Original Research Article

Fabrication of zero-valent iron nanoparticles by green and chemical reduction methods: Application in the field of antibacterial activities for medicinal point of view

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A B S T R A C T

In the area of life sciences, iron nanoparticles (Fe NPs) have many applications. In this paper, the unique properties of iron nanoparticles as antimicrobials are studied. In this study, nanoparticles of iron have been fabricated by green and chemical reduction method. With the help of FESEM analysis and Zeta size analysis, the usual value of nanoparticles was found to be 10-30 nm in size. Furthermore, the prepared nanoparticles were examined for antibacterial perspective aligned with gram-positive and negative strains namely Staphylococcus aureus & Bacillus subtilis and Escherichia coli, using agar plate method and IC50 was also estimated using tube dilution assay.

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1. Introduction

Nanotechnology is an extremely convincing technology with its approaching medical properties in early disease detection and treatment. At present, there is a growing concern in the research of nano-metals as these signify an intermediate dimension between atoms/molecules and bulk particles.1 Among metal nanoparticles, iron oxide has received more attention because of its variety of scientific and technological applications such as antimicrobial activity2, biosensor3, food preservation4, ferrofluids, magnetic resonance imaging, magnetic refrigeration, magnetic storage media, cell sorting, targeted drug delivery, and hyperthermia cancer treatments.5–7

The size of nanoparticles makes them an attractive applicant for in vivo and in vitro biomedical research. The incorporation of nanoparticles in the medical field has led to their special use in the field of sensing, drug delivery, imaging, and artificial implants. An interesting new approach for their investigation in the medical field is their application in antimicrobial activities that aim at drug-resistant and highly pathogenic microbes.8

There is also an increasing interest in the application of Iron and other metal nanoparticles in blood platelet sanitization. Ag+ ions may slow down some imperative transport processes such as succint and phosphate uptake, by contacting the cellular oxidation process in the respiratory string. The high absorption of Ag+ not only causes toxicity but may also lead to electrolyte imbalances in patients.9

In recent research, it has been shown that the better antibacterial properties of iron nanoparticles (Fe NPs) are dependent upon the high surface area and high volume of nanoparticles that allow proficient bacterial disinfection.10,11 Iron nanoparticles (Fe NPs) have a larger surface area compared to silver so that they possess better antimicrobial properties as compared to Ag+ ions.12–15

Despite this, in an aqueous solution zero-valent iron
nanoparticles (Fe NPs) can inactivate E. coli. Which increases the interest of iron nanoparticles in antimicrobial activity. In the deficiency of oxygen, iron nanoparticles (Fe NPs) show high antimicrobial activity analogous to that of silver nanoparticles (Ag NPs). The strong antibacterial activity of iron nanoparticles (Fe NPs) implies that Fe NPs can serve as a cost-effective biocide for many of the applications in which silver is being used. In the case of antibacterial activity, iron nanoparticles (Fe NPs) involve the release of Fe ions on the biological obligatory site that hold upon to the surface of the targeted atom and developed a bacterial interaction with the NPs that starts from the surface layer.

2. Materials and Methods

2.1. Green method

2.1.1. Fabrication of Fe NPs using S. oleracea aqueous leaf extract

Fe NPs were fabricated by using fresh leaves of S. oleracea extract as a reducing agent. For the fabrication of Fe NPs, fresh leaves of S. oleracea were washed, dried at room temperature, and cut into small pieces. Dried-out leaves are crushed into powdered form and from which 4g was mixed in 50 ml of double distilled water and then boiled for 15 min at 100°C. Filter the extract and stored it at 4°C. Furthermore, 10ml of S. oleracea extract was added into 10ml aqueous solution of 1.0 mM ferric chloride. The mixture was boiled for about half an hour and after which black color was observed, representing the fabrication of Fe NPs.

2.2. Chemical reduction method

2.2.1. Fabrication of Fe NPs using NaBH₄ as reducing agent

In a three-neck, round-bottom flask 0.1M FeCl₃.6H₂O (4.1703 g) and 0.05M EDTA (3.7224 g) were mixed in 100 ml Milli-Q water by using appropriate mixing. Then the solution of NaBH₄ (0.75M NaBH₄(2.837 g) in 100 ml Milli-Q water) was added dropwise into the above solution. Slowly the solution changed to black color.

3. Results and Discussion

3.1. Characterization of Fe NPs

3.1.1. Analysis by ultraviolet-Visible spectra

An ultraviolet-Visible spectrum is an effectual technique that illustrates the development of varieties of metals in the creation of colloidal M NPs. As exposed in Fig. 1a the optical absorbance Plasmon band of the Fe NPs using S. oleracea extract in the colloidal suspension is about 273 & 224. In both cases, the black color appeared due to the formation of Fe NPs.

3.1.2. Zeta size measurement

The zeta potential value (21.9) shows the stability of Fe NPs using S. oleracea extract and confirms the size of Fe NPs particles to be 80.37(d. nm). On the other hand, the zeta potential value (-21.2) of Fe NPs using classical reduction method with EDTA and found the size of Fe NPs to be 51.37 (d. nm) (Fig-2: A&B).

3.1.3. XRD of Fe NPs by chemical reduction and green method

The aqueous solution of Fe NPs was frequently centrifuged at 10,000 rpm for 30 min, redispersed with distilled water, and lyophilized to get pure Fe NPs pellets. The Debye-Scherer principle has been used to evaluate the standard particle size that could be understood as:

\[ D = \frac{0.91}{\beta \cos \theta} \]

An average dimension of Fe NPs was found in the green method is 55.18 nm and in the chemical reduction method 23.84 nm by XRD methods. (Fig-3: A&B)
3.1.4. EDX of Fe NPs by chemical reduction and green method

EDX is Energy Dispersive X-Rays (aka EDAX - energy dispersive x-ray analysis) and is reliant on the atomic mass of the elements being detected. In the fabrication of Fe NPs in the chemical reduction method the wt% value of Fe is 31.7, O is 25.3 and Cl is 0.7 and in the case of the green method the wt% value of Fe is 30.9, O is 16.2 and Cl is 2.7. The peak of Fe presents in the spectrum shows the fabrication of Fe NPs.

3.1.5. FE SEM analysis of Fe NPs

The FE SEM image shows that the size of indusial Fe NPs was found to be around 50-60 nm using S. oleracea extract confirms the size of Fe NPs particles to be 51.22 (d. nm). On the other hand, the Fe NPs using the classical reduction method with EDTA was found to be 47.97 (d. nm) (Fig-5).

3.2. In-vitro antibacterial studies of nanoparticles

The iron nanoparticles produced by the liquid phase reduction technique were subjected to the in-vitro antibacterial activity of iron nanoparticles aligned with gram-negative (E. coli) and gram-positive (S. aureus and B. subtilis) respectively. Iron nanoparticles showed excellent and dose-dependent activity when compared to a standard antibiotic. The effects obtained after 24h incubation are presented in table 1 and fig6. Test compound the highest region of prohibition at 500μg/mL was 21 ± 0.64 mm as compared to standard ampicillin.

<table>
<thead>
<tr>
<th>Test compound</th>
<th>E. coli Zone of inhibition (mm)</th>
<th>S. aureus Zone of inhibition (mm)</th>
<th>B. subtilis Zone of inhibition (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ampicillin (Standard) (80μg/mL)</td>
<td>20 ± 0.56</td>
<td>15 ± 0.23</td>
<td>18 ± 0.85</td>
</tr>
<tr>
<td>200μg/mL</td>
<td>12 ± 0.54</td>
<td>13 ± 0.47</td>
<td>14 ± 0.76</td>
</tr>
<tr>
<td>300μg/mL</td>
<td>16 ± 0.36</td>
<td>15 ± 0.22</td>
<td>15 ± 0.51</td>
</tr>
<tr>
<td>400μg/mL</td>
<td>18 ± 0.76</td>
<td>17 ± 0.61</td>
<td>17 ± 0.33</td>
</tr>
<tr>
<td>500μg/mL</td>
<td>19 ± 0.47</td>
<td>20 ± 0.72</td>
<td>21 ± 0.64</td>
</tr>
</tbody>
</table>

3.2.1. Tube dilution assay

In tube dilution assay, prepared nanoparticles showed significant antibacterial activities against targeted bacterial strains. IC$_{50}$ of test compound was found to be 350 μg/mL, 300 μg/mL, and 400 μg/mL against bacterial strains compared with IC$_{50}$ of standard ampicillin was found to be 40 μg/mL, 50 μg/mL, and 50 μg/mL against pathogenic E. coli, S. aureus and B. subtilis respectively (Fig. 2a & 2b). Test compounds have shown better activities against B. subtilis and it was observed that it showed maximum ZOI and minimum IC$_{50}$ (300 μg/mL) in both in vitro models. It indicates that it can be useful in the management of various B. subtilis mediated infections. The results of biological activity are revealed that iron nanoparticles found active against both strains but slightly better against gram-positive
strains. The findings of the study were found in agreement with previously published reports (Behera et al. 2012).

Fig. 7: Statistical significance was assessed by ANOVA test and compared with control (p < 0.0001) which denoted as (***) and (p > 0.05) which denoted as (*).

3.3. Conclusion

Today, iron nanoparticles’ production with the help of plant materials or in an eco-friendly manner is considered attractive. The attrition of metallic ions into base metals by the biological method is a rapid and more efficient method than conventional methods. This method is eco-friendly and can be conducted willingly at room temperature and is also amplified easily. The reducing agents drawn in this method than conventional methods. This method is eco-friendly and can be conducted willingly at room temperature and is also amplified easily. The reducing agents drawn in this

4. Source of Funding

None.

5. Conflict of Interest

None.

References