

# Nanotoxic Effects of Silver Nanoparticles on Normal HEK-293 Cells in Comparison to Cancerous HeLa Cell Line

This article was published in the following Dove Press journal:  
International Journal of Nanomedicine

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**Purpose:** Biomimetic approaches for the synthesis of silver nanoparticles (AgNPs) had created a substantial impression among the research community that focuses on nano-bio interactions. In this study, an eco-friendly method using *Rhizophora apiculata* aqueous leaf extract as a reductant-rich hydrosol was followed to synthesize AgNPs and test its cytotoxicity.

**Methods:** To optimise the parameters for the synthesis of AgNPs, central composite design based on response surface methodology was used. The particles synthesized at a nano-scale were characterized in our previously published report. The present report further characterizes the nanoparticles by X-ray diffraction, SEM and TEM at varying sites and magnifications. The characterized AgNPs were tested for their cytotoxic effects on HEK-293 and HeLa cells.

**Results:** The cytotoxicity on the cell lines was dose-dependent. At a concentration of 2.5  $\mu\text{L/mL}$  of the AgNPs-containing hydrosol, 100% inhibition of HEK-293 cells and 75% inhibition of the HeLa cells were observed. The  $\text{IC}_{50}$  value for AgNPs on HEK-293 was 0.622  $\mu\text{L/mL}$  (12.135 ng), whereas, for HeLa cells, it was 1.98  $\mu\text{L/mL}$  (38.629 ng).

**Conclusion:** The nanoparticles were three-fold toxic towards the HEK-293 cells in comparison to the HeLa cells. Therefore, the therapeutic index is low for *R. apiculata* derived AgNPs on HeLa cells when tested in comparison with the HEK-293 cells. The nanotoxicity profile of the synthesized AgNPs seems more prominent than the nanotherapeutic index. According to our knowledge, this is the first-ever report on the optimization of synthesis of AgNPs using response surface methodology and identifying the therapeutic index of mangrove leaf-derived AgNPs.

**Keywords:** AgNPs, XRD, HEK-293, HeLa, nanotoxicity

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## Introduction

Nanoparticles are usually synthesized using two approaches: top-down and bottom-up. Physical methods break particles from bulk to fine forms by a top-down approach. Biological and chemical methods build particles from the bottom-up. The atoms self-assemble into nuclei which later lead to the formation of particles in the size of the nanometer via covalent or supramolecular contact.<sup>1,2</sup> Green approaches for the synthesis of AgNPs using different sources such as plants are a blooming area of research because of their antioxidant and the resultant cytotoxic effects.<sup>3-5</sup> Since plants are a rich source of secondary metabolites that can act as reductants and capping agents, the advantages of biological synthesis are

numerous.<sup>6</sup> The mechanism of nanoparticle synthesis using plants has been elucidated by several reports. The metal ions interact with the phytochemicals of the plant which eventually reduce the metal ions. The reduced ions nucleate and combine to form larger materials from tiny-sized ones by Ostwald ripening. Later, the particles integrate to form a definite and final shape.<sup>7</sup>

Although the advantages of nanomedicine appear plenty, risk assessment and analysis of the exposure of these particles obtained using rapid synthesis procedures towards normal or cancer cells are critical at specific doses for enhanced applications in nanomedicine.<sup>8,9</sup> The toxicity assessment for health hazards of AgNPs is still in its infancy although their role in nanomedicine is well known.<sup>10–12</sup> Few reports indicate both local and systemic toxicity of AgNPs, with mechanisms that relate to toxicity at genome levels, induction of immune cells, and oxidative stress. Various modes of exposure can lead to size, dose and encapsulation-dependent cellular uptake of AgNPs. Further enduring studies using several models can indicate the specific modes of toxicity.<sup>13–16</sup>

Human embryonic kidney 293 (HEK-293) cells are one among the widely used standards for normal human cells. It is believed to be derived from embryonic kidney cells using adenoviral DNA in the 1970s.<sup>17</sup> Nanoparticles, especially nanosilver, are known to be toxic to HEK-293 and other cell lines which are considered to be normal.<sup>18–20</sup> HeLa is the first immortalised human cervical adenocarcinoma cell line to be cultured in vitro in 1951 and stabilised in 1953 at Johns Hopkins Hospital in Baltimore, Maryland, US. It is a key cell line that revolutionised cancer research by accounting for more than sixteen thousand scientific publications related to oncology.<sup>21</sup>

MTT assay, which is a widely applied or standard method to elucidate the cytotoxic potential of agents at the preclinical level, was used to determine the dose related to cell death.<sup>22</sup> In pharmacology, half-maximal inhibitory concentration (IC<sub>50</sub>) is a determinant of how potent an antagonistic drug is.<sup>23</sup> AgNPs are well known for their cytotoxicity on cancer cells, with a special mention to HeLa cells.<sup>24,25</sup> Therefore, MTT assay was used in this study to determine the IC<sub>50</sub> value which is a measure of the efficacy of the tested material.

With this background, the present research intends to test the nanotoxic effects of AgNPs synthesized using the leaf extract of *R. apiculata*. This intention is the basis for recognizing the therapeutic index of such particles in normal and malignant cell lines for applications at the nanoscale.

According to the knowledge of the authors, this is the first-ever international report to optimize the synthesis of AgNPs using response surface methodology (RSM) and to identify the therapeutic index of mangrove tree leaf-derived AgNPs towards HeLa cells in comparison with HEK-293 cells.

## Materials and Methods

### Synthesis of AgNPs

One gram of *R. apiculata* leaf powder was dissolved in 100 mL of Millipore water and incubated at 60 °C for 5 minutes. Five millilitres of the resultant aqueous extract was mixed with 95 mL of 1 mM aqueous AgNO<sub>3</sub> solution and incubated at varying temperatures from 30 °C to 95 °C. After visual observation of colour change to brown, the hydrosol was centrifuged at 10,000 rpm for 20 minutes at 4 °C to derive a pellet that contains AgNPs. The synthesis is performed similarly to the method followed by Song and Kim et al with slight modifications.<sup>26</sup>

### Characterization of AgNPs

The synthesized nanoparticles were characterized by our previous reports using spectroscopic techniques such as ultraviolet-visible, X-ray photoelectron and fourier-transform infrared spectroscopy. Transmission (TEM) and scanning (SEM) electron microscopes were used to determine the morphology of AgNPs. Techniques such as energy dispersive X-ray analysis and inductively coupled plasma-optical emission spectrometry (ICP-OES) were used for establishing the elemental composition. Zeta potential, particle size and polydispersity index were determined using dynamic light scattering.<sup>27–30</sup> The nanoparticles were further characterized by X-ray diffraction (XRD) analysis using PANalytical X'Pert<sup>3</sup> Powder instrument (Malvern Panalytical Inc., Westborough, MA, USA) in this study. SEM and TEM observations were made at different sites and magnifications in comparison to our published report and presented here.

### Cytotoxicity of AgNPs

The cell lines were purchased from National Centre for Cell Science (NCCS), Pune and used for the study. To validate the use of AgNPs for biomedical applications, toxicity assessment against normal HEK-293 and human cervical cancer cell line HeLa was performed, in the present report, in triplicate. Minimum Essential medium-Eagle supplemented with 10% fetal bovine serum was used for maintaining the cells at 37 °C, 5% CO<sub>2</sub>, 95% air and 100% relative

humidity. One hundred microlitres of the cell suspension was implanted at a concentration of 10,000 cells in every well and incubated for 24 hours. After incubation, the medium seeded with HEK-293 and HeLa cells were treated with different concentrations of the AgNPs-containing hydrosol (0.005 to 2.5  $\mu\text{L/mL}$ ). After 48 hours of treatment, the cells were washed and the rate of cellular proliferation was determined by observation through an inverted microscope. 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was performed and the percentage of cell inhibition was calculated using the following formula:

$$\% \text{ cell inhibition} = 100 - \frac{\text{Abs (Test)}}{\text{Abs (Control)}} \times 100$$

Cell viability was plotted in relation to logarithmic concentrations of AgNPs and  $\text{IC}_{50}$  was determined using GraphPad Prism software (Version 6.01).<sup>31,32</sup>

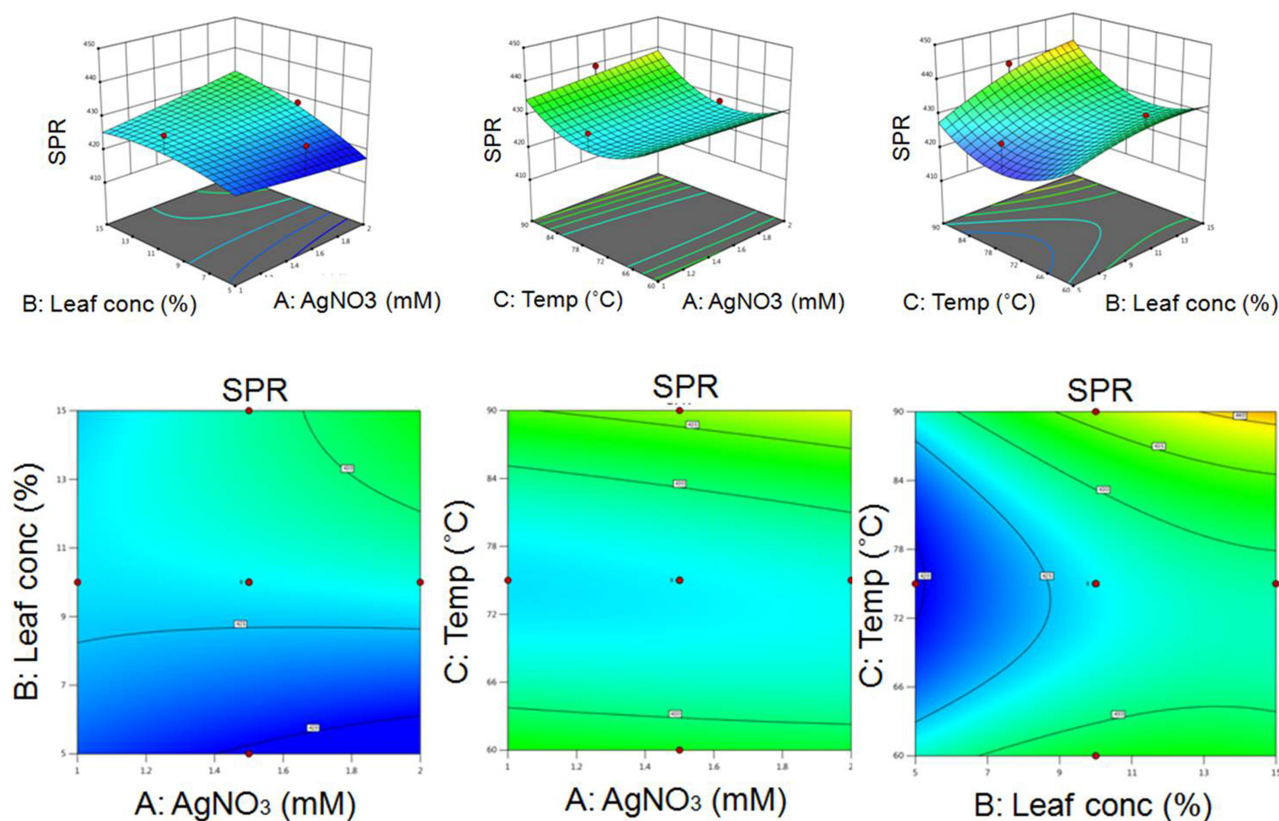
## Data Analysis

The Central composite design (CCD) based on RSM was applied for the study. The optimum responses were calculated

and the Design-Expert software Version 13 (State-Ease Inc., Minneapolis, MN, USA) was used to interpret the results.

## Results and Discussion

Regression analysis is a widely used statistical technique that employs models such as Langmuir.<sup>33–35</sup> According to the statistical analysis, the Model  $F$ -value of 3.18 indicates that the model is significant. The  $F$ -value indicates that there is an extremely low chance (4.28%) of this phenomenon occurring as a resultant of noise. The Adeq Precision indicates a signal-to-noise ratio of 8.067. A ratio larger than 4 is usually desired. Therefore, the ratio observed in this study seems adequate to navigate the design space.<sup>36,37</sup> The  $R^2$  value near to 1 determines an ideal connection between the mean and the data.<sup>38</sup> The  $R^2$  value obtained in this study (0.7413) correlates to this connection better. The mean of surface plasmon resonance (SPR) of AgNPs in this study (429.45 nm) was in the range of 410 to 450 nm indicating spherical nanoparticles of sizes less than 100 nm that are effectively cytotoxic in comparison to microparticles (Figure 1, Tables 1 and 2).<sup>39,40</sup>



**Figure 1** Response surface plots and contour plots for the synthesized AgNPs.

**Table 1** ANOVA, Lack of Fit Test and the Significance of Response Surface Model for the Synthesized AgNPs Using CCD

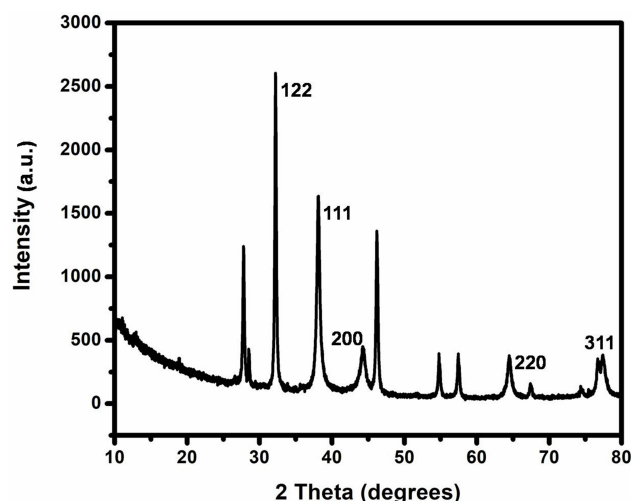
Source	Sum of Squares	df	Mean Square	F-value	p-value	
<b>Model</b>	656.59	9	72.95	3.18	0.0428	Significant
A-AgNO <sub>3</sub> (mM)	4.69	1	4.69	0.2047	0.6606	
B-Leaf conc (%)	215.57	1	215.57	9.41	0.0119	
C-Temp (° C)	45.84	1	45.84	2.00	0.1877	
AB	70.45	1	70.45	3.07	0.1101	
AC	18.42	1	18.42	0.8038	0.3910	
BC	44.94	1	44.94	1.96	0.1917	
A2	0.0799	1	0.0799	0.0035	0.9541	
B2	13.18	1	13.18	0.5752	0.4657	
C2	183.58	1	183.58	8.01	0.0178	
<b>Residual</b>	229.19	10	22.92			
Lack of Fit	229.19	5	45.84			
Pure Error	0.0000	5	0.0000			
<b>Cor Total</b>	885.78	19				

**Table 2** Regression Analysis for the Synthesized AgNPs Using CCD

<b>Std. Dev.</b>	<b>4.79</b>	<b>R<sup>2</sup></b>	<b>0.7413</b>
<b>Mean</b>	429.45	<b>Adjusted R<sup>2</sup></b>	0.5084
<b>C.V. %</b>	1.11	<b>Predicted R<sup>2</sup></b>	-1.4496
		<b>Adeq Precision</b>	8.0669

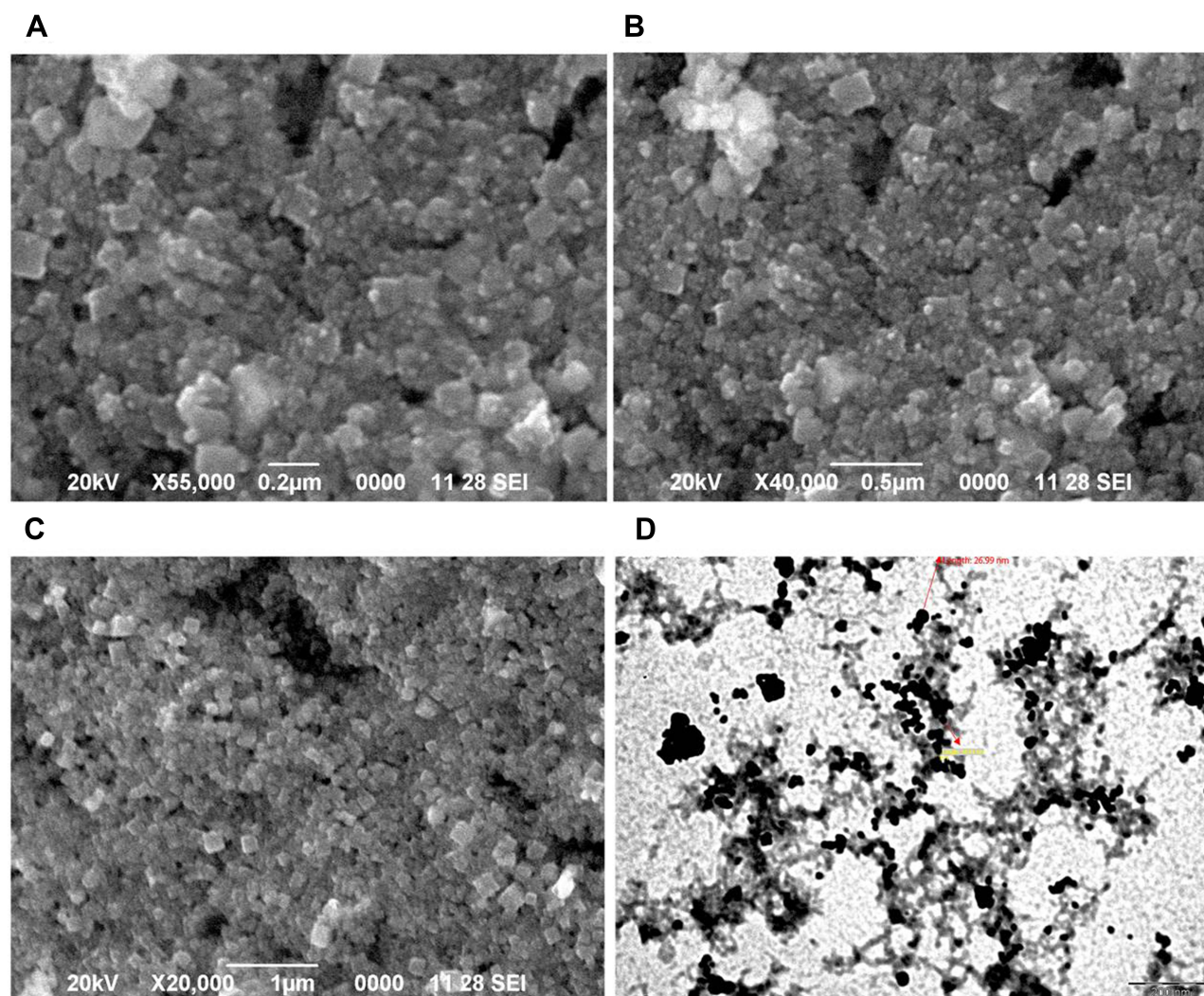
The examination of the structure and crystalline size of the biosynthesized AgNPs was performed using XRD. The pattern indicates diffraction peaks at (2θ) 32.19°, 38.07°, 44.25°, 64.43° and 77.38°. The peaks could be allocated to the (122), (111), (200), (220), and (311) typical planes of face-centered, cubic, and crystalline silver synthesized using green methods (JCPDS file number: 04-0783). The average crystalline size of the AgNPs was calculated using the Debye–Scherrer's equation:  $D = K\lambda/\beta \cos\theta$ .  $K$  is equivalent to 0.94, whereas,  $\lambda$  is equivalent to 1.54178.  $\beta$  is the line broadening in radians and  $\theta$  is the Bragg's angle. According to Bragg's reflection, the estimated average size of the particles was 31.12 nm. The broadening of Bragg's peaks positioned specifies the formation of smaller sized nanoparticles. The unassigned peaks could be correlated to the existence of phytoconstituents<sup>41–44</sup> (Figure 2). SEM and TEM are electron microscopic techniques used to study the morphology of nanomaterials at various magnifications.<sup>45,46</sup> The representative images are presented in Figure 3A–D.

Chemotherapeutic drugs lack the capacity to segregate normal forms from cells that are malignant.<sup>47</sup> Hence, particles are being fabricated at the nano-regime

**Figure 2** XRD pattern of the synthesized nanoparticles.

to specifically target cancer cells with limited toxicity. Such medical systems that can deter the harmful after-effects of conventional therapeutic methods are being approved for clinical practice recently. These nanoparticles can enter the tumor microenvironment efficiently and inhibit the cancer cells with the ability to metastasize, from spreading to other sites.<sup>48–51</sup> Therapeutic index is the measurement or comparison of the ratio of inhibition of normal and cancer cells by any medication. This can lead to the identification of a safety window intended for the treatment of neoplasms.<sup>52</sup> Consequently, assessment of toxicity on various types of cells is a critical point to warrant the safety of such nano-carriers.<sup>53,54</sup> Toxicity associated with kidneys is usually tested to identify a drug as they determine the





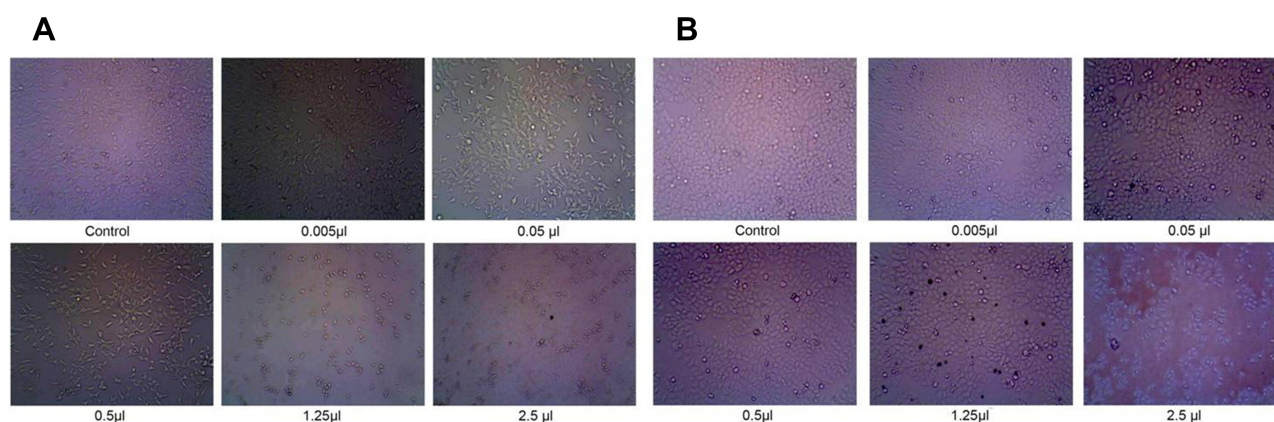
**Figure 3** Electron microscopic images of the synthesized AgNPs (A) SEM image taken at 0.2  $\mu\text{m}$  (B) SEM image taken at 0.5  $\mu\text{m}$  (C) SEM image taken at 1  $\mu\text{m}$  (D) TEM image taken at 200 nm.

homeostasis of the body.<sup>55</sup> Assessment of toxicity to kidneys is critical in clinical practice.<sup>56</sup> Therefore, HEK-293 cells were chosen for this study with regard to the identification of nanotoxicity.

AgNPs are renowned for their biomedical applications.<sup>57</sup> Yet, extensive exposure to AgNPs can cause systemic nanotoxic effects including argyria in humans and intensify the hostile effects towards his environment.<sup>58–60</sup> Supportive of the aforementioned properties, AgNPs are known to be cytotoxic and genotoxic towards normal cells like HEK-293. Although its origin is still unclear, this normal cell line is used in various biological experiments. The purpose of these trials is to identify the toxicity profile and therapeutic effects with an intention of screening a drug.<sup>61–63</sup> Therefore, we analysed the

cytotoxicity of AgNPs on normal HEK-293 and HeLa cells, in order to indicate its therapeutic window. As an initial analysis, microscopic observations indicated the annihilating effect of AgNPs towards HEK-293 and HeLa cells (Figure 4A and B).

Nanotoxicity of AgNPs is dependent on quite a lot of characteristics such as size, surface, shape, agglomeration, dose and route of administration. Dose determination of an anticancer drug is very critical and foremost for formulating therapeutic strategies. This is due to the fact that an increased dose can cause a decline in the antitumor effect and an upsurge in unintended toxicity. Observations from cytotoxicity experiments indicate that 0.4  $\mu\text{g/mL}$  is the minimal  $\text{IC}_{50}$  value for AgNPs among cell lines which are considered to be normal. The highest value analysed was 250  $\mu\text{g/mL}$ . Twenty-five ppm



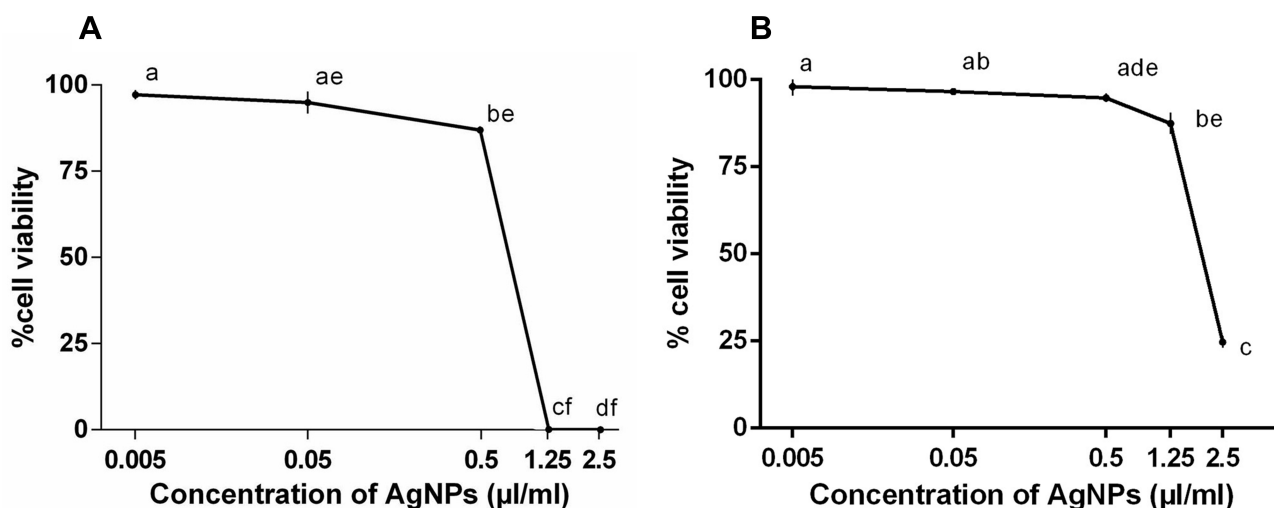
**Figure 4** Microscopic observations of the cytotoxic effects of AgNPs on (A) HEK-293 cells (B) HeLa cells.

was the most toxic dose for liver cells. However, genotoxic effects were observed at 0.01–10 mg/mL in BEAS-2B, a normal lung epithelial cell line. It is interesting to note that the human body can tolerate 0.4 to 27 µg of AgNPs per day when consumed through the oral route. Therefore, the nanotoxicity profile can differ based on the dose and the origin of cell lines subjected to a specific toxicity analysis.<sup>64–67</sup>

The results of the present study reveal 100% cell death of HEK-293 cells at 2.5 µL/mL. At the same concentration of 2.5 µL/mL required for 100% killing of HEK-293 cells, only 75% cell death was observed among HeLa cells. The percentage of viability is depicted in Figure 5A and B. AgNPs used in this study were less effective against the HeLa cells in comparison to the cytotoxicity exerted by AgNPs on other cancer cell lines like HepG2, NIH-3T3,

PC-12, A-549, HCT116 and SiHa cells. In previous such reports, the IC<sub>50</sub> value for AgNPs on cancerous cell lines did range from 3 to 99 ppm.<sup>68–71</sup> Explicitly, according to existing reports, the IC<sub>50</sub> values for AgNPs on HeLa cell lines range from 19 to 51 ppm.<sup>72–74</sup>

In the current study, IC<sub>50</sub> value for AgNPs on HeLa cells was 1.98 µL/mL (1980 ppm). The concentration of AgNPs in the hydrosol was equivalent to 38.629 ng, as per ICP-OES analysis (19.51 µg/mL). The IC<sub>50</sub> value for the hydrosol against HEK-293 cells was 0.622 µL/mL (622 ppm). The concentration of AgNPs in 0.622 µL of the hydrosol was equivalent to 12.135 ng. The IC<sub>50</sub> values determine that the AgNPs were three-fold toxic towards HEK-293 cells in comparison to toxicity exerted on HeLa cells. Therefore, based on the IC<sub>50</sub> value, this study determines that AgNPs



**Figure 5** Cytotoxicity of AgNPs on (A) HEK-293 cells (B) HeLa cells. Results were expressed as Mean ± SEM (n = 3) and statistically analysed using one-way ANOVA along with Tukey's post hoc test of significance; different alphabets denote significant difference (p < 0.05).

synthesized using *R. apiculata* were more toxic towards HEK-293 cells in comparison to HeLa cells.

A general expectancy in screening an anticancer drug is to sensitize and destroy the cancer cells rather than the normal cells.<sup>75</sup> The mechanism of cytotoxicity of nanoparticles is dependent on mechanisms that involve (i) an increase in calcium levels, (ii) genotoxic effects that lead to cell cycle arrest at G<sub>2</sub>/M phase, and (iii) ROS, JNK signaling and mitochondria-dependent apoptosis.<sup>76</sup> To conclude, the therapeutic index is low for the metastatic HeLa cells, in comparison to the normal cells (HEK-293), as analysed through the present report.

## Conclusion

Conferring to the results of the present study and the supportive conclusions of previously published reports, the AgNPs synthesized using *R. apiculata* could be considered to be nanotoxic against HEK-293 cells while comparing the IC<sub>50</sub> values with those of HeLa cells. The cell inhibiting effects were dependent on the dose used. The AgNPs were less effective against HeLa cells in comparison to HEK-293 cells. Therefore, this study identified that the therapeutic index for HeLa cells is poor and the therapeutic window is not extensive. The nanotoxic effects emerge more towards HEK-293 cells. Although HEK-293 and HeLa cells are of various origins, the work was based on the ideology for future analysis of the systemic toxicity of AgNPs. This means that the AgNPs intended for the therapy of cervical cancer cell model HeLa was toxic towards the normal cell line HEK-293 of the renal system. Further comparative nanotoxicity analysis and mechanistic studies on such toxic effects may provide an insight into the use of these AgNPs for applications in cancer nanomedicine.

## Acknowledgments

We thank Nanome Consulting for the technology transfer of nanoparticle synthesis.

## Disclosure

The authors report no conflicts of interest in this work.

## References

- Salem SS, Fouda A. Green synthesis of metallic nanoparticles and their prospective biotechnological applications: an overview. *Biol Trace Elem Res*. 2020.
- Iqbal P, Preece JA, Mendes PM. Nanotechnology: the "Top-Down" and "Bottom-Up" Approaches. In: *Supramolecular Chemistry: From Molecules to Nanomaterials*. 2012.
- Sathishkumar RS, Sundaramanickam A, Srinath R, et al. Green synthesis of silver nanoparticles by bloom forming marine microalgae *Trichodesmium erythraeum* and its applications in antioxidant, drug-resistant bacteria, and cytotoxicity activity. *J Saudi Chem Soc*. 2019;23:1180–1191. doi:10.1016/j.jscs.2019.07.008
- Algebaly AS, Mohammed AE, Abutaha N, Elobeid MM. Biogenic synthesis of silver nanoparticles: antibacterial and cytotoxic potential. *Saudi J Biol Sci*. 2020;27:1340–1351. doi:10.1016/j.sjbs.2019.12.014
- Hemlata MPR, Singh AP, Tejavath KK. Biosynthesis of silver nanoparticles using cucumis prophetarum aqueous leaf extract and their antibacterial and antiproliferative activity against cancer cell lines. *ACS Omega*. 2020;5:5520–5528. doi:10.1021/acsomega.0c00155
- Huang H, Shan K, Liu J, et al. Synthesis, optimization and characterization of silver nanoparticles using the catkin extract of piper longum for bactericidal effect against food-borne pathogens via conventional and mathematical approaches. *Bioorg Chem*. 2020;103:104230. doi:10.1016/j.bioorg.2020.104230
- Singh J, Dutta T, Kim K-H, Rawat M, Samddar P, Kumar P. 'Green' synthesis of metals and their oxide nanoparticles: applications for environmental remediation. *J Nanobiotechnol*. 2018;16:84.
- Mousavi SM, Hashemi SA, Ghasemi Y, et al. Green synthesis of silver nanoparticles toward bio and medical applications: review study. *Artif Cells, Nanomed Biotechnol*. 2018;46:S855–S72. doi:10.1080/21691401.2018.1517769
- Singh AV, Laux P, Luch A, et al. Review of emerging concepts in nanotoxicology: opportunities and challenges for safer nanomaterial design. *Toxicol Mech Methods*. 2019;29:378–387. doi:10.1080/15376516.2019.1566425
- Calderón-Jiménez B, Johnson ME, Montoro Bustos AR, Murphy KE, Winchester MR, Vega Baudrit JR. Silver nanoparticles: technological advances, societal impacts, and metrological challenges. *Front Chem*. 2017;5:6.
- McClements DJ, Xiao H. Is nano safe in foods? Establishing the factors impacting the gastrointestinal fate and toxicity of organic and inorganic food-grade nanoparticles. *Npj Science of Food*. 2017;1:6. doi:10.1038/s41538-017-0005-1
- Lee SH, Jun B-H. Silver nanoparticles: synthesis and application for nanomedicine. *Int J Mol Sci*. 2019;20:865. doi:10.3390/ijms20040865
- Stensberg MC, Wei Q, McLamore ES, Porterfield DM, Wei A, Sepúlveda MS. Toxicological studies on silver nanoparticles: challenges and opportunities in assessment, monitoring and imaging. *Nanomedicine (London, England)*. 2011;6:879–898. doi:10.2217/nnm.11.78
- Ferdous Z, Nemmar A. Health impact of silver nanoparticles: a review of the biodistribution and toxicity following various routes of exposure. *Int J Mol Sci*. 2020;21:2375. doi:10.3390/ijms21072375
- Yang Y, Qin Z, Zeng W, et al. Toxicity assessment of nanoparticles in various systems and organs. *Nanotechnol Rev*. 2017;6:279–289. doi:10.1515/ntrev-2016-0047
- Chen L, Wu M, Jiang S, et al. Skin toxicity assessment of silver nanoparticles in a 3D epidermal model compared to 2D keratinocytes. *Int J Nanomedicine*. 2019;14:9707. doi:10.2147/IJN.S225451
- Kavsan VM, Iershov AV, Balynska OV. Immortalized cells and one oncogene in malignant transformation: old insights on new explanation. *BMC Cell Biol*. 2011;12:23. doi:10.1186/1471-2121-12-23
- Sooklert K, Chattong S, Manotham K, et al. Cytoprotective effect of glutaraldehyde erythropoietin on HEK293 kidney cells after silver nanoparticle exposure. *Int J Nanomedicine*. 2016;11:597. doi:10.2147/IJN.S95654
- Peng H, Zhang X, Wei Y, et al. Cytotoxicity of silver nanoparticles in human embryonic stem cell-derived fibroblasts and an L-929 cell line. *J Nanomater*. 2012;2012:160145. doi:10.1155/2012/160145



20. Sahu SC, Hayes AW. Toxicity of nanomaterials found in human environment: a literature review. *Toxicol Res Appl*. 2017;1:2397-847317726352. doi:10.1177/2397847317726352
21. Mirabelli P, Coppola L, Salvatore M. Cancer cell lines are useful model systems for medical research. *Cancers (Basel)*. 2019;11:1098. doi:10.3390/cancers11081098
22. Bahuguna A, Khan I, Bajpai VK, Kang SC. MTT assay to evaluate the cytotoxic potential of a drug. *Bangladesh J Pharmacol*. 2017;12: Online: Apr. doi:10.3329/bjp.v12i2.30892
23. Aykul S, Martinez-Hackert E. Determination of half-maximal inhibitory concentration using biosensor-based protein interaction analysis. *Anal Biochem*. 2016;508:97–103. doi:10.1016/j.ab.2016.06.025
24. Yuan Y-G, Zhang S, Hwang J-Y, Kong I-K. Silver nanoparticles potentiate cytotoxicity and apoptotic potential of camptothecin in human cervical cancer cells. *Oxid Med Cell Longev*. 2018;2018:6121328. doi:10.1155/2018/6121328
25. Ratan ZA, Haidere MF, Nurunnabi M, et al. Green chemistry synthesis of silver nanoparticles and their potential anticancer effects. *Cancers (Basel)*. 2020;12:855. doi:10.3390/cancers12040855
26. Song JY, Kim BS. Rapid biological synthesis of silver nanoparticles using plant leaf extracts. *Bioprocess Biosyst Eng*. 2009;32:79–84. doi:10.1007/s00449-008-0224-6
27. Antony JJ, Sivalingam P, Siva D, et al. Comparative evaluation of antibacterial activity of silver nanoparticles synthesized using *Rhizophora apiculata* and glucose. *Colloids Surf B Biointerfaces*. 2011;88:134–140. doi:10.1016/j.colsurfb.2011.06.022
28. Jacob JA, Shanmugam A. Silver nanoparticles provoke apoptosis of Dalton's ascites lymphoma in vivo by mitochondria dependent and independent pathways. *Colloids Surf B Biointerfaces*. 2015;136:1011–1016. doi:10.1016/j.colsurfb.2015.11.004
29. Zhang H, Jacob JA, Jiang Z, et al. Hepatoprotective effect of silver nanoparticles synthesized using aqueous leaf extract of *Rhizophora apiculata*. *Int J Nanomedicine*. 2019;14:3517–3524. doi:10.2147/IJN.S198895
30. Wen X, Wang Q, Dai T, et al. Identification of possible reductants in the aqueous leaf extract of mangrove plant *Rhizophora apiculata* for the fabrication and cytotoxicity of silver nanoparticles against human osteosarcoma MG-63 cells. *Mater Sci Eng C*. 2020;116:111252. doi:10.1016/j.msec.2020.111252
31. Mosmann T. Rapid colorimetric assay for cellular growth and survival: application to proliferation and cytotoxicity assays. *J Immunol Methods*. 1983;65:55–63. doi:10.1016/0022-1759(83)90303-4
32. Monks A, Scudiero D, Skehan P, et al. Feasibility of a high-flux anticancer drug screen using a diverse panel of cultured human tumor cell lines. *J Natl Cancer Inst*. 1991;83:757–766. doi:10.1093/jnci/83.11.757
33. Kumari K, Yadav S. Linear regression analysis study. *J Pract Cardiovasc Sci*. 2018;4:33. doi:10.4103/jpcs.jpcs\_8\_18
34. Marković DD, Lekić BM, Rajaković-Ognjanović VN, Onjia AE, Rajaković LV, New A. Approach in regression analysis for modeling adsorption isotherms. *Sci World J*. 2014;2014:930879. doi:10.1155/2014/930879
35. Rashtbari Y, Arfaeina H, Ahmadi S, et al. Potential of using green adsorbent of humic acid removal from aqueous solutions: equilibrium, kinetics, thermodynamic and regeneration studies. *Int J Environ Anal Chem*. 2020;1–18. doi:10.1080/03067319.2020.1796993
36. Ahmadi S, Mohammadi L, Rahdar A, et al. Acid dye removal from aqueous solution by using neodymium(III) oxide nanoadsorbents. *Nanomaterials*. 2020;10:556. doi:10.3390/nano10030556
37. Sandhya M, Rajkumar K, Burgula S. Efficient eco-friendly approach towards bimetallic nanoparticles synthesis and characterization using *Exiguobacterium aestuarii* by statistical optimization. *Green Chem Lett Rev*. 2019;12:420–434. doi:10.1080/17518253.2019.1687762
38. Saunders LJ, Russell RA, Crabb DP. The coefficient of determination: what determines a useful R2 statistic? *Invest Ophthalmol Vis Sci*. 2012;53:6830–6832. doi:10.1167/iov.12-10598
39. Suriyakalaa U, Antony JJ, Suganya S, et al. Hepatocurative activity of biosynthesized silver nanoparticles fabricated using *Andrographis paniculata*. *Colloids Surf B Biointerfaces*. 2013;102:189–194. doi:10.1016/j.colsurfb.2012.06.039
40. Sahu D, Kannan G, Tailang M, Vijayaraghavan R. In vitro cytotoxicity of nanoparticles: a comparison between particle size and cell type. *J Nanosci*. 2016;2016:1–9. doi:10.1155/2016/4023852
41. Jemal K, Sandeep B, Pola S. Synthesis, characterization, and evaluation of the antibacterial activity of *allophylus serratus* leaf and leaf derived callus extracts mediated silver nanoparticles. *J Nanomater*. 2017;2017:1–11. doi:10.1155/2017/4213275
42. Anandalakshmi K, Venugobal J, Ramasamy V. Characterization of silver nanoparticles by green synthesis method using *Pedicular murex* leaf extract and their antibacterial activity. *Appl Nanosci*. 2016;6:399–408. doi:10.1007/s13204-015-0449-z
43. Sree GS, Botsa SM, Reddy BJM, Ranjitha KVB. Enhanced UV–Visible triggered photocatalytic degradation of Brilliant green by reduced graphene oxide based NiO and CuO ternary nanocomposite and their antimicrobial activity. *Arab J Chem*. 2020;13:5137–5150. doi:10.1016/j.arabjc.2020.02.012
44. Botsa SM, Naidu GP, Ravichandra M, Rani SJ, Anjaneyulu RB, Ramana CV. Flower like SnO<sub>2</sub>-Fe<sub>2</sub>O<sub>3</sub>-rGO ternary composite as highly efficient visible light induced photocatalyst for the degradation of organic pollutants from contaminated water. *J Mater Res Technol*. 2020;9:12461–12472. doi:10.1016/j.jmrt.2020.08.087
45. Zhang XF, Liu ZG, Shen W, Gurunathan S. Silver nanoparticles: synthesis, characterization, properties, applications, and therapeutic approaches. *Int J Mol Sci*. 2016;17:1534. doi:10.3390/ijms17091534
46. Su D. Advanced electron microscopy characterization of nanomaterials for catalysis. *Green Energy Environ*. 2017;2:70–83. doi:10.1016/j.jee.2017.02.001
47. Yuan M, Huang -L-L, Chen J-H, Wu J, Xu Q. The emerging treatment landscape of targeted therapy in non-small-cell lung cancer. *Signal Transduct Target Ther*. 2019;4:61. doi:10.1038/s41392-019-0099-9
48. Parasuraman S. Toxicological screening. *J Pharmacol Pharmacother*. 2011;2:74–79. doi:10.4103/0976-500X.81895
49. Sutradhar KB, Amin ML. Nanotechnology in cancer drug delivery and selective targeting. *ISRN Nanotechnol*. 2014;2014:939378. doi:10.1155/2014/939378
50. Miao L, Huang L. Exploring the tumor microenvironment with nanoparticles. *Cancer Treat Res*. 2015;166:193–226.
51. Fernandes C, Soares D, Yegeri MC. Tumor microenvironment targeted nanotherapy. *Front Pharmacol*. 2018;9.
52. Deepa PR, Vandhana S, Jayanthi U, Krishnakumar S. Therapeutic and toxicologic evaluation of anti-lipogenic agents in cancer cells compared with non-neoplastic cells. *Basic Clin Pharmacol Toxicol*. 2012;110:494–503. doi:10.1111/j.1742-7843.2011.00844.x
53. Li W, Zhou J, Xu Y. Study of the in vitro cytotoxicity testing of medical devices. *Biomed Rep*. 2015;3:617–620. doi:10.3892/br.2015.481
54. Patra JK, Das G, Fraceto LF, et al. Nano based drug delivery systems: recent developments and future prospects. *J Nanobiotechnol*. 2018;16:71.
55. Kim SY, Moon A. Drug-induced nephrotoxicity and its biomarkers. *Biomol Ther (Seoul)*. 2012;20:268–272. doi:10.4062/biomolther.2012.20.3.268
56. Basile DP, Anderson MD, Sutton TA. Pathophysiology of acute kidney injury. *Compr Physiol*. 2012;2:1303–1353. doi:10.1002/cphy.c110041
57. A-C B, Gherasim O, Grumezescu AM, Mogoantă L, Fica A, Andronescu E. Biomedical applications of silver nanoparticles: an up-to-date overview. *Nanomaterials (Basel)*. 2018;8:681. doi:10.3390/nano8090681
58. Antony JJ, Sivalingam P, Chen B. Toxicological effects of silver nanoparticles. *Environ Toxicol Pharmacol*. 2015;40:729–732. doi:10.1016/j.etap.2015.09.003



59. Paknejadi M, Bayat M, Salimi M, Razavilar V. Concentration- and time-dependent cytotoxicity of silver nanoparticles on normal human skin fibroblast cell line. *Iran Red Crescent Med J.* **2018**;20.
60. Murphy M, Ting K, Zhang X, Soo C, Zheng Z. Current development of silver nanoparticle preparation, investigation, and application in the field of medicine. *J Nanomater.* **2015**;2015:696918. doi:10.1155/2015/696918
61. Jiang X, Lu C, Tang M, et al. Nanotoxicity of silver nanoparticles on HEK293T cells: a combined study using biomechanical and biological techniques. *ACS Omega.* **2018**;3:6770–6778. doi:10.1021/acsomega.8b00608
62. Bethu MS, Netala VR, Domdi L, Tartte V, Janapala VR. Potential anticancer activity of biogenic silver nanoparticles using leaf extract of *Rhynchosia suaveolens*: an insight into the mechanism. *Artif Cells, Nanomed Biotechnol.* **2018**;46:104–114. doi:10.1080/21691401.2017.1414824
63. Stepanenko AA, Dmitrenko VV. HEK293 in cell biology and cancer research: phenotype, karyotype, tumorigenicity, and stress-induced genome-phenotype evolution. *Gene.* **2015**;569:182–190. doi:10.1016/j.gene.2015.05.065
64. Yildirim L, Thanh NTK, Loizidou M, Seifalian AM. Toxicology and clinical potential of nanoparticles. *Nano Today.* **2011**;6:585–607. doi:10.1016/j.nantod.2011.10.001
65. Budama-Kilinc Y, Cakir-Koc R, Zorlu T, et al. Assessment of nanotoxicity and safety profiles of silver nanoparticles. In: Khan M, editor. *Silver Nanoparticles-Fabrication, Characterization and Applications*. London: Intechopen; **2018**:185.
66. Akter M, Sikder MT, Rahman MM, et al. A systematic review on silver nanoparticles-induced cytotoxicity: physicochemical properties and perspectives. *J Adv Res.* **2018**;9:1–16. doi:10.1016/j.jare.2017.10.008
67. Gurney H. How to calculate the dose of chemotherapy. *Br J Cancer.* **2002**;86:1297–1302. doi:10.1038/sj.bjc.6600139
68. Sambale F, Wagner S, Stahl F, Khaydarov R, Scheper T, Bahnmann D. Investigations of the toxic effect of silver nanoparticles on mammalian cell lines. *J Nanomater.* **2015**;2015:1–9. doi:10.1155/2015/136765
69. Mishra A, Mehdi SJ, Irshad M, et al. Effect of biologically synthesized silver nanoparticles on human cancer cells. *Sci Adv Mat.* **2012**;4:1200–1206. doi:10.1166/sam.2012.1414
70. Sufyani A, Moslah N, Hussien NA, Hawsawi YM. Characterization and anticancer potential of silver nanoparticles biosynthesized from *olea chrysophylla* and *lavandula dentata* leaf extracts on HCT116 colon cancer cells. *J Nanomater.* **2019**;2019:1–9. doi:10.1155/2019/7361695
71. Aziz N, Faraz M, Sherwani MA, Fatma T, Prasad R. Illuminating the anticancerous efficacy of a new fungal chassis for silver nanoparticle synthesis. *Front Chem.* **2019**;7:65. doi:10.3389/fchem.2019.00065
72. Chanthini AB, Balasubramani G, Ramkumar R, et al. Structural characterization, antioxidant and in vitro cytotoxic properties of seagrass, *Cymodocea serrulata* (R. Br.) Asch. & Magnus mediated silver nanoparticles. *J Photochem Photobiol B.* **2015**;153:145–152. doi:10.1016/j.jphotobiol.2015.09.014
73. Jeyaraj M, Rajesh M, Arun R, et al. An investigation on the cytotoxicity and caspase-mediated apoptotic effect of biologically synthesized silver nanoparticles using *Podophyllum hexandrum* on human cervical carcinoma cells. *Colloids Surf B Biointerfaces.* **2013**;102:708–717. doi:10.1016/j.colsurfb.2012.09.042
74. Ahmed MJ, Murtaza G, Rashid F, Iqbal J. Eco-friendly green synthesis of silver nanoparticles and their potential applications as antioxidant and anticancer agents. *Drug Dev Ind Pharm.* **2019**;45:1682–1694. doi:10.1080/03639045.2019.1656224
75. Vazquez A. The universe of normal and cancer cell line responses to anticancer treatment: lessons for cancer therapy. *Nat Preced.* **2011**;1.
76. Kumar V, Sharma N, Maitra SS. In vitro and in vivo toxicity assessment of nanoparticles. *Int Nano Lett.* **2017**;7:243–256.

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