In-vitro cytotoxicity of java tea mediated selenium nanoballs against L6 cell lines

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ABSTRACT

The emergence of Nanotechnology has provided a catholic research in recent years by intersecting with assorted branches of science and forming collision on all forms of life. At present there has been a prodigious excitement in the field of Nano pharmacology to seek the role of Nano selenium in individual healthcare and explains how selenium is a double edged sword in the pathologies of chronic diseases like Diabetes mellitus because of their inferior toxicity and ability to gradually release the bioactive principle and free selenium after ingestion. Thus the present study is an embryonic attempt and was aimed at the adroit synthesis of selenium nanoballs from the aqueous extract of well known herbal tea leaves namely Java tea (Orthosiphon stamineus) using selenious acid solution. The synthesized Selenium nanoparticles were then subjected to various characterization techniques such as UV, FTIR, FESEM, EDAX and Zeta potential respectively. Finally the green synthesized SeNps were tested for their cytotoxic effect against L6 rat skeletal muscle cell lines. The pre clinical studies are underway to prove the insulin mimic activity of the selenium nanoparticles.

Keywords: Selenium Nanoparticles, Java tea, L6 cell lines.

1. INTRODUCTION

The field of Nanotechnology devours great enthusiasm in recent years because of its anticipated impact on science, industry, economy and our everyday life. Today nanoparticles of both metallic and non metallic origin are under research and development in various fields of biology and therapeutics 1. Due to the inimitable characteristics exhibited by nanoparticles, they are employed in nanomedicine and nanotherapeutics which are the supreme aspects of nanotechnology to be implemented in human health.

Metal nanoparticles are vital part of future nanotherapeutics which has wide-ranging applications in diverse areas such as chemistry, physics, and biomedical and material sciences2. They have tremendous applications in the area of catalysis, optoelectronics, diagnostic probes and display devices 3. Production of nanoparticles can be achieved through conventional chemical methods and physical methods4,5. While chemical approaches are the most popular methods for the production of nanoparticles some chemical methods cannot avoid the use of toxic chemicals in the synthesis protocol. Engaging plants in the synthesis of nanoparticles has drawn more interest of workers because it provides single step biosynthesis process, can be advantageous over other biological processes by eliminating the elaborate process of maintaining cell culture6. Plants also render a superior option for synthesis of nanoparticle, as the protocols involving plant sources are free from toxicants; furthermore, natural capping agents are readily supplied by the plants. Selenium (Se), belonging to group 16 of the periodic table is well known for its photoelectric and semiconductor properties. These nanoparticles show biological activity and good adsorptive ability due to interaction between the nanoparticles and NH, C O, COO– and C N groups of proteins7. Selenium nanoparticles have also been developed for applications in medical diagnostics8. Thus selenium nanoparticles caused the great interest of researchers and a variety of synthesis methods have been exploited9.
2. MATERIALS AND METHODS

2.1. Collection of Plant Material

The fresh plant of *(Orthosiphon stamineus)* was collected and was authenticated by Dr. S. Sahaya Sathish, Associate Professor, Department of Botany, St. Joseph’s College, Trichy. Dried Java tea leaves were purchased from Organic farm, Salem and was used as the sample for the biosynthesis of the Selenium nanoparticles.

2.2. Preparation of Extract

The dried leaves were blended using a blender and the powder was stored in a clean glassware container for further analysis. 50 gms of powdered leaves were mixed in 300 ml of distilled water in a clean beaker and was stored in a glass container. The contents were mixed by stirring for 15 minutes. It was kept undisturbed for overnight. After 24 hours the mixture was then filtered using filter paper (whatman no1). The filtrates were then used for further assay. The aqueous extract of the leaves were screened qualitatively for the presence of secondary metabolites using the routine phytochemical analysis.

2.3. Synthesis of Selenium Nanoparticles

For the green synthesis of selenium nanoparticles, 1 ml of plant extract was mixed with 10 ml of 30 mM selenious acid solution along with 200 ul of 40 mM ascorbic acid which was used as an initiator of reduction reaction. Standard positive control was maintained using 1 ml of 0.2% sodium alginate + 10 ml of 30 mM selenious acid and 200 ul of 40 mM ascorbic acid for the synthesis of selenium nanoparticles. 1% *Orthosiphon* leaf extract +200 µl of 40 mM ascorbic acid was used as negative control. The preparations were incubated at room temperature. After 24 hrs of incubation the preparation was centrifuged at 10000 rpm for 30 minutes. The pellet was washed with double distilled water and then with absolute alcohol three times. This washed ethanol pellet was dried overnight. The red selenium nanoparticles were suspended in PBS (pH-7.4) by ultra sonication and then centrifuged. The powder form of the extract was used for further analysis.

2.4. Characterization of Nanoparticles

The formation of selenium nanoparticles was confirmed by the visual observation. The synthesized SeNps were analyzed for the rate of absorption by using UV-Visible spectrometer. The UV-Visible spectral study was done in the range of 200 to 400 nm. FT-IR measurements was carried out for the synthesized SeNps to identify the possible bioactive molecules responsible for the reduction of the selenium and the capping ability of the bio reduced selenium nanoparticles by the aqueous leaf extract of *O. Stamineus* using KBr pellets and the spectra was recorded in the wavelength interval 4000 to 400 cm⁻¹. The FESEM and EDAX analysis revealed average shape, size and elemental composition of the synthesized SeNps. The charge distribution of selenium nanoparticles was confirmed with zeta potential analysis.

2.5. In-vitro Cytotoxicity Analysis

**MTT assay**

The monolayer cell culture of L6 rat skeletal muscle cell lines was trypsinized and the cell count was adjusted to 1.0 x 10⁵ cells/ml using respective media containing 10% FBS. To each well of the 96 well micro titer plates, 100 µl of the diluted cell suspension (50,000 cells/well) was added. After 24 h, when a partial monolayer was formed, the supernantant was flicked off, washed the monolayer once with medium and 100 µl of different test concentrations (10,20,40,80,160,320 µg/ml) of selenium nanoparticles were added on to the partial monolayer in micro titer plates. The plates were then incubated at 37°C for 24hrs in 5% CO₂ atmosphere. After incubation the test solutions in the wells were discarded and 100 µl of MTT (5 mg/10 ml of MTT in PBS) was added to each well. The plates were incubated for 4 h at 37°C in 5% CO₂ atmosphere. The supernantant was removed and 100 µl of DMSO was added and the plates were gently shaken to solublize the formed formazan. The absorbance was measured using a micro plate reader at a wavelength of 590 nm. The percentage of cell viability was calculated using the following formula.

% Cell Viability = (OD of sample/OD of Control) x 100.

3. RESULTS AND DISCUSSION

3.1. Bioactive profiling of Aqueous extracts (O. Stamineus)

**Table 1: Phytochemical screening of Aqueous extract of O.stamineus**

<table>
<thead>
<tr>
<th>TEST</th>
<th>AQUEOUS EXTRACT</th>
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<tbody>
<tr>
<td>ALKALOIDS</td>
<td>++ ++</td>
</tr>
<tr>
<td>FLAVONOIDS</td>
<td>++ ++</td>
</tr>
<tr>
<td>STEROIDS</td>
<td>+</td>
</tr>
<tr>
<td>CARDIAC GLYCOSIDES</td>
<td>+</td>
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<tr>
<td>TERPENOIDS</td>
<td>+</td>
</tr>
<tr>
<td>TANNINS</td>
<td>+++</td>
</tr>
<tr>
<td>SAPONINS</td>
<td>++ ++</td>
</tr>
<tr>
<td>REDUCING SUGAR</td>
<td>+</td>
</tr>
<tr>
<td>ANTHRAQUINONE</td>
<td>++ ++</td>
</tr>
<tr>
<td>PHENOL</td>
<td>+++</td>
</tr>
<tr>
<td>CARBOHYDRATES</td>
<td>++ ++</td>
</tr>
</tbody>
</table>

Bioactive profiling was performed in aqueous extract of *O.Stamineus* leaves. The phytochemical screening showed the presence of alkaloids, flavonoids, amino acids and...
protein, carbohydrate, cardiac glycosides and saponins in aqueous extract of Java tea leaves (Table-1). It is rich in flavanoids, terpenoids and other secondary metabolites having reducing functional groups which might have played a role in reducing selenious acids to SeNPs. *O. stamineus* has wide traditional and pharmacological uses in various pathophysiological conditions. It has been shown that plant extract containing phenol and flavonol derivatives act as reducing agents and nanoparticle stabilizer. FTIR spectrum of java tea extract shows the presence of various functional groups (Figure-3).

### 3.2. Visual Observation

![Visual Observation of Formation of SeNps](image)

Figure 1: Visual Observation of Formation of SeNps

The bio reduction of the selenious acid using aqueous leaf extract of *O. stamineus* was monitored and the appearance of brownish orange colour indicates the formation selenium nanoparticles. Initially the plant extract was light brown in colour. But after the addition of colourless selenious acid solution the colour gradually changes from light brown to dark brownish orange as shown in the (Figure-1).The appearance of brownish orange colour indicates the reduction of selenious acid and formation of SeNps similar to the results reported earlier.

### 3.3. UV-Visible spectroscopy

![UV-Visible Spectra of Se Nanoparticles Synthesized by Using Java tea leaf Extract](image)

Figure 2: UV–Vis Spectra of Se Nanoparticles Synthesized by Using Java tea leaf Extract

The above (Figure-2) shows the UV-Vis Spectra of the as-formed SeNps exhibiting the maximum absorption peak at about 209 nm. This is due to the surface plasmon resonance of selenium nanoparticles. The red color of the colloids implied that selenium form was either amorphous or monoclinic, since trigonal selenium is known to be black.
3.4. Fourier Transform Infrared Spectroscopy (FTIR) analysis

FTIR measurements of both the aqueous java tea leaves extract and the synthesized dried SeNps were carried out to identify the possible biomolecules responsible for the reduction, capping and efficient stabilization of the bio reduced SeNps. The FTIR spectra of both the leaf extract as well as SeNps are shown in (Figure-3). The aqueous java tea leaf extract displays a number of absorption peaks reflecting its complex nature. The spectrum was recorded in the wavelength region between 400 cm\(^{-1}\) to 4000 cm\(^{-1}\). The spectrum of aqueous java tea leaves extract shows the peaks at wave numbers 3388 cm\(^{-1}\), 2928 cm\(^{-1}\), 2856 cm\(^{-1}\), 1637 cm\(^{-1}\), 1385 cm\(^{-1}\), 1321 cm\(^{-1}\), 1265 cm\(^{-1}\), 1158 cm\(^{-1}\), 1069 cm\(^{-1}\) respectively. The broad peak at 3388 was due to the presence of alcohol (O-H) stretching of Phenolic compound. The peaks at 2928 cm\(^{-1}\) and 2856 cm\(^{-1}\) showed a sharp peak and strong alkane C-H stretching. The peak at 1636 cm\(^{-1}\) and 1384 cm\(^{-1}\) showed a variable alkene C=C and alkane (C-H) stretch. The other peaks at 1265 cm\(^{-1}\), 1158 cm\(^{-1}\), 1069 cm\(^{-1}\) were due to amine (C-N), ether, alkene, C-Cl stretching. After the reduction process the absorption peaks at 3381 cm\(^{-1}\), 2960 cm\(^{-1}\), 1718 cm\(^{-1}\), 1629 cm\(^{-1}\), 1516 cm\(^{-1}\) were observed as shown in (Figure-3).Corresponds to alcohol(O-H), alkene (C-H), carbonyl (C=O), alkene C=C and aromatic C=C stretching of Phenolic compound. The peaks at 2928 cm\(^{-1}\) and 2856 cm\(^{-1}\) were disappeared and shifted to 2960 cm\(^{-1}\). The peak corresponds to alkene C=C was disappeared and shifted to 1718 cm\(^{-1}\). A sharp peak at 1384 cm\(^{-1}\) was disappeared and was shifted to 1441 cm\(^{-1}\) corresponds to aromatic C=C stretch. Based on the above observations it is inferred that the biosynthesized SeNps might be surrounded by any one of these organic molecules such as polyphenols, alkaloids, terpenoids which are in accordance with the facts already reported in the earlier reports\(^{16}\).

3.5. FESEM and EDAX analysis

The surface morphology and size of the nanoparticles were obtained by Frontier Scanning Electron Microscopy analysis. The above (Figure-4) shows the SeNPs synthesized by the Java tea leaves extract. The electrostatic interactions and hydrogen bond between the bio-organic capping molecules bond are responsible for the synthesis.
of selenium nanoparticles using plant extract. The image clearly indicates the spherical nature of the selenium nanoparticles as shown in the above figure and they merely look like balls. The relatively uniform shape of the SeNps was confirmed in the range of 88 nm -141nm. The obtained results were also supported in the earlier studies.

An elemental composition analysis employing FESEM-EDAX showed the presence of a strong signal from Se atoms (12.62%) (Figure-5). This analysis indicated that the nanostructures were composed of selenium. Other EDAX peaks such as C, K, Na, Mg, O, Si were also found, suggesting that they were mixed precipitates of the selenium salt.

3.6. Zeta potential value of Selenium Nanoparticles:

The zeta potential measurements indicate negative charge (-34.9 mV) on the selenium nanoparticles as shown in the above (Figure-6). If all the particles in suspension have a negative or positive zeta potential, they will tend to repel each other and there is little tendency for the particles come together. The slightly negative charge on Se particles is probably resulting in the high stability of the selenium nanoparticles without forming aggregates and these particles do not transform to black amorphous form when kept for prolonged period of time of more than a month.

3.7. MTT Assay

Thus the successfully synthesized SeNp was screened for the cytotoxicity against L6 Cell lines by MTT Assay. From the above Table-2, it was understood that the optical density decreases as the concentration of selenium nanoparticles increases, indicating the increase in cell viability. Thus the L6 rat skeletal muscle cells showed...
nontoxic effect and the % of cell viability was notably increased. These results clearly demonstrate that the phytochemicals within these herbs provide nontoxic coating on SeNPs and corroborate the results of the internalization studies discussed above. The lack of any noticeable toxicity of Java tea mediated selenium nanoparticles provides new opportunities for the safe application in molecular imaging and therapy.

In modern science, biomedical technology is well developed and growing in recent times. Biological synthesis of metal nanoparticles is an important role in the field of modern nanotechnology. Green synthesis of nanomedicine plays key role in bio Medical science. In this study, biological synthesis of selenium using java tea leaves and their cytotoxic study is taken into consideration. The nanoparticles obtained are characterized by UV-spectral analysis, FESEM, EDAX, Zeta potential. The functional groups of the selenium nanoparticles in Java tea leaves extract were characterized by Fourier Transform Infrared Spectroscopy. The in vitro cytotoxicity analysis of selenium nanoparticles showed good cell viability against L6 rat skeletal muscle cell line. From this study, it is found that the biomedical properties are present in Selenium nanoparticles.

4. CONCLUSION

In the present study, selenium nanoballs were successfully synthesized using the plant extract of Orthosiphon stamineus. The synthesized nanoparticles were found to increase the cell viability in L6 cell lines. The preclinical studies for treating diabetes mellitus and to prove that the insulin mimic activity of selenium nanoparticles are underway. Thus the green synthesized nanoparticles were found to be cost-effective, simpler, and environmentally safe. As the nanotechnology is an emerging field in medicine, the biological synthesis of nanoparticles helps in the other way. In Future, it is a promising contender for various clinical applications in a safe and ecofriendly manner.

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