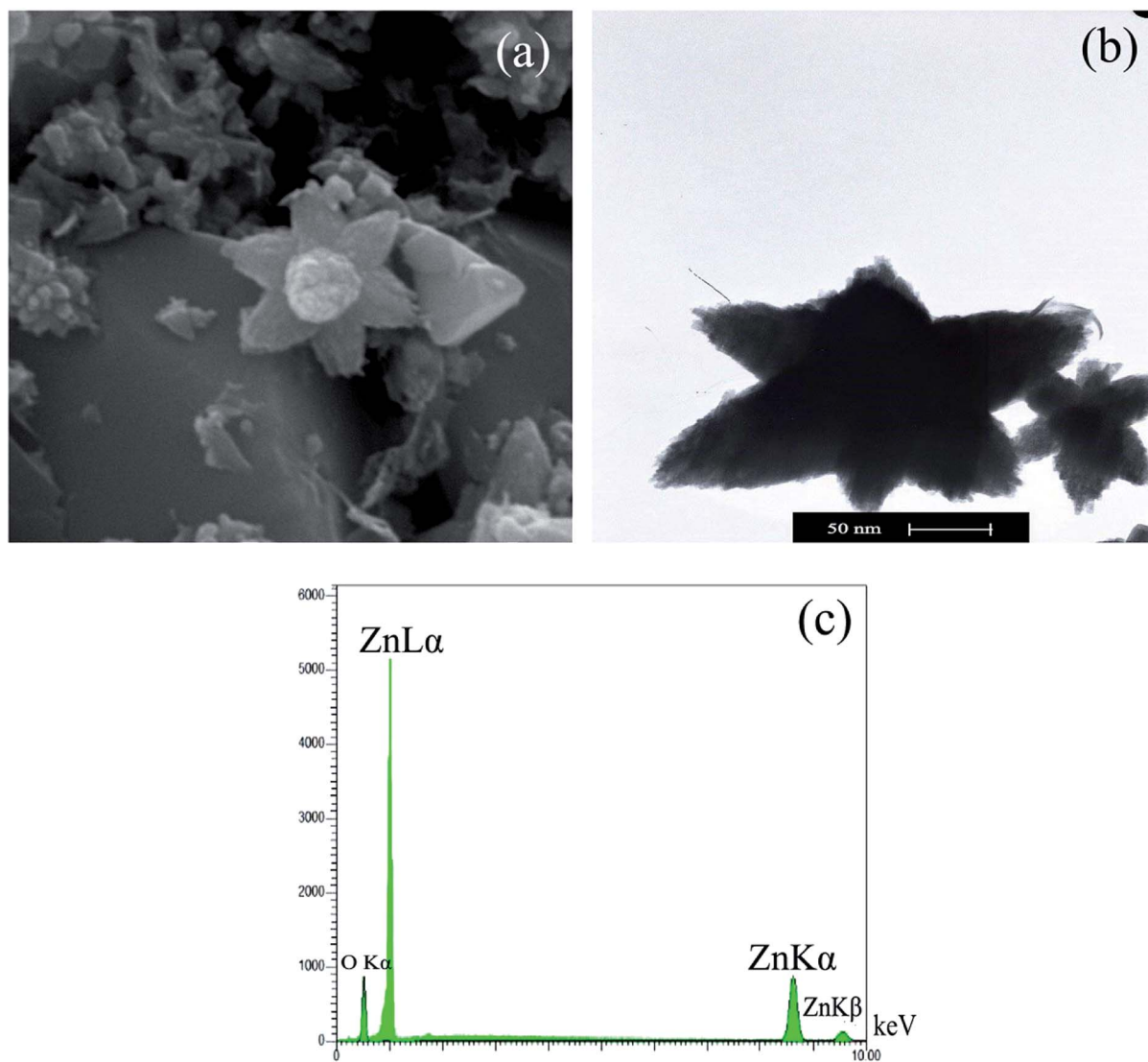


Fig. 6 (a) Scanning electron microscopy (SEM) image of CEN-ZnO NPs; (b) transmission electron microscopy (TEM) image of CEN-ZnO NPs; (c) the analysis of elemental composition of CEN-ZnO NPs recorded by EDX.

ml^{-1} and $2500 \mu\text{g ml}^{-1}$ concentrations, the cell integrity of *P. aeruginosa* PAO1 was 4.54% and 7.92% (Fig. 10d), for *E. coli* 11.9% and 14.1% (Fig. 10e), and for *S. aureus* at a concentration of $32 \mu\text{g ml}^{-1}$, $64 \mu\text{g ml}^{-1}$ was 14.9% and 17.6% (Fig. 10f), respectively.

Table 2 MIC and MBC values against tested bacterial strains according to CLSI protocol

Clinical strains	ZnONPs		CEN-ZnONPs	
	MIC ($\mu\text{g ml}^{-1}$)	MBC ($\mu\text{g ml}^{-1}$)	MIC ($\mu\text{g ml}^{-1}$)	MBC ($\mu\text{g ml}^{-1}$)
<i>E. coli</i> ATCC 25922	6500	6600	2000	2500
<i>S. aureus</i> ATCC 25923	3000	3500	64	128
<i>P. aeruginosa</i> PAO1	6500	6700	2000	2500

4. Discussion

The biosynthesis of CEN-ZnO NPs was environmentally friendly and did not require high temperatures, high pressures, or harmful chemical compounds. The biosynthesized CEN-ZnO NPs had a higher antimicrobial activity in comparison with unalloyed ZnONPs and cyanobacterial cell extract. In a previous report, it was shown that ZnONPs has more toxic effects against Gram-positive bacteria (*S. aureus*) than Gram-negative bacteria (*P. aeruginosa* and *E. coli*).²⁷ They concluded that the difference in the toxicity of nanoparticles might be due to differentiation of the bacterial cell membrane. The results also confirmed that *P. aeruginosa* and *E. coli* are identical in terms of their sensitivity to the equal dose of nanoparticles. The antimicrobial activity of the biosynthesized CEN-ZnO might be due to its star-like shape and its surface-to-volume ratio. In addition, the effectiveness of CEN-ZnO nanoparticles increases with increasing nanoparticle



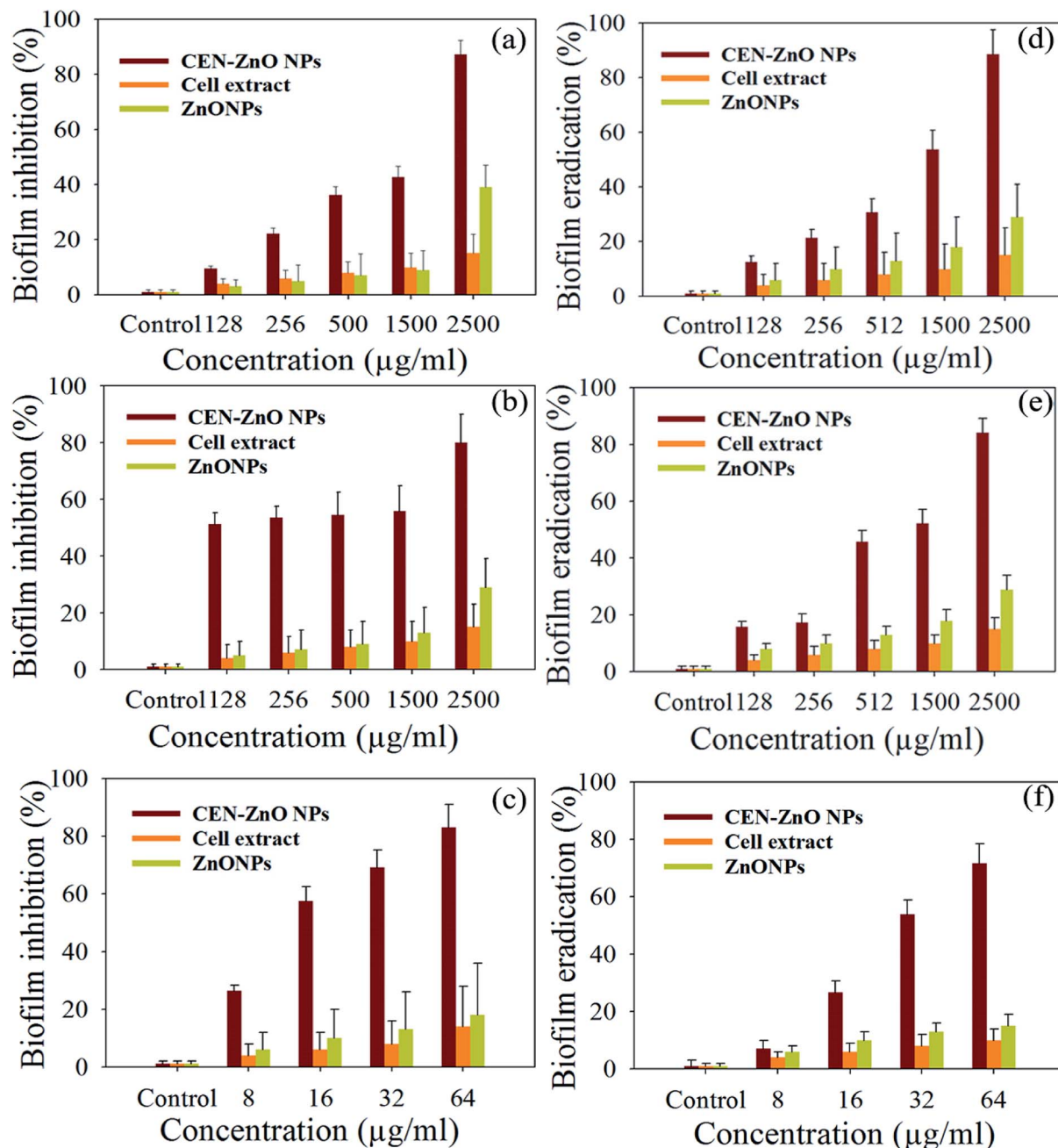


Fig. 7 The inhibitory effects of ZnONPs, cell extract and CEN-ZnO NPs on biofilm formation by (a) *E. coli*, (b) *P. aeruginosa*, (c) *S. aureus*. * $P < 0.05$; bars represent the mean of values and error bars represent mean \pm SD. Inhibitory effect of ZnONPs, cell extract and CEN-ZnO NPs on biofilm eradication by (d) *P. aeruginosa*, (e) *E. coli*, and (f) *S. aureus*. * $P < 0.05$; bars represent the mean of values and error bars represent mean \pm SD.

concentration. Adams *et al.*²⁸ showed that by increasing the concentration of zinc oxide nanoparticles, the antibacterial effects of the nanoparticle against *Bacillus subtilis* and *E. coli* increases. In another study, Khursheed *et al.*²⁹ reported that the antibacterial performance (MIC, MBC) of the biosynthesized zinc oxide nanoparticle using Aloe vera extract against *Pseudomonas aeruginosa*, *E. coli* (ESBL) and MRSA *Staphylococcus aureus* were $2200 \mu\text{g ml}^{-1}$, $2400 \mu\text{g ml}^{-1}$ and $2700 \mu\text{g ml}^{-1}$, respectively. However, as a result of our research, the

biosynthesized CEN-ZnO were reported to have antibacterial activity against *S. aureus* at a lower dose of fewer than $64 \mu\text{g ml}^{-1}$. Concerning *Pseudomonas aeruginosa*, the result was in agreement with our study.²⁹ The higher antibacterial effects of biosynthesized nanoparticles using *Nostoc* sp. EA03 is supposedly due to the existence of the bioactive compounds, making an advantage compared to the unalloyed zinc oxide particles. In addition, the antibacterial behavior of ZnONPs can possibly be attributed to the chemical interaction between hydrogen



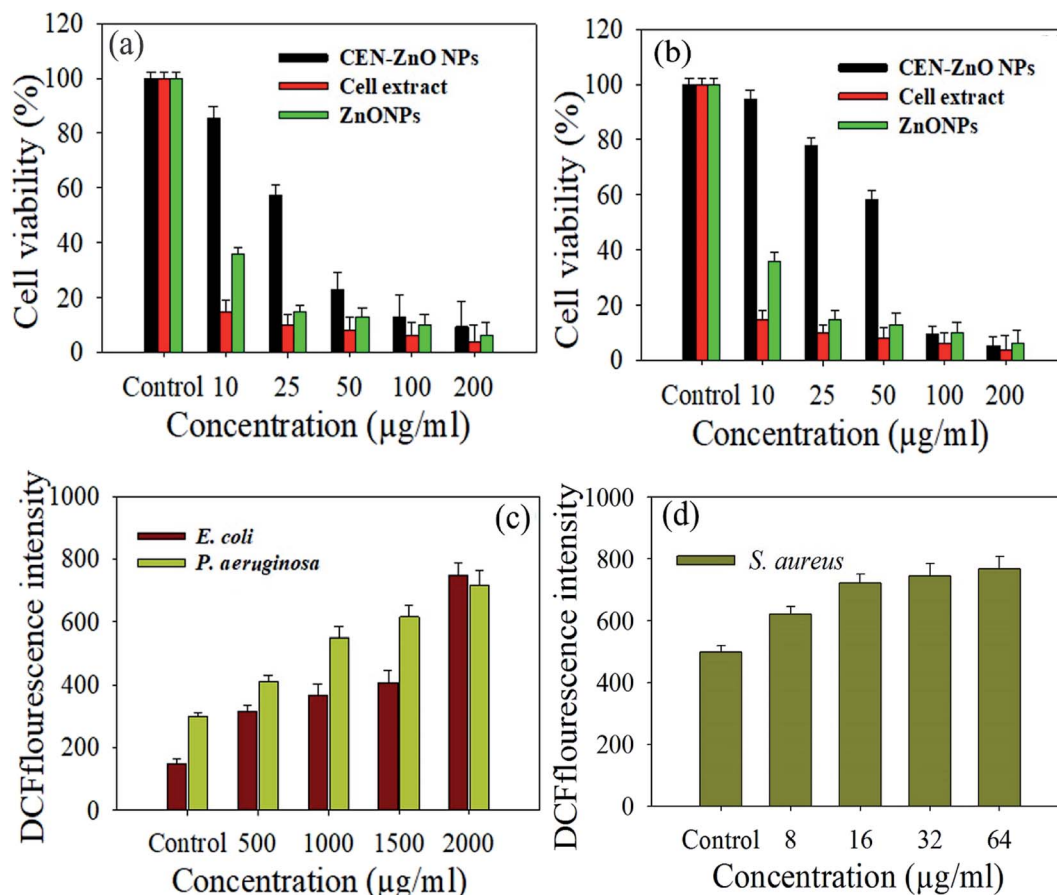


Fig. 8 Histogram representing cell viability based on MTT assay for (a) A549, and (b) MRC-5 cells treated with different concentrations of ZnONPs, CEN-ZnO NPs and cyanobacterial cell extract. Error bars were calculated based on standard deviations of three replications per group ($N = 3$). *In vivo* measurement of reactive oxygen species (ROS) generation induced by indicated concentrations of CEN-ZnO NPs in bacteria cells, (c) *P. aeruginosa*, and *E. coli*, (d) *S. aureus*. Error bars were calculated based on standard deviations of three replications per group ($N = 3$).

peroxide and membrane proteins or lipid bilayer of bacteria. The produced hydrogen peroxide gets through the bacterial cell membrane and destroys the bacterium.³⁰ Rauf *et al.* Reported the ability of antibacterial activity of biomimetically synthesized ZnO nanoparticles against *S. aureus* to induce less-sensitive skin infections in experimental animals. They showed that the synthesized ZnONPs could effectively inhibit less susceptible Gram-positive and Gram-negative bacterial isolates. The synthesized ZnONPs also inhibited the biofilm synthesis of tested isolates *in vitro* and successfully treated the *S. aureus*-mediated experimental skin infection in BALB/c mice.³¹ The penetration of CEN-ZnO NPs into the bacterial cells (in line with the results obtained from flow cytometry analysis), increases intracellular accumulation of nanoparticles, thereby destroying bacterial cells.³² Malaikozhundan *et al.*³³ tested the cytotoxic effects of Pp-ZnO NPs on MCF-7 breast cancer cells and found that single-treatment with Pp-ZnO at concentrations greater than 50 $\mu\text{g ml}^{-1}$ reduces cell viability. The toxic effect of bio-synthesized CEN-ZnO NPs on A549 lung cancer cells starts at 25 $\mu\text{g ml}^{-1}$, has a more toxic effect than ZnONPs biosynthesized by *P. pinnata* seed extract. The biosynthesized CEN-ZnO NPs, at 50 μg , destroyed 77% of cancerous cells (A549) (*i.e.*, 23% of cells

were alive), whereas $\sim 59\%$ of the cells were remained alive after treatment of the healthy lung cells (MRC-5) with a similar concentration. This means that CEN-ZnO NPs, in a similar concentration, eliminate cancerous cells twice as higher than healthy cells. Our results were almost close to those findings that showed a 50% reduction in human lung cancer cells A549 and breast cancer cells MCF-7 after treatment with ZnONPs at a concentration of 31.2 $\mu\text{g ml}^{-1}$.³⁴ Sharma *et al.* (2016) reported that zinc oxide in nanoscale dimensions damage DNA through lipid peroxidation and oxidative stress in human epidermal cells. They showed that the nanostructured materials can be used as a drug carrier, reducing the non-specific toxicity of strong anticancer drugs.³⁵ Earlier studies have described that ZnONPs can reduce cell viability due to their potential for cancer treatment.^{36–38} Our findings indicated that the treatment of bacterial cells with CEN-ZnO NPs results in an elevated ROS production. The treatment of *E. coli* and *P. aeruginosa* with CEN-ZnO NPs showed higher ROS production compared to *S. aureus*, which could be explained by the different potentials that are used to change the polarity of the surface for both Gram-negative and Gram-positive bacteria.^{39,40} The production of ROS results in the disruption of the bacterial cell membrane,



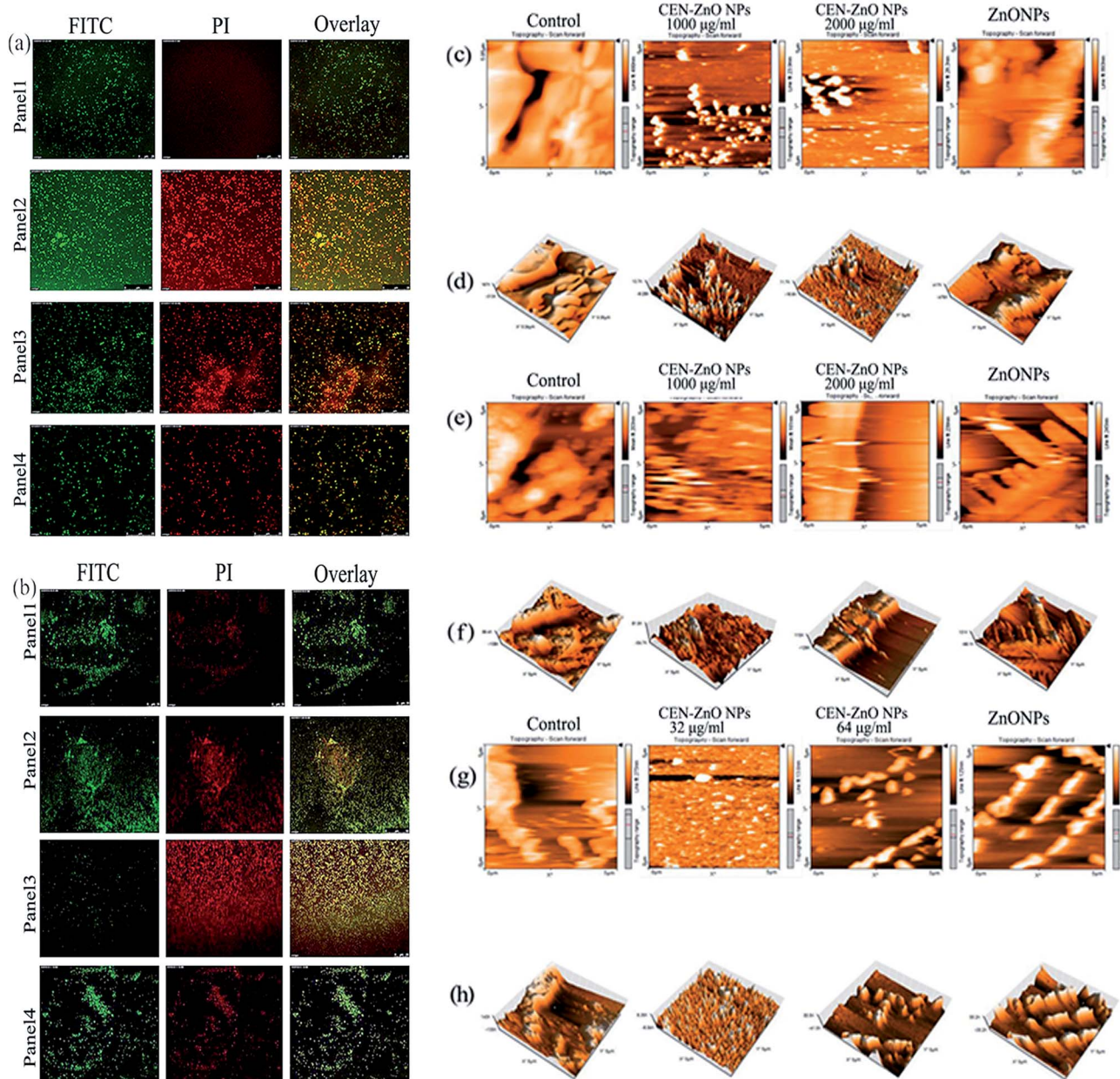


Fig. 9 CLSM micrograph of *P. aeruginosa* (a), *S. aureus* (b) biofilm. Panel 1 from left to right represent CLSM images of a native biofilm (without CEN-ZnONPs). Panel 2 and panel 3 from left to right represent CLSM images of *P. aeruginosa* (a), *S. aureus* (b) biofilms treated with CEN-ZnONPs ($1500 \mu\text{g mL}^{-1}$) and ($2500 \mu\text{g mL}^{-1}$) for *P. aeruginosa* and ($32 \mu\text{g mL}^{-1}$), ($64 \mu\text{g mL}^{-1}$) for *S. aureus*. Panel 4 from left to right show CLSM images of *P. aeruginosa* (a), *S. aureus* (b) biofilm treated with ZnO NPs ($6500 \mu\text{g mL}^{-1}$) for *P. aeruginosa* and ($5000 \mu\text{g mL}^{-1}$) for *S. aureus*. AFM images of (c) *E. coli* topography view and (d) 3D view, (e) *P. aeruginosa* topography view, (f) 3D view, (g) *S. aureus* topography view, and (h) 3D view biofilm. Right images shows native biofilm (no treated with CEN-ZnONPs), whereas, middle images represent AFM images of (c) *E. coli*, (e) *P. aeruginosa* and (g) *S. aureus* biofilms treated with CEN-ZnO NPs. Left images represent AFM images of (c) *E. coli*, (e) *P. aeruginosa* and (g) *S. aureus* biofilm treated with ZnO NPs.

protein denaturation and damage to other macromolecules that ultimately cause cell death.³⁶ In a previous report, Xia *et al.*⁴¹ compared the mechanism of toxicity of zinc oxide and cerium oxide nanoparticles based on their dissolution and oxidative stress features. They concluded that ZnO-induced toxicity in RAW 264.7 and BEAS-2B cell lines leads to the generation of reactive oxygen species (ROS), oxidant injury, excitation of inflammation, and cell death. In contrast, CeO (2) suppressed

ROS production and induced cellular resistance to an exogenous source of oxidative stress. Guo *et al.*⁴² also investigated the role of oxidative stress in toxicity and possible contribution of mitochondria in the production of reactive oxygen species (ROS) upon exposure of retinal ganglion cells (RGC-5) to ZnO nanoparticles. They found that ZnO nanoparticles decrease the mitochondrial membrane potential, increase the production of ROS, leading to the RGC-5 cell damage and induction of the



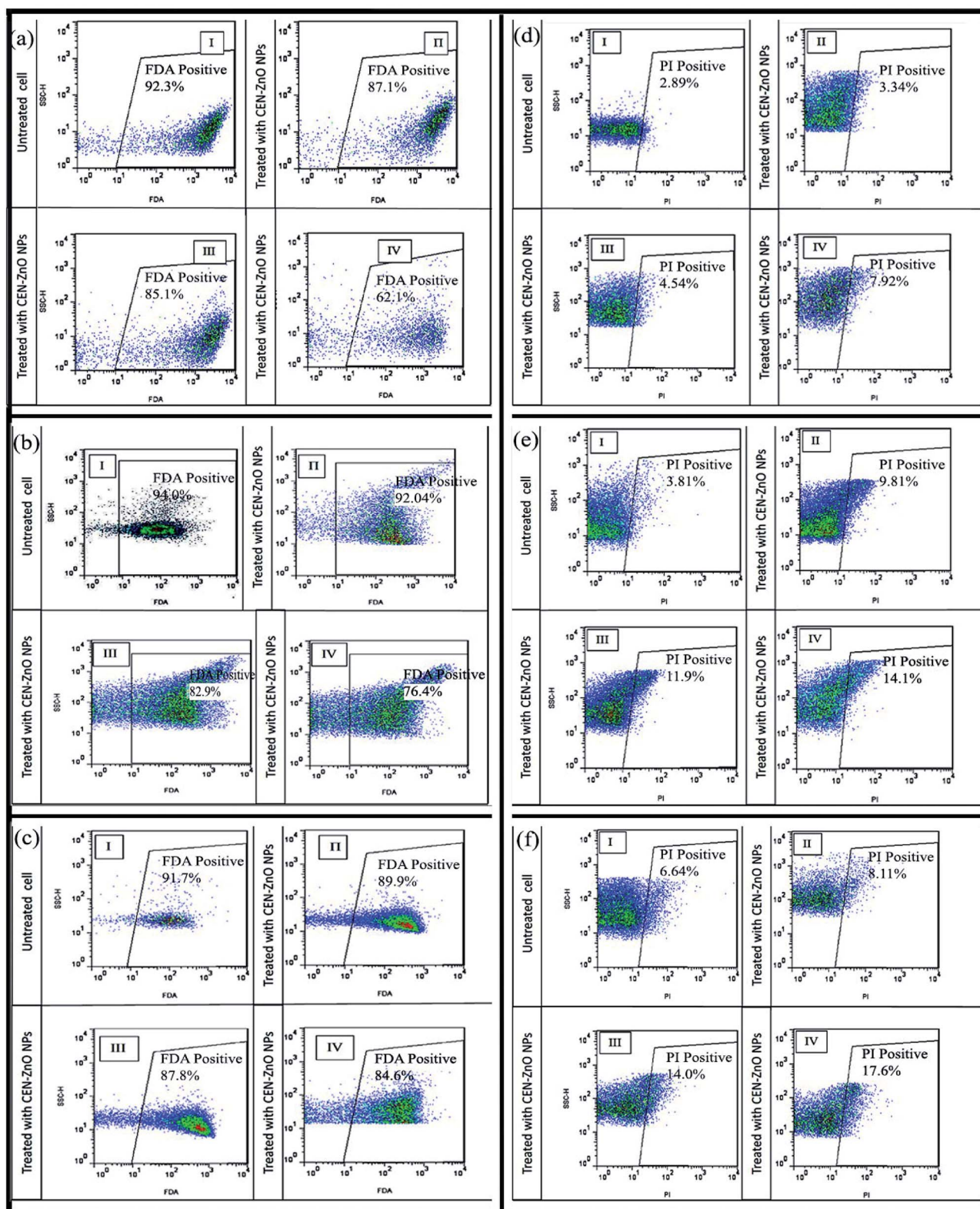


Fig. 10 Flow cytometric analysis of (a) *P. aeruginosa*, (b) *E. coli* and (c) *S. aureus*. The cell viability level as indicated by fluorescein diacetate (FDA) staining (a) and (b) cytograms I: control (untreated sample); II: treated with CEN-ZnO NPs (500 $\mu\text{g mL}^{-1}$), III: (1500 $\mu\text{g mL}^{-1}$) and IV: (2500 $\mu\text{g mL}^{-1}$). (c) Cytograms I: control (untreated sample); II: *S. aureus* treated with CEN-ZnO NPs (16 $\mu\text{g mL}^{-1}$), III: (32 $\mu\text{g mL}^{-1}$) and IV: (64 $\mu\text{g mL}^{-1}$). Flow cytometric analysis of (d) *P. aeruginosa*, (e) *E. coli* and (f) *S. aureus*. Membrane internalization staining using prodium iodide (PI). (a) and (b) Cytograms I: control (untreated sample); II: treated with CEN-ZnO NPs (500 $\mu\text{g mL}^{-1}$), III: (1500 $\mu\text{g mL}^{-1}$) and IV: (2500 $\mu\text{g mL}^{-1}$). (c) Cytograms I: control (untreated sample); II: *S. aureus* treated with CEN-ZnO NPs (16 $\mu\text{g mL}^{-1}$), III: (32 $\mu\text{g mL}^{-1}$) and IV: (64 $\mu\text{g mL}^{-1}$).



apoptosis/necrosis. Liu *et al.*⁴³ investigated the cytotoxicity of two types of ZnO NPs of various sizes and indicate that the reactive oxygen species (ROS) played an essential role in ZnO NP-induced cytotoxicity, possibly due to the leaching of zinc ions from ZnO NPs. They showed that ZnO NPs induce toxic responses in the human neuroblastoma SHSY5Y cell line in a size-dependent manner, which might be related to their ion-shedding ability.

The results obtained from flow cytometry analysis showed that the cell viability decreases with increase in CEN-ZnO NP concentration. Unlike the results obtained here, Bao *et al.*⁴⁴ used PI dye to analyze the dead fractions of *E. coli* bacterial cells after treatment with 5 $\mu\text{g ml}^{-1}$ and 10 $\mu\text{g ml}^{-1}$ AgNPs and found that the PI-positive percentages of the bacterial cells were much higher than control groups. However, this increasing PI positive percentage is not correlated to the concentration of the AgNPs, and the highest positive ratio of the bacterial cells detected in the 5 $\mu\text{g ml}^{-1}$ group.⁴⁴ Accurate measurement of nanoparticle entering is important for measuring the physical and chemical properties of nanoparticle uptake into target cells. Also, by increasing the concentration of CEN-ZnO NPs at a fixed time, the penetration rate of nanoparticles into the cell wall of the bacteria increases. It can be speculated that Zn^{2+} ion release from zinc oxide may result in the transfer of Zn^{2+} ion into the cell wall, leading to bacteriostatic effects. For example, the interaction between the cell coating components and the Zn^{2+} ion (from ZnO), as reported before, was consistent with the results obtained in our research.⁴⁵ The evidence obtained from confocal laser scanning microscopy (CLSM) analysis showed that while exopolysaccharides with green fluorescence were not detected, many dead cells were observable after treatment of biofilm samples with CEN-ZnO NPs. Since there are water channels (hole) to transport nutrients in biofilms, nanoparticles may penetrate directly through the exopolysaccharide layer.²³ ZnO nanoparticles (ZNPs) are considered as promising inorganic nanoparticles for use in catalysis, sunscreens, cosmetics, coatings, pigments, and food additives. However, there are ever more concerns for scientists regarding their environmental threats. For example, Li *et al.*⁴⁶ studied the uptake and depuration performance of *n*ZnO and Zn ions in a freshwater cladoceran *Daphnia magna*. They found that the uptake of radio-synthesized *n*ZnO by *D. magna* is related to *n*ZnO concentration. The *n*ZnO could enter the body of daphnids and reached the highest level within a short period of time, followed by a quick release. The depuration of *n*ZnO was quick and controlled by *n*ZnO dissolution. They finally suggested that both dissolved Zn and nanoparticles had better to be considered essential for studying the toxicology of *n*ZnO. Rajput *et al.*⁴⁷ investigated the possible threats caused by toxicity of various NPs on plants and microbial diversity. The exposure of soil to nanoparticles led to a decrease in soil microbial biomass and enzymatic activity, affecting microbial community composition including yeasts, bacteria, fungi, and biological diversity. They concluded that it is expected to divulge whether NPs could compromise crop yield, soil properties, soil organisms, and functional activities of soil.

5. Conclusion

The use of cyanobacteria has provided the basis for the development of green nanotechnology, thus reducing the production of unnecessary and impure chemical substances. The present study gives new insight into the use of cyanobacterial cell extracts, as a reducing and capping agent in the synthesis of zinc oxide nanoparticles. Our results showed that the bio-synthesized zinc oxide nanoparticles have a favorable biological potential through antibacterial, anti-biofilm, and anticancer activities. The current study has also made significant findings concerning the capacity of the biosynthesized nanoparticles to destroy the lung cancer cell. The number of normal cells was approximately two-fold higher than that of cancer cells upon exposure to CEN-ZnO NPs, at a concentration of 50 $\mu\text{g ml}^{-1}$. In addition to anti-cancer activity, the CEN-ZnO NPs was able to eliminate 24 hour-old biofilms, up to 80 to 90%, in tested bacterial strains.

Altogether, zinc oxide nanoparticles synthesized using the cyanobacterium *Nostoc* sp. EA03 can be a good candidate for cancer treatment as well as a favorable alternative to common antibiotics for controlling microbial pathogens.

Conflicts of interest

There are no conflict to declare.

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