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A green chemistry approach for synthesizing biocompatible gold nanoparticles

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Abstract

Gold nanoparticles (AuNPs) are a fascinating class of nanomaterial that can be used for a wide range of biomedical applications, including bio-imaging, lateral flow assays, environmental detection and purification, data storage, drug delivery, biomarkers, catalysis, chemical sensors, and DNA detection. Biological synthesis of nanoparticles appears to be simple, cost-effective, non-toxic, and easy to use for controlling size, shape, and stability, which is unlike the chemically synthesized nanoparticles. The aim of this study was to synthesize homogeneous AuNPs using pharmaceutically important *Ganoderma* spp. We developed a simple, non-toxic, and green method for water-soluble AuNP synthesis by treating gold (III) chloride trihydrate (HAuCl₄) with a hot aqueous extract of the *Ganoderma* spp. mycelia. The formation of biologically synthesized AuNPs (bio-AuNPs) was characterized by ultraviolet (UV)-visible absorption spectroscopy, X-ray diffraction (XRD), Fourier transform infrared spectroscopy (FTIR), energy dispersive X-ray (EDX), dynamic light scattering (DLS), and transmission electron microscopy (TEM). Furthermore, the biocompatibility of as-prepared AuNPs was evaluated using a series of assays, such as cell viability, lactate dehydrogenase leakage, and reactive oxygen species generation (ROS) in human breast cancer cells (MDA-MB-231). The color change of the solution from yellow to reddish pink and strong surface plasmon resonance were observed at 520 nm using UV-visible spectroscopy, and that indicated the formation of AuNPs. DLS analysis revealed the size distribution of AuNPs in liquid solution, and the average size of AuNPs was 20 nm. The size and morphology of AuNPs were investigated using TEM. The biocompatibility effect of as-prepared AuNPs was investigated in MDA-MB-231 breast cancer cells by using various concentrations of AuNPs (10 to 100 μM) for 24 h. Our findings suggest that AuNPs are non-cytotoxic and biocompatible. To the best of our knowledge, this is the first report to describe the synthesis of monodispersed, biocompatible, and soluble AuNPs with an average size of 20 nm using *Ganoderma* spp. This study opens up new possibilities of using an inexpensive and non-toxic mushroom extract as a reducing and stabilizing agent for the synthesis of size-controlled, large-scale, biocompatible, and monodispersed AuNPs, which may have future diagnostic and therapeutic applications.

Keywords: AuNPs; Biocompatibility; *Ganoderma* spp; Human breast cancer cells; Transmission electron microscopy; UV-visible spectroscopy

Background

In recent years, gold nanoparticles (AuNPs) have been of great research interest because of their unique properties, such as size- and shape-dependent optoelectronic, physiochemical, and biological properties as well as various potential therapeutic applications. AuNPs possess distinct physical and chemical properties that make them excellent tools for creating novel chemical and biological

sensors [1-3]. First, AuNPs can be synthesized using a simple method and made highly stable. Second, they possess unique optoelectronic properties. Third, they provide high surface-to-volume ratios with excellent biocompatibility when using appropriate ligands [1]. Fourth, these AuNP properties can be readily tuned by varying their size and shape as well as the surrounding chemical environment [3].

Because of their stability, oxidation resistance, and biocompatibility, AuNPs have a wide range of potential applications, such as in electronics and photonics, catalysis, information storage, chemical sensing and imaging, drug

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delivery, and biological labeling [4,5]. The tuning of AuNPs is an important process to enhance versatility in defining and controlling the shape [5]. Thus, new methodologies are essential for designing shape-controlled synthesis of AuNPs [6-8].

Several synthetic chemical methods have been adopted for AuNP synthesis, including physical methods, such as attrition and pyrolysis, which were previously utilized for the synthesis of metallic nanoparticles [9]. Alternatively, chemical methods are the most widely and traditionally used methods and incorporate various reducing agents, such as hydrazine [9] and sodium borohydride [10]. However, many of these methods can be cumbersome and involve the use of toxic chemicals, high temperatures, and pressures and, most importantly, can cause the particles to become unstable or aggregate upon interaction with biological media or biomolecules [11]. At the same time, these approaches produce multi-shaped nanoparticles that require purification by differential centrifugation, and consequently have a low yield [12,13]. Therefore, there is a constant need for new methodologies to produce shape-controlled synthesis of AuNPs.

Recently, several labs have been interested in developing methodologies for synthesis of nanomaterials using a green chemistry approach, which is an alternate approach to biosynthesizing nanomaterials that relies on natural organisms for the reduction of metal ions into stable nanocrystals [14-21]. Biological methods are supposed to yield novel and complex structural entities, unlike those obtained using conventional techniques [14,15,22]. A number of microbial species have been used for synthesis of metal nanoparticles but without much success in achieving shape control. The shape-controlled microbial synthesis of nanostructures is an exciting new area with considerable potential for development. Recently, Das et al. reported the synthesis of single-crystalline AuNPs [19] and different nanostructures from HAuCl₄ using *Rhizopus oryzae* [5].

Biological methods exhibit size and shape control over a diverse array of materials, and they also facilitate mass production, high yield, and reproducibility [23,24]. Biosynthesis of AuNPs and silver nanoparticles (AgNPs) have been reported in different prokaryotic organisms, including *Bacillus licheniformis* [20], *Brevibacterium casei* [21], *Bacillus subtilis* [25], *Escherichia coli* [26], *Lactobacillus* [27], *Pseudomonas aeruginosa* [28], and *Rhodospseudomonas capsulate* [29]. Several researchers exploited fungi as reducing agents for AgNP synthesis, including fungi such as *Verticillium* [14], *Fusarium oxysporum* [16], *Aspergillus fumigatus* [30], *Penicillium fellutanum* [31], *Volvariella volvacea* [32], *Pleurotus florida* [33], *Candida* [34], *Ganoderma lucidum* [35], and *Neurospora crassa* [36].

Among nanoparticles, AuNPs have immense potential for cancer diagnosis and therapy. Conjugation of AuNPs

to ligands on cancer cells allows molecular imaging and detection of cancer [37]. Further, AuNPs have potential applications in electronics, catalysis, biological sensors, cancer diagnostics, therapeutics, nanomedicine, and environmental work, because they have several merits, such as the fact that they are easy to synthesize, cost effective, and non-toxic, and they have easy functionalization, optical properties, facile surface chemistry, and biocompatibility [37,38]. Moreover, biological processes could provide significant yield and are free from downstream processing; therefore, many researchers are interested in synthesizing nanoparticles with green manufacturing technology that uses bacteria, fungi, plants, and plant products.

In most studies, either AuNPs or AgNPs were synthesized using bacteria. Many fungi have not been explored, including those mentioned above, and only a few fungi have been investigated for AuNP and AgNP synthesis. Among fungi that have not been tested, *Ganoderma* spp. have long been used as medicinal mushrooms in Asia, and they have an array of pharmacological properties, including immunomodulatory activity and pharmacological properties [39]. *Ganoderma* spp. have several advantages: They have large quantities of viable mycelia, they are cost-effective, there is easy downstream processing, and they are non-pathogenic. Moreover, the synthesized AuNPs are highly soluble in water. Therefore, the aim of this study was to investigate the possible use of *Ganoderma* spp. as green producers for AuNP synthesis and to further evaluate the biocompatibility effect of as-prepared AuNPs in human breast cancer cells (MDA-MB-231).

Methods

Reagents

Gold (III) chloride trihydrate was purchased from Sigma (St. Louis, MO, USA). Penicillin-streptomycin solution, trypsin-EDTA solution, Dulbecco's modified Eagle's medium (DMEM/F-12), and 1% antibiotic-antimycotic solution were obtained from Life Technologies GIBCO (Grand Island, NY, USA). All the other chemicals and reagents were purchased from Sigma (St. Louis, MO, USA), unless otherwise specified.

Culturing and maintenance of *Ganoderma* spp.

The culture of *Ganoderma* spp. was collected from a tropical forest near Pollachi, Tamilnadu, India. Culturing and maintenance were conducted as described in previous studies, with suitable modifications [40,41]. Briefly, the mycelia were cultured on potato dextrose agar (PDA) and incubated at 28°C ± 2°C for 7 days. The mycelia were then transferred to glucose yeast malt peptone broth (GYMP). The inoculated medium was incubated at 28°C ± 2°C and agitated at 150 rpm for 10 days. After incubation, the mycelia were harvested, washed with distilled water,

freeze-dried, and stored at 4°C in air-tight containers, prior to use.

Preparation of mycelia hot aqueous extract

The preparation of mushroom extract was carried out according to a method described in previous studies [40,41], with suitable modifications. In brief, the freeze-dried mycelia were soaked in distilled water at a ratio of 1:20 and double boiled for 45 min, left to cool, and filtered through Whatman filter paper No. 4. The hot aqueous extract was then freeze-dried at $-70^{\circ}\text{C} \pm 2^{\circ}\text{C}$ for 48 h and stored at 4°C in airtight containers. The freeze-dried hot aqueous extract of the mycelia was used as the reducing and stabilizing agent for AuNP synthesis.

Synthesis of AuNPs

Synthesis of AuNPs was carried out according to the method described earlier [21]. In a typical reaction, 1 mg/mL of freeze-dried hot aqueous mushroom mycelia extract was mixed with an aqueous solution of 1 mM HAuCl_4 solution and kept at room temperature for 24 h. Synthesis was observed using ultraviolet (UV)-visible spectroscopy. The color change observed was from pale yellow to purple. To compare the efficiency of biologically prepared AuNPs, we used citrate-mediated synthesis of AuNPs (chem-AuNPs) from Sigma.

Characterization of AuNPs

Characterization of synthesized AuNPs was carried out according to previously described methods [20]. The nanoparticles were primarily characterized by UV-visible spectroscopy, which has proven to be a very useful technique for nanoparticle analysis [26]. UV-visible (UV-vis) spectra were obtained using a WPA (Biowave II, Biochrom, Cambridge, UK). Dispersion particle size was measured by Zetasizer Nano ZS90 (Malvern Instruments Limited, Malvern, UK).

The synthesized AuNPs were freeze dried, powdered, and used for X-ray diffraction (XRD) analysis. The spectra were evaluated using an X-ray diffractometer (PHILIPS X'Pert-MPD diffractometer, Amsterdam, the Netherlands) and $\text{Cu-K}\alpha$ radiation (1.5405 \AA) over an angular range of 10° to 80° at 40 kV and 30 mA. The dried powder was diluted with potassium bromide at a the ratio of 1:100, and the results were recorded using the Fourier transform infrared spectroscopy (FTIR; PerkinElmer Inc., Walham, MA, USA) and spectrum GX spectrometry within the range of 500 to $4,000 \text{ cm}^{-1}$.

Transmission electron microscopy (TEM, JEM-1200EX, JEOL Ltd., Tokyo, Japan) was used to determine the size and morphology of AuNPs. AuNPs were prepared by dropping a small amount of aqueous dispersion on copper grids, which were dried and then examined in the TEM. Further, the presence of Au metals in the sample was

analyzed by energy dispersive X-ray analysis (EDX) combined with a field emission SEM.

Cell culture

MDA-MB-231 human breast cancer cells were kindly provided by Kyung Jin Lee, Institute for Life Sciences, ASAN Medical Center, University of Ulsan College of Medicine. MDA-MB-231 breast cancer cell lines were grown adherently and maintained in DMEM containing 10% fetal calf serum (FCS) and 1% antibiotic solution containing penicillin and streptomycin at 37°C under 5% CO_2 . All the experiments were performed in six-well plates, unless stated otherwise. Cells were seeded onto plates at a density of 1×10^6 cells per well and incubated for 24 h prior to the experiments. The cells were washed with phosphate buffered saline (PBS, pH 7.4) and incubated in fresh medium containing different concentrations of AuNPs dissolved in water.

Cell viability assay

In order to evaluate the biocompatibility of the as-prepared AuNPs, we carried out cell viability assay in breast cancer cells (MDA-MB-231) by using MTT reagents. In addition, to compare the biocompatibility effect of bio-AuNPs, we used chemical-mediated synthesis of chem-AuNPs as a positive control. Cell viability was measured using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide dye reduction assay performed to determine the cytotoxic effect of the AuNPs at various concentrations. Briefly, the cells were plated onto 96-well flat-bottom culture plates with various concentrations of AuNPs (0 to $100 \mu\text{M}$). All the cultures were incubated for 24 h at 37°C in a humidified incubator. After 24 h of incubation (37°C , 5% CO_2 in a humid atmosphere), $10 \mu\text{L}$ of MTT (5 mg/mL in PBS) was added to each well, and the plate was incubated for another 4 h at 37°C . The resulting formazan was dissolved in $100 \mu\text{L}$ of DMSO with gentle shaking at 37°C , and the absorbance was measured at 595 nm by using an ELISA reader (Spectra MAX; Molecular Devices, Sunnyvale, CA, USA). The results are shown as the mean of three independent experiments.

Membrane integrity

Cell membrane integrity of MDA-MB-231 cells was evaluated by determining the activity of lactate dehydrogenase (LDH) leaking out of the cell, according to the manufacturer's instructions (*in vitro* toxicology assay kit, TOX7, Sigma, USA). The LDH assay is based on the release of the cytosolic enzyme LDH from cells with damaged cellular membranes. Thus, in cell culture, AuNPs induced cytotoxicity and were quantitatively analyzed by measuring the activity of LDH in the supernatant. Briefly, cells were exposed to various concentrations of AuNPs for 24 h, and then $100 \mu\text{L}$ per well of each cell-free supernatant was

transferred in triplicates into wells in a 96-well plate, and 100 μ L of LDH-assay reaction mixture was added to each well. After 3-h incubation under standard conditions, the optical density of the generated color was determined at a wavelength of 490 nm using a Microplate Reader.

Determination of ROS

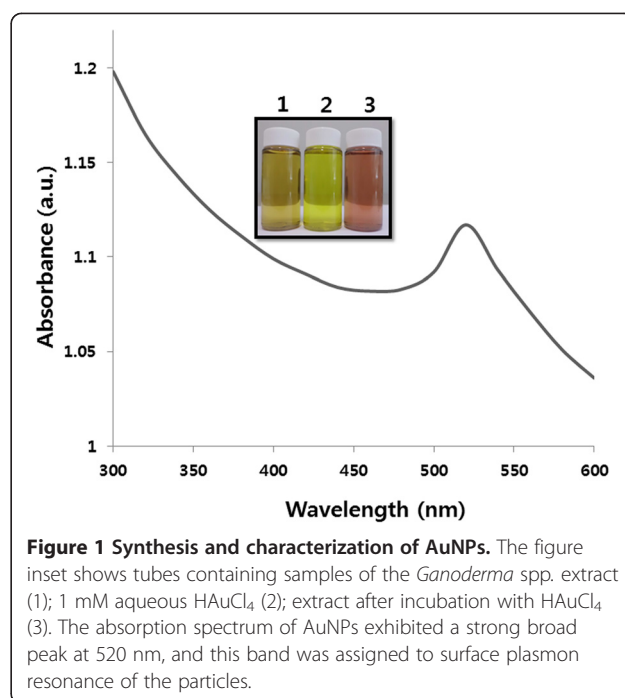
Intracellular reactive oxygen species (ROS) were measured based on the intracellular peroxide-dependent oxidation of 2',7'-dichlorodihydrofluorescein diacetate (DCFH-DA, Molecular Probes, Eugene, OR, USA) to form the fluorescent compound 2',7'-dichlorofluorescein (DCF), as previously described. Cells were seeded onto 24-well plates at a density of 5×10^4 cells per well and cultured for 24 h. After washing twice with PBS, fresh medium containing 100 μ M of AuNPs, 1 mM H_2O_2 , or AgNPs (5 μ g/mL) was added, and the cells were incubated for 24 h. For the control, cells were added to 20 μ M of DCFH-DA and incubation continued for 30 min at 37°C. The cells were rinsed with PBS, 2 mL of PBS was added to each well, and fluorescence intensity was determined with a spectrofluorometer (Gemini EM) with excitation at 485 nm and emission at 530 nm. For the control, had antioxidant *N*-acetyl-L-cystein (NAC, 5 mM) was added to the cells grown in 24-well plates (for 24 h) for 1 h prior to exposure to AuNPs, 1 mM H_2O_2 , or AgNPs (5 μ g/mL) for 24 h. We then added 20 μ M of DCFH-DA, and the cells were incubated for 30 min at 37°C before measuring DCF fluorescence changes as described.

Results and discussion

Extracellular synthesis of AuNPs

Primary characterization of the ability of *Ganoderma* spp. mushroom extract for AuNP synthesis was analyzed. The Figure 1 inset shows tubes with the *Ganoderma* spp. mycelia extract [1], $H AuCl_4$ [2], and extract after reaction with $H AuCl_4$ ions for 24 h [3]. As expected, the color changed from pale yellow to deep purple in the presence of the extract, which indicates AuNP formation and is evidence of synthesis.

Further, to determine if there was AuNP synthesis, the biosynthesis of nanoparticle reactions was monitored by UV-vis spectroscopy. UV-vis absorption spectroscopy is the most widely used technique for characterizing the optical properties and electronic structure of nanoparticles, because the absorption bands are related to the diameter and different aspect ratios of metal nanoparticles, including size and shape [42]. As shown in Figure 1, the spectra of AuNP synthesis showed a gradual increase in the surface plasmon resonance (SPR) excitation peak centered at 520 nm, which is characteristic of AuNPs [11,43]. This further indicates that the mushroom extract could be useful as a reducing agent for AuNP synthesis. Control reactions in the absence of mushroom extract



exhibited no change in color or absorbance at 520 nm, clearly indicating that the protein and polysaccharides found in the extract are responsible for biosynthesis of AuNPs.

Previous studies demonstrated that metal biotransformation might involve a complex of either capping proteins/peptides and reductases, quinines, cytochromes, phytochelatins, or electron shuttles that are known to reduce various metals and metal oxides [11,43-46]. Das et al. [47] proposed possible mechanisms of AuNP synthesis in *Rhizopus oryzae*. The first mechanism is binding of Au (III) on the cell wall through electrostatic interaction followed by reduction to AuNPs by proteins/enzymes present on the cell wall, and the second is diffusion or transportation of Au (III) into the cytoplasm and protein/enzymatic reduction to form AuNPs. Taken together, these results indicate that AuNP synthesis could be facilitated by the presence of proteins in the extract.

XRD analysis of AuNPs

The crystalline nature of as-prepared AuNPs was confirmed using XRD. The XRD spectrum shows two predominant peaks that agree with Bragg's reflection of AuNPs reported in a previous study, which used extracellular and intracellular culture supernatant of *Aspergillus fumigatus* and *Aspergillus flavus* [48]. The diffraction peaks, which appeared at 31.6°C and 45.4°C corresponded to the (111) and (200) planes, respectively (Figure 2). No extra peak was observed in the diffraction peaks, which indicates that the as-prepared AuNPs were highly purified without any contamination.

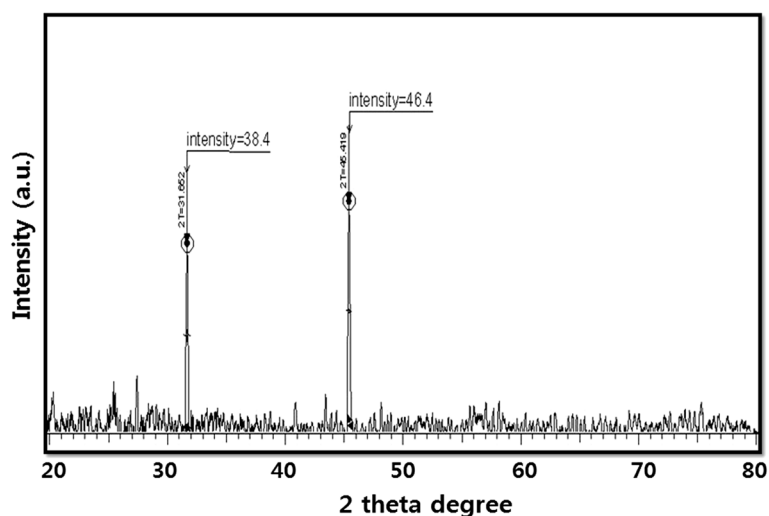


Figure 2 X-ray diffraction spectra of AuNPs.

Gupta and Bector [48] observed four different intense peaks at 2θ angle: 38.22, 44.42, 64.71, and 77.62 with Bragg reflections corresponding to (111), (200), (220), and (311) in biomass-associated AuNPs. Alternatively, only a single prominent peak was observed at 2θ angle: 38.22 with a Bragg reflection corresponding to (111) in extracellular AuNPs. Our present findings are consistent with earlier studies that used biological methods to synthesize AuNPs using plant extracts [49-51], yeast [16], and bacteria [20].

FTIR analysis

The AuNPs synthesized by *Ganoderma* spp. mushroom extract were subjected to FTIR analysis to identify the biomolecules involved in stabilizing the nanoparticles in solution [51]. The AuNPs synthesized by mushroom extract yielded strong bands at 602, 1096, 1201, 1388, and 1636 cm^{-1} (Figure 3). These bands correspond to the

amide I, II, and III bands of polypeptides/proteins, and are consistent with previous reports [51,52]. As suggested by Sastry et al., the polypeptides found in the mushroom extracts served as capping agents in AuNPs, particularly glutathione, which is known to be produced by yeast cells [53].

It is well known that proteins can bind to AuNPs either through free amine groups or cysteine residues in the proteins [54]; therefore, stabilization of the AuNPs by surface-bound proteins is a possibility in the case of AuNPs synthesized by *Ganoderma* spp. Additionally, the bands at 1,636 cm^{-1} can be assigned to the vibrational modes of C=C double bonds of these molecules. The large peak between 1,500 and 1,700 cm^{-1} falls in the region of C=O stretching frequency, and the bands at 3,492 cm^{-1} correspond to carbonyl and hydroxyl functional groups in alcohols and phenol derivatives [11,16,55]. The FTIR results show that the surface capping of AuNPs synthesized

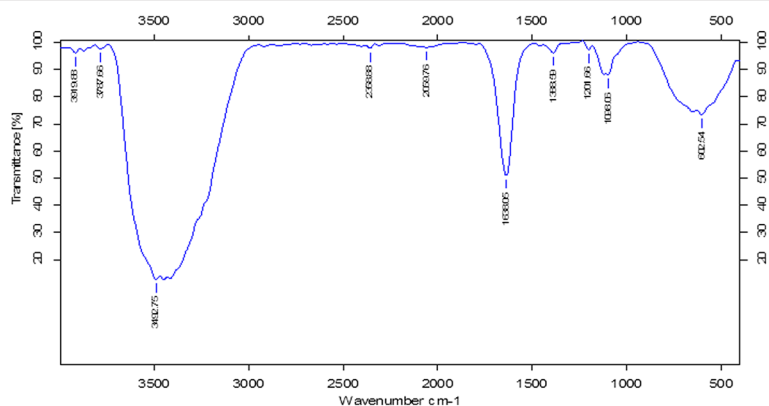


Figure 3 FTIR spectra of AuNPs.

by the mushroom extract is predominantly by proteins. Moreover, our results are consistent with those reported earlier for biosynthesized nanoparticles [11,16,50,51,55].

AuNP synthesis by the *Ganoderma* spp. extract was confirmed using EDS and spectra, as represented in Figure 4. The EDS profile shows a strong gold signal along with weak oxygen and carbon peaks, which may have originated from the biomolecules of the mushroom extract that bound to the AuNP surfaces.

Particle size analysis

Further characterization was carried out to determine the particle size distributions using dynamic light scattering (DLS) technique, which reveals the average hydrodynamic diameter of particles in a liquid suspension. These particle sizes are well within the range reported for photoluminescence of AuNPs [15]. Figure 5 shows the DLS analysis of mushroom extract-mediated synthesis of AuNPs; the average size (20 nm) is within the expected range of particle sizes between 15 to 30 nm and is very similar to the size that was observed in TEM (20 nm). However, for particle sizes larger than 25 nm, the bandwidth increases with the increase in particle size [42], and nanoparticles such as gold and silver have also been shown to exhibit size-dependent optical properties. Husseiny et al. [28] observed the absorption spectra of AuNPs using three different strains of *P. aeruginosa* ATCC 90271, *P. aeruginosa*, and *P. aeruginosa*, and the maximum absorption peaks observed were 543, 540, and 531 nm corresponding to particle sizes of 30 ± 10 , 25 ± 15 , and 15 ± 5 nm, respectively.

In our results, we found that the absorption peak is at 520 nm, which corresponds to a particle size of 20 nm. Moreover, our results are consistent with the absorption spectra and particle size analysis data obtained for chemically prepared AuNPs that have a characteristic band at 524 nm, corresponding to a 20-nm particle size. To

confirm the particular size and shape, synthesized AuNPs were further analyzed using TEM.

TEM analysis

TEM micrographs of the AuNPs revealed distinct, uniform molecules that were spherical in shape and well separated from each other (Figure 6). The average particle size was estimated from counting more than 200 particles from TEM images, and the average size of homogeneous, spherical AuNPs was 20 nm. Interestingly, the AuNPs synthesized by *Ganoderma* spp. are spherical and smaller than those synthesized by other fungi, such as *Colletotrichum* spp. [51] and edible mushrooms [32]; most importantly, the prepared nanoparticles were homogeneous and spherical in shape.

Homogeneous nanoparticles with specific shapes are important for applications in biological and chemical sensing as well as for optical, medical, and electronic devices because the optical properties of AuNPs are dependent on the size and shape [56]. Several studies have reported synthesis of various size AuNPs using different fungi. *Fusarium oxysporum* produced spherical and triangular morphologies of particles with a size range of 20 to 40 nm [15]. Honary et al. [57] reported that *Penicillium aurantiogriseum*, *Penicillium citrinum*, and *Penicillium waksmanii* synthesized AuNPs that were fairly uniform with spherical shapes and had average diameters of 153.3, 172, and 160.1 nm, respectively. Alternatively, the fungi *Aspergillus fumigates* [30] and *Neurospora crassa* [36] produced average AuNP sizes of 25 and 32 nm, respectively.

Effect of AuNPs on cell viability

The use of nontoxic and biocompatible nanoparticles with capping materials is an important aspect of biomedical applications. Consequently, the cytotoxic effects and future health problems caused by nanoparticles must be considered in the engineering of such materials. It is essential to

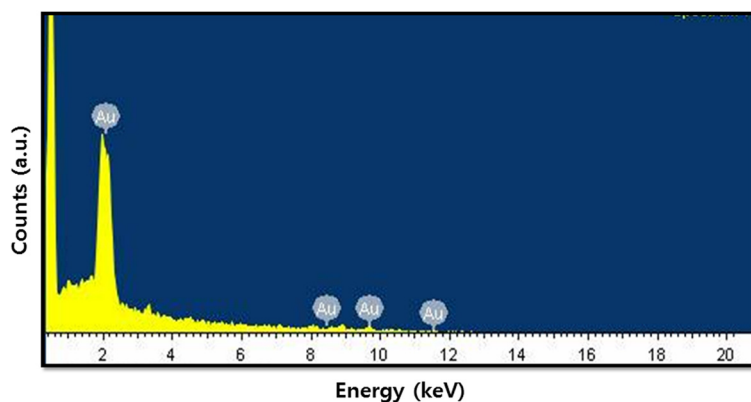


Figure 4 EDX spectra of AuNPs.

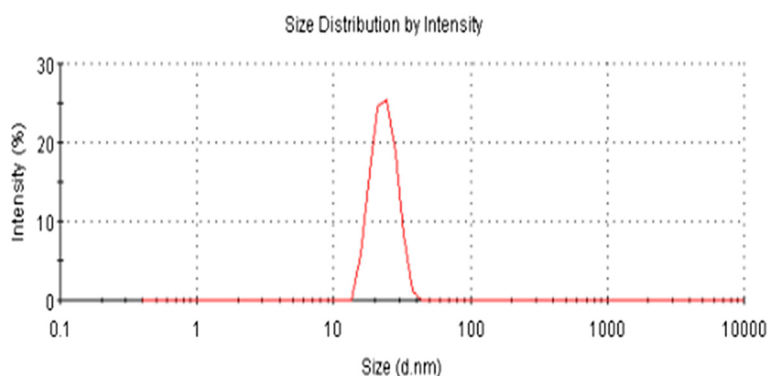


Figure 5 Size distribution analysis of AuNPs by DLS. The particle-size distribution revealed that the average particle size was 20 nm.

validate whether as-prepared AuNPs are toxic or biocompatible, because biomedical applications of any nanomaterial involves intentional exposure to nanoparticles. Therefore, understanding the properties of nanoparticles and their effects on the human body are crucial before they are clinically applied [58].

The biocompatibility of both AuNPs was assessed by a proliferation assay, using mitochondrial functional activity as an indicator of cell viability. The cells were treated with different concentrations of both bio- and chem-AuNPs for 24 h, using the cell viability assay. We found that cells treated with bio-AuNPs showed no toxic effect, even at higher concentrations (Figure 7A), which suggests that as-prepared AuNPs possess biocompatible properties, whereas the chem-AuNPs showed a slight decrease of cell viability at higher concentrations. However, the results are not statistically different from those of the controls. It was also confirmed by incubating AuNPs with medium only and checking the absorption at a wavelength used for MTT assay that the presence of all tested AuNPs did not interfere with the assay.

Shukla *et al.* [59] suggested that AuNPs are not cytotoxic, reduce the production of reactive oxygen and nitrite species, and do not stimulate secretion of proinflammatory cytokines, such as TNF-alpha and IL1-beta, making them suitable candidates for nanomedicine. Using a human leukaemia cell line, gold nanospheres of different sizes (4, 12, and 18 nm in diameter) and capping agents were found to be nontoxic based on the MTT assay [60]. Similarly, Arnida *et al.* [61] found that plain spherical AuNPs and PEGylated spheres and rods did not interfere with the proliferation of PC-3 cells when cells were exposed to as high as 1.5 nM of AuNPs for a period of over two population doubling times (88 h). Plain spherical particles that were 50 and 90 nm in diameter slightly stimulated the proliferation of PC-3 cells. Parab *et al.* [58] investigated the biocompatibility effect of sodium hexametaphosphate (HMP)-stabilized AuNPs (Au-HMPs) in tumor and fibroblast cells. Synthesized Au-HMP nanoparticles and their surface-modified counterparts revealed non-cytotoxic properties in tested cells and showed biocompatibility.

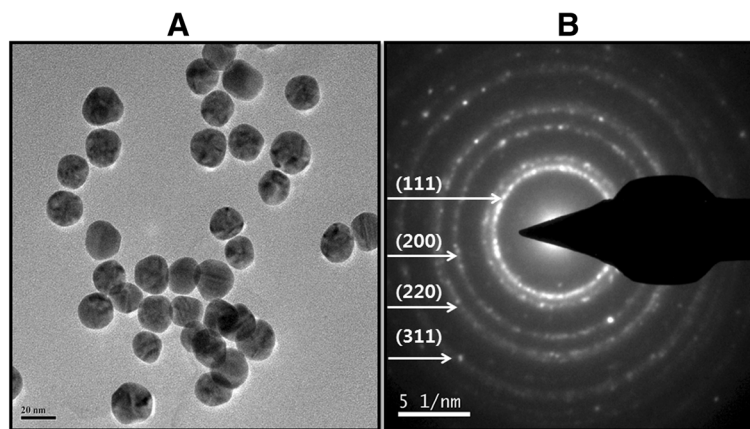


Figure 6 Size and shape analysis of AuNPs by TEM. Several fields were photographed and used to determine the size and morphology of AuNPs (A). Selected area of electron diffraction pattern (B).

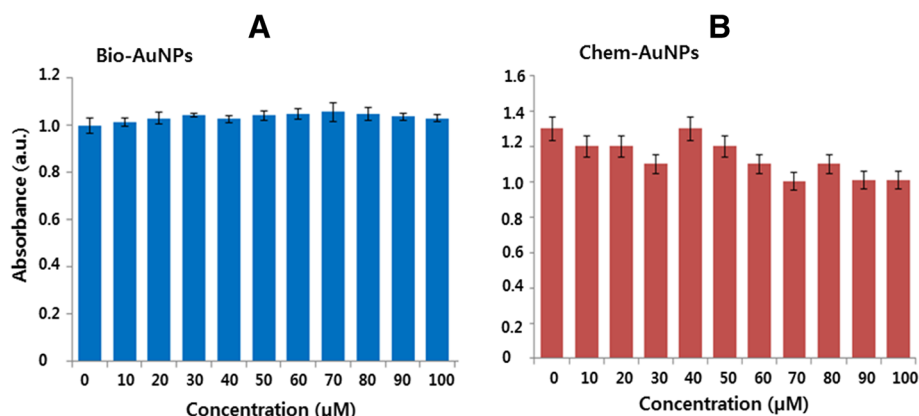


Figure 7 The effect of AuNPs on cell viability of MDA-MB-231 human breast cancer cells. MDA-MB-231 human breast cancer cells were treated with bio-AuNPs (A) or chem-AuNPs (B) at various concentrations from 0 to 100 μM/mL for 24 h, and cell viability was determined by the MTT method. The results are expressed as the mean ± SD of three separate experiments, each of which contained three replicates. Treated groups were not statistically different from the control group based on the Student's *t* test.

Mukherjee et al. [38] designed and developed an AuNP-based drug delivery system (DDS) (Au-DOX) containing doxorubicin (DOX). Administration of this DDS to breast cancer cells (MCF-7 and MDA-MB-231) showed significant inhibition of breast cancer cell proliferation compared with pristine doxorubicin. The viability of the bovine retinal pigment epithelial cells was not affected with an AuNP concentration of up to 300 nM, and increasing the concentrations above 300 nM resulted in significant cell death [62]. AuNPs have anti-oxidative and anti-hyperglycemic activities in streptozotocin-induced diabetic mice by balancing or inhibiting ROS generation in hyperglycemic conditions by scavenging free radicals and leading to increased anti-oxidant defense enzymes in mice.

Effect of AuNPs on LDH leakage

To determine whether AuNPs have an effect on LDH leakage, which is a vital marker of cell death, the cells were treated with various concentrations of bio- and chem-AuNPs (0 to 100 μM) for 24 h. It is interesting to observe that, when treated with both AuNP solutions, there was no change in the leakage of LDH level for up to 24 h in comparison with the untreated control (Figure 8). The LDH and cell viability data are consistent and show no reduction of cell proliferation at higher gold concentrations after 24 h.

Pan et al. [63] found that 1.4-nm gold nanospheres triggered necrosis and mitochondrial damage and induced oxidative stress in endothelial and epithelial cells. In contrast, they found no evidence of cellular damage for

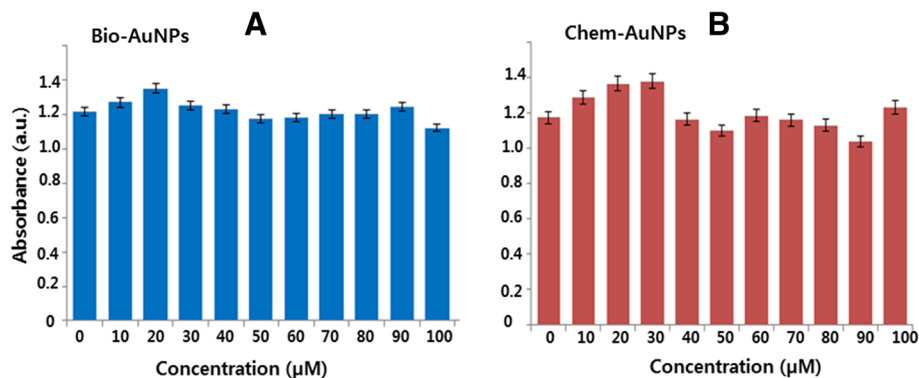


Figure 8 The effect of AuNPs on membrane integrity of cells. MDA-MB-231 human breast cancer cells were treated with bio-AuNPs (A) or chem-AuNPs (B) at various concentrations from 0 to 100 μM/mL for 24 h, and LDH leakage was estimated as described in the 'Methods' section. The results are expressed as the mean ± SD of three separate experiments, each of which contained three replicates. Treated groups were not statistically different from the control group based on the Student's *t* test ($p > 0.05$).

15-nm gold nanospheres bearing the same surface group [63], and these results also suggest that the toxicity of AuNPs depends on size. Interestingly, citrate-capped AuNPs (13 nm in diameter) were found to be toxic to a human carcinoma lung cell line but not to a human liver carcinoma cell line at the same dosage [64].

Uboldi et al. [65] reported that, after 24 to 48 h of exposure, AuNPs induced a mild LDH release in the human A769-like cell line A549, independent of the presence or absence of surface contaminants. Additionally, after 72 h of exposure to AuNPs, there was a dose-dependent release of LDH in the supernatant, and the amount of LDH released was significantly higher compared with shorter exposure times. Zhang et al. [66] reported that chloroplast-mediated synthesis of AuNPs retained up to 85% better viability in both GES-1 and MGC-803 cells, even up to 150 $\mu\text{g}/\text{mL}$ after 36 h of treatment. Freese et al. [67] studied the effect of AuNPs on the amount of LDH released into the supernatant, and they suggest that up to 100 μM of AuNPs did not induce cytotoxicity in human dermal microvascular endothelial cells (HDMECs) and human cardiac microvascular endothelial cells (hCMECs). Altogether, our findings suggest that the biologically derived AuNPs with an average size of 20 nm are biocompatible.

ROS generation

ROS, which are a specific type of oxygen-containing reactive molecule, play important roles in various cellular processes and are known to be essential for basal cell proliferation [68]. Higher concentrations of ROS lead to cell death [69,70]. Several studies suggested that

nanoparticle-mediated cytotoxicity is associated with ROS production.

In this case, we further examined the effect of AuNPs on oxidative stress utilizing the fluorescent dye H_2DCFDA , which does not exhibit enhanced fluorescence in the presence of AuNPs. Results from exposure of MDA-MB-231 to 100 μM of AuNP solution indicate that there are no noticeable increases of ROS generation at the treated concentrations. The results indicate that both bio- and chem-AuNPs are largely ineffective at inducing ROS generation in MDA-MB-231 cells, whereas H_2O_2 - and AgNP-treated groups showed remarkable increase in ROS generation (Figure 9).

Chuang et al. [71] extensively studied the exposure of three different-sized AuNPs in human gastric carcinoma (AGS) and human lung adenocarcinoma epithelial (A549) cells. Their results suggest that significant increases of ROS generation occur with certain concentrations of AuNPs in AGS cells. Conversely, no obvious increases were observed for A549 cells in any of the three sizes of AuNPs. The authors eventually concluded that ROS signaling may play a role in AuNP-induced apoptotic cell death in AGS cells.

Furthermore, western blot analyses revealed that the expression of proteins involved in the anti-oxidative defense system was not significantly modulated any of the three sizes of AuNPs in both lines, except for a modest increase in TrxR-1 and SOD-1 in AGS cells [71]. Altogether, our results suggest that biologically synthesized AuNPs have significant biocompatibility and could possibly be used for ultrasensitive detection, gene transfer, biomolecular imaging, drug delivery, and cancer therapy.

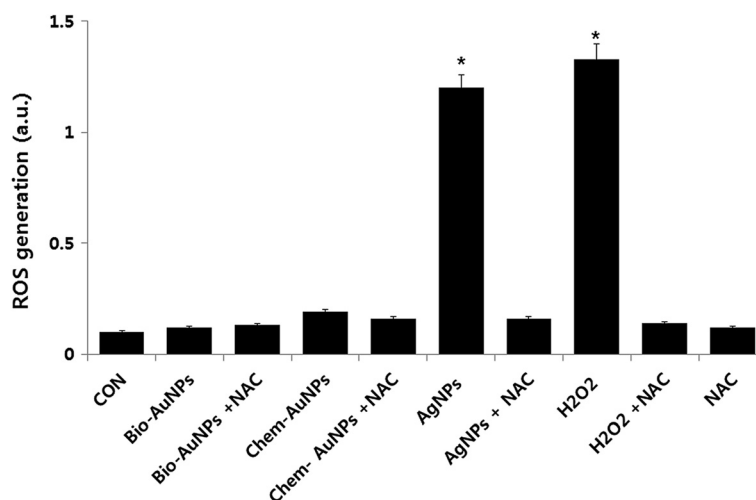


Figure 9 The effect of AuNPs in ROS generation. Relative fluorescence of DCF was measured using a spectrofluorometer with excitation at 485 nm and emission at 530 nm. The results are expressed as the mean \pm SD of three separate experiments, each of which contained three replicates. Treated groups with bio- and chem-AuNPs were not statistically different from the control group based on the Student's *t* test. ($p > 0.05$). H_2O_2 - and AgNP-treated groups were statistically different from the control group based on the Student's *t* test ($*p < 0.05$).

Conclusion

Synthesis of nanoparticles using biological systems is an important area of nanobiotechnology. Here we show a simple, rapid, clean, efficient, cost-effective, and green method for the synthesis of biocompatible AuNPs using *Ganoderma* spp. extract as a reducing and stabilizing agent. The as-prepared AuNPs were characterized via UV-vis, XRD, FTIR, EDX, DLS, and TEM. The biologically derived AuNPs were spherical, discrete, and the average size was 20 nm. The biocompatibility effect of AuNPs was investigated using cell viability, LDH, and ROS assays. The results indicate that biologically derived AuNPs are biocompatible. Finally, this eco-friendly method provides an alternative route for large-scale production of biocompatible AuNPs that can be used in catalysis, sensors, electronics, and biomedical applications, especially for cancer therapy.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

SG came up with the idea and participated in the design, preparation of AuNPs, and writing of the manuscript. JWH performed characterization of nanoparticles. JHP participated in culturing, cell viability, LDH, and ROS assay. SG and JHK participated in coordination of this study. All authors read and approved the final manuscript.

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References

1. Daniel MC, Astruc D: Gold nanoparticles: assembly, supramolecular chemistry, quantum-size-related properties, and applications toward biology, catalysis, and nanotechnology. *Chem Rev* 2004, **104**(1):293–346.
2. Boisselier E, Astruc D: Gold nanoparticles in nanomedicine: preparations, imaging, diagnostics, therapies and toxicity. *Chem Soc Rev* 2009, **38**(5):1759–1782.
3. Saha K, Agasti SS, Kim C, Li XN, Rotello VM: Gold nanoparticles in Chemical and Biological Sensing. *Chem Rev* 2012, **112**(5):2739–2779.
4. Corti CW, Holiday RJ: Commercial aspects of gold applications: from materials science to chemical science. *Gold Bull* 2004, **37**(1–2):20–26.
5. Das SK, Das AR, Guha AK: Microbial synthesis of multishaped gold nanostructures. *Small* 2010, **6**(9):1012–1021.
6. Wong SS, Joselevich E, Woolley AT, Cheung CL, Lieber CM: Covalently functionalized nanotubes as nanometre-sized probes in chemistry and biology. *Nature* 1998, **394**(2):52–55.
7. Tang Z, Kotov NA, Giersig M: Spontaneous organization of single CdTe nanoparticles into luminescent nanowires. *Science* 2002, **297**(12):237–240.
8. Grubbs RB: Nanoparticle assembly: solvent-tuned structures. *Nat Mater* 2007, **6**(8):553–555.
9. Li C, Price JE, Milas L, Hunter NR, Ke S, Yu DF, Charnsangavej C, Wallace S: Antitumor activity of poly(L-glutamic acid)-paclitaxel on syngeneic and xenografted tumors. *Clin Cancer Res* 1999, **5**:891–897.
10. Kim JS, Kuk E, Yu KN, Kim JH, Park SJ, Lee HJ, Kim SH, Park YK, Park YH, Hwang CY, Kim YK, Lee YS, Jeong DH, Cho MH: Antimicrobial effects of silver nanoparticles. *Nanomedicine* 2007, **3**(1):95–101.
11. Suresh AK, Pelletier DA, Wang W, Broich ML, Moon JW, Gu B, Allison DP, Joy DC, Phelps TJ, Doktycz MJ: Biofabrication of discrete spherical gold nanoparticles using the metal-reducing bacterium *Shewanella oneidensis*. *Acta Biomater* 2011, **7**(5):2148–2152.
12. Punties VF, Krishnan KM, Alivisatos AP: Colloidal nanocrystal shape and size control: the case of cobalt. *Science* 2001, **291**(16):2115–2117.
13. Murphy CJ: Nanocubes and nanoboxes. *Science* 2002, **298**(5601):2139–2141.
14. Mukherjee P, Ahmad A, Mandal D, Senapati S, Sainkar SR, Khan MI, Ramani R, Parischa R, Ajayakumar PV, Alam M, Sastry M, Kumar R: Bioreduction of AuCl₄⁻ ions by the fungus, *Verticillium* sp. and surface trapping of the gold nanoparticles formed. *Angew Chem Int Ed Engl* 2001, **40**(19):3585–3588.
15. Mukherjee P, Senapati S, Mandal D, Ahmad A, Khan MI, Kumar R, Sastry M: Extracellular synthesis of gold nanoparticles by the fungus *Fusarium oxysporum*. *ChemBioChem* 2002, **3**(5):461–463.
16. Ahmad A, Mukherjee P, Senapati S, Mandal D, Khan MI, Kumar R, Sastry M: Extracellular biosynthesis of silver nanoparticles using the fungus *Fusarium oxysporum*. *Colloids Surf B: Biointerf* 2003, **28**(4):313–318.
17. Kumar V, Yadav SK: Plant-mediated synthesis of silver and gold nanoparticles and their applications. *J Chem Technol Biot* 2009, **84**(2):151–157.
18. Bhambure R, Bule M, Shaligram N, Kamat M, Singhal R: Extracellular biosynthesis of gold nanoparticles using *Aspergillus niger* – its characterization and stability. *Chem Eng Technol* 2009, **32**(7):1036–1041.
19. Das SK, Das AR, Guha AK: Gold nanoparticles: microbial synthesis and application in water hygiene management. *Langmuir* 2009, **25**(14):8192–8199.
20. Kalishwaralal K, Deepak V, Pandian Ram Kumar S, Gurunathan S: Biological synthesis of gold nanocubes from *Bacillus licheniformis*. *Bioresour Technol* 2009, **100**(21):5356–5358.
21. Kalishwaralal K, Deepak V, Pandian SRK, Kottaisamy M, BarathManiKanth S, Kartikeyan B, Gurunathan S: Biosynthesis of silver and gold nanoparticles using *Brevibacterium casei*. *Colloids Surf B: Biointerf* 2010, **77**(2):257–262.
22. Klaus T, Joerger R, Olsson E, Granqvist CG: Silver-based crystalline nanoparticles, microbially fabricated. *Proc Natl Acad Sci USA* 1999, **96**(24):13611–13614.
23. Jin Y, Li H, Bai J: Homogeneous selecting of a quadruplex-binding ligand-based gold nanoparticle fluorescence resonance energy transfer assay. *Anal Chem* 2009, **81**(14):5709–5715.
24. Narayanan KB, Sakthivel N: Biological synthesis of metal nanoparticles by microbes. *Adv Colloid Interface Sci* 2010, **156**(1–2):1–13.
25. Beveridge TJ, Murray RG: Sites of metal deposition in the cell wall of *Bacillus subtilis*. *J Bacteriol* 1980, **141**(2):876–887.
26. Gurunathan S, Kalishwaralal K, Vaidyanathan R, Venkataraman D, Pandian SR, Muniyandi J, Hariharan N, Eom SH: Biosynthesis, purification and characterization of silver nanoparticles using *Escherichia coli*. *Colloids Surf B: Biointerf* 2009, **74**(1):328–335.
27. Nair B, Pradeep T: Coalescence of nanoclusters and formation of submicron crystallites assisted by *Lactobacillus* strains. *Cryst Growth Des* 2002, **2**(4):293–298.
28. Husseiny MI, El-Aziz MA, Badr Y, Mahmoud MA: Biosynthesis of gold nanoparticles using *Pseudomonas aeruginosa*. *Spectrochim Acta A* 2007, **67**(3–4):1003–1006.
29. He S, Guo Z, Zhang Y, Zhang S, Wang J, Gu N: Biosynthesis of gold nanoparticles using the bacteria *Rhodospseudomonas capsulata*. *Mater Lett* 2007, **61**(18):3984–3987.
30. Bhainsa KC, D'Souza SF: Extracellular biosynthesis of silver nanoparticles using the fungus *Aspergillus fumigatus*. *Colloids Surf B: Biointerf* 2006, **47**(2):160–164.
31. Kathiresan K, Manivannan S, Nabeel MA, Dhivya B: Studies on silver nanoparticles synthesized by a marine fungus, *Penicillium fellutanum* isolated from coastal mangrove sediment. *Colloids Surf B: Biointerf* 2009, **71**(1):133–137.
32. Philip D: Biosynthesis of Au, Ag and Au–Ag nanoparticles using edible mushroom extract. *Spectrochim Acta A Mol Biomol Spectrosc* 2009, **73**(2):374–381.
33. Bhat R, Deshpande R, Ganachari SV, Huh DS, Venkataraman A: Photo-irradiated biosynthesis of silver nanoparticles using edible mushroom *Pleurotus florida* and their antibacterial activity studies. *Bioinorg Chem Appl* 2011, **2011**(2011):7.
34. Ahmad T, Wani IA, Manzoor N, Ahmed J, Asiri AM: Biosynthesis, structural characterization and antimicrobial activity of gold and silver nanoparticles. *Colloids Surf B: Biointerf* 2013, **107**:227–234.

35. Karwa A, Gaikwad S, Rai MK: **Mycosynthesis of silver nanoparticles using Lingzhi or Reishi medicinal mushroom, *Ganoderma lucidum* (W. Curt.:Fr.) P. Karst. and their role as antimicrobials and antibiotic activity enhancers.** *Int J Medicinal Mushrooms* 2011, **13**(5):483–491.
36. Castro-Longoria E, Vilchis-Nestor AR, Avalos-Borja M: **Biosynthesis of silver, gold and bimetallic nanoparticles using the filamentous fungus *Neurospora crassa*.** *Colloids Surf B: Biointerf* 2011, **83**(1):42–48.
37. Jain PK, El-Sayed IH, El-Sayed MA: **Au nanoparticles target cancer.** *Nanotoday* 2007, **2**(1):18–29.
38. Mukherjee S, Sushma V, Patra S, Barui AK, Bhadra MP, Sreedhar B, Ranjan Patra C: **Green chemistry approach for the synthesis and stabilization of biocompatible gold nanoparticles and their potential applications in cancer therapy.** *Nanotechnology* 2012, **23**(45):455103.
39. Seow SLS, Naidu M, David P, Wong KH, Sabaratnam V: **Potential of neurotogenic activity of medicinal mushrooms in rat pheochromocytoma cells.** *BMC Complement Altern Med* 2013, **13**:157.
40. Lin ES, Chen YH: **Factors affecting mycelial biomass and exopolysaccharide production in submerged cultivation of *Antrodia cinnamomea* using complex media.** *Bioresour Technol* 2007, **98**(13):2511–2517.
41. Wong KH, Sabaratnam V, Abdullah N, Naidu M, Keynes R: **Activity of aqueous extracts of lion's mane mushroom *Hericium erinaceus* (Bull.: Fr.) Pers. (Aphyllphoromycetidae) on the neural cell line NG108-15.** *Int J Medicinal Mushrooms* 2007, **9**(1):57–65.
42. Phillips RL, Miranda OR, You CC, Rotello VM, Bunz UHF: **Rapid and efficient identification of bacteria using gold-nanoparticle-poly(paraphenyleneethynylene) constructs.** *Angew Chem Int Ed Engl* 2008, **47**(14):2590–2594.
43. Anil Kumar S, Abyaneh MK, Gosavi SW, Kulkarni SK, Pasricha R, Ahmad A, Khan MI: **Nitrate reductase-mediated synthesis of silver nanoparticles from AgNO₃.** *Biotechnol Lett* 2007, **29**(3):439–445.
44. Marsili E, Baron DB, Shikhare ID, Coursolle D, Gralnick JA, Bond DR: ***Shewanella* secretes flavins that mediate extracellular electron transfer.** *Proc Natl Acad Sci USA* 2008, **105**(10):3968–3973.
45. Fredrickson JK, Romine MF, Beliaev AS, Auchtung JM, Driscoll ME, Gardner TS, Nealon KH, Osterman AL, Pinchuk G, Reed JL, Rodionov DA, Rodrigues JL, Saffarini DA, Serres MH, Spormann AM, Zhulin IB, Tiedje JM: **Towards environmental systems biology of *Shewanella*.** *Nat Rev Microbiol* 2008, **6**(8):592–603.
46. Kumar SA, Peter YA, Nadeau JL: **Facile biosynthesis, separation and conjugation of AuNPs to doxorubicin.** *Nanotechnology* 2008, **19**(49):495101.
47. Das SK, Dickinson C, Lafir F, Brougham DF, Marsili E: **Synthesis, characterization and catalytic activity of AuNPs biosynthesized with *Rhizopus oryzae* protein extract.** *Green Chem* 2012, **14**(5):1322–1334.
48. Gupta S, Bector S: **Biosynthesis of extracellular and intracellular AuNPs by *Aspergillus fumigatus* and *A. flavus*.** *Antonie Van Leeuwenhoek* 2013, **103**(5):1113–1123.
49. Gardea-Torresdey JL, Parsons JG, Gomez E, Peralta-Videa J, Troiani HE, Santiago P, Jose Yacaman M: **Formation and growth of Au nanoparticles inside live Alfalfa plants.** *Nano Lett* 2002, **2**(4):397–401.
50. Shankar SS, Rai A, Ankamwar B, Singh A, Ahmad A, Sastry M: **Biological synthesis of triangular gold nanoprism.** *Nat Mater* 2004, **3**(7):482–488.
51. Shankar SS, Ahmad A, Pasricha R, Sastry M: **Bioreduction of chloroaurate ions by geranium leaves and its endophytic fungus yields gold nanoparticles of different shapes.** *J Mater Chem* 2003, **13**(7):1822–1826.
52. Caruso F, Furlong DN, Ariga K, Ichinose I, Kunitake T: **Characterization of polyelectrolyte-protein multilayer films by atomic force microscopy, scanning electron microscopy, and Fourier transform infrared reflection-absorption spectroscopy.** *Langmuir* 1998, **14**(16):4559–4565.
53. Mehra RK, Winge DR: **Metal ion resistance in fungi: molecular mechanisms and their regulated expression.** *J Cell Biochem* 1991, **45**(1):30–40.
54. Gole A, Dash C, Ramachandran V, Mandale AB, Sainkar SR, Rao M, Sastry M: **Pepsin-gold colloid conjugates: preparation, characterization, and enzymatic activity.** *Langmuir* 2001, **17**(5):1674–1679.
55. Suresh AK, Pelletier DA, Wang W, Moon JW, Gu B, Mortensen NP, Allison DP, Phelps TJ, Doktycz MJ: **Silver nanocrystallites: biofabrication using *Shewanella oneidensis*, and an evaluation of their comparative toxicity on gram-negative and gram-positive bacteria.** *Environ Sci Technol* 2010, **44**(13):5210–5215.
56. Rao CNR, Cheetham AK: **Science and technology of nanomaterials: current status and future prospects.** *J Mater Chem* 2001, **11**(12):2887–2894.
57. Honary S, Gharaei-Fathabad E, Barabadi H, Naghibi F: **Fungus-mediated synthesis of gold nanoparticles: a novel biological approach to nanoparticle synthesis.** *J Nanosci Nanotechnol* 2013, **13**(2):1427–1430.
58. Parab HJ, Huang JH, Lai TC, Jan YH, Liu RS, Wang JL, Hsiao M, Chen CH, Hwu YK, Tsai DP, Chuang SY, Pang JH: **Biocompatible transferrin-conjugated sodium hexametaphosphate-stabilized AuNPs: synthesis, characterization, cytotoxicity and cellular uptake.** *Nanotechnology* 2011, **22**(39):395706.
59. Shukla R, Bansal V, Chaudhary M, Basu A, Bhonde RR, Sastry M: **Biocompatibility of AuNPs and their endocytotic fate inside the cellular compartment: a microscopic overview.** *Langmuir* 2005, **21**(23):10644–10654.
60. Connor EE, Mwamuka J, Gole A, Murphy CJ, Wyatt MD: **Gold nanoparticles are taken up by human cells but do not cause acute cytotoxicity.** *Small* 2005, **1**(3):325–327.
61. Arnida, Malugin A, Ghandehari H: **Cellular uptake and toxicity of gold nanoparticles in prostate cancer cells: a comparative study of rods and spheres.** *J Appl Toxicol* 2010, **30**(3):212–217.
62. Karthikeyan B, Kalishwaralal K, Sheikpranbabu S, Deepak V, Haribalaganesh R, Gurunathan S: **Gold nanoparticles downregulate VEGF- and IL-1 β -induced cell proliferation through Src kinase in retinal pigment epithelial cells.** *Exp Eye Res* 2010, **91**(5):769–778.
63. Pan Y, Leifert A, Ruau D, Neuss S, Bornemann J, Schmid G, Brandau W, Simon U, Jahnhen-Dechent W: **AuNPs of diameter 1.4 nm trigger necrosis by oxidative stress and mitochondrial damage.** *Small* 2009, **5**(18):2067–2076.
64. Patra HK, Banerjee S, Chaudhuri U, Lahiri P, Dasgupta AK: **Cell selective response to gold nanoparticles.** *Nanomed* 2007, **3**(2):111–119.
65. Uboldi C, Bonacchi D, Lorenzi G, Hermanns MI, Pohl C, Baldi G, Unger RE, Kirkpatrick CJ: **AuNPs induce cytotoxicity in the alveolar type-II cell lines A549 and NCIH441.** *Part Fibre Toxicol* 2009, **6**:18.
66. Zhang YX, Zheng J, Gao G, Kong YF, Zhi X, Wang K, Zhang XQ, Cuida X: **Biosynthesis of gold nanoparticles using chloroplasts.** *Int J Nanomedicine* 2011, **6**:2899–2906.
67. Freese C, Uboldi C, Gibson MI, Unger RE, Weksler BB, Romero IA, Couraud PO, Kirkpatrick CJ: **Uptake and cytotoxicity of citrate-coated gold nanospheres: comparative studies on human endothelial and epithelial cells.** *Part Fibre Toxicol* 2012, **9**:23.
68. Burdon RH: **Superoxide and hydrogen peroxide in relation to mammalian cell proliferation.** *Free Radic Biol Med* 1995, **18**(4):775–794.
69. Fiers W, Beyaert R, Declercq W, Vandenaebelle P: **More than one way to die: apoptosis, necrosis and reactive oxygen damage.** *Oncogene* 1999, **18**(54):7719–7730.
70. Matés JM, Segura JA, Alonso FJ, Márquez J: **Intracellular redox status and oxidative stress: implications for cell proliferation, apoptosis, and carcinogenesis.** *Arch Toxicol* 2008, **82**(5):273–299.
71. Chuang SM, Lee YH, Liang RY, Roam GD, Zeng ZM, Tu HF, Wang SK, Chueh PJ: **Extensive evaluations of the cytotoxic effects of gold nanoparticles.** *Biochim Biophys Acta* 2013, **1830**(10):4960–4973.

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