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ANTIMICROBIAL ACTIVITY AND PHYTOCHEMICALS
STUDY OF AVERRHOA CARAMBOLA AND SYNTHESIS
OF SILVER NANOPARTICLES

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ABSTRACT
Starfruit botanically known as Averrhoa carambola and it belongs to family Oxalidaceae, grows in tropical and subtropical regions throughout the world. The Starfruit can be used raw as vegetable and ripe as fruit. The Starfruit are sweet tasting fruit that possesses high nutritional value. This study was aimed to understand the antimicrobial activities and phytochemical study of tropical fruits Averrhoa carambola (starfruit). The edible parts of the fruits were analyzed for different phytochemicals were found in starfruits. In the present study, information regarding the extracts of Averrhoa carambola were evaluated to investigate antimicrobial activity against Gram positive and Gram negative bacteria by well diffusion method. Phytochemical were present such as saponins, phenols, tannins, steroids, proteins, carbohydrates and flavonoids. In this study well defined silver nanoparticles were synthesized by using carambola fruit extract. The synthesized NPs were analyzed by ultraviolet spectroscopy. The antimicrobial activity of the synthesized AgNPs was obtained against Escherichia coli and Pseudomonas aeruginosa by agar well diffusion method.

KEY WORDS: Averrhoa carambola, Phytochemicals, Nanoparticles.

1. INTRODUCTION
Averrhoa carambola, commonly known as starfruit bears a grt significance in traditional medicine. Five-lobbed fleshy, yellow-greenish, edible fruits of Averrhoa carambola of Oxalidaceae are native of South-East Asia and it cultivated in some parts of India. Averrhoa carambola is a small, slow-growing evergreen tree with a short-trunk or a shrub. The compound leaves are soft, medium-green, they are spirally arranged around the branches in an alternate fashion. The fruits are showy with an oblong shape. The fruits have a thin, waxy skin that is orange-yellow colored. A yellow or green tropical fruit with smooth skin and five pointed, curved parts, making a star shape when you cut through it. The juicy fruits are yellow inside when ripe and have a crisp texture and when cut in cross-section are star shaped. Starfruit generally stored at room temperature for maximum of two to five days. Carambola are not too a particular soil, it grows well on sand, heavy clay or limestone and in rich loam.

The fruits are good source of antioxidants and used traditionally in mouth ulcers, toothache, nausea, diarrhea, ascites etc. Pharmacological investigations on Averrhoa carambola have demonstrated anti-inflammatory, anti-microbial, anti-fungal, anti-tumor and anti-ulcer activities. In addition, the plant possesses hypocholesterolemic investigations have shown the presence of Phytochemical such as saponins, tannins, alkaloids and flavonoids. Anti-
bacterial activities of extracts of different plants against various microorganisms have been reported. Plants have always been a significant source of natural products having therapeutic potential. In India, rich resource of wild or underutilized fruits is available. These underutilized fruits have recently drawn attention of many researchers as a natural source of treatment for curing various diseases. Some studies on underutilized fruits have claimed them to be better sources of nutrients. Such underutilized tropical fruits provide limitless opportunities for screening of novel drugs. Phenolics possess a wide spectrum of biochemical activities such as antioxidant, anti-mutagenic, anti-carcinogenic. Flavonoids are the largest group of naturally occurring phenolic compounds, which occurs in different plant parts both in free state and as glycosides. The Ascorbic acid levels of the starfruit is believed to be responsible for its sweet or sour taste.

Present study has been aimed to understand the characterization of antimicrobial activity and phytochemicals study of tropical Averrhoa carambola (starfruit) as well as In this study well defined silver nanoparticles were synthesized by using carambola fruit extract. we were reporting the synthesis and characterization of silver nanoparticles by using carambola fruit extract. The synthesized AgNPs were characterized by using characterization techniques and tested against several pathogenic microorganisms.

2. MATERIALS AND METHODS

2.1 Sample collection and preparation

Fruits of Averrhoa carambola (starfruits) were collected in September 2017, from markets of Dadar, Mumbai, Maharashtra. The fruits were cleaned. The edible portions of the fruits were dried at 50 ºC for 2-3 days, then separately grind into fine powder using a mechanical grinder. The powder was kept in dark coloured glass bottles and subsequently used.

2.2 Preparation of starfruit extract:

5 g of dry powder was mixed in 50 ml sterile methanol (80%). Keep 2 hours at room temperature using orbital shaker at 150 Rpm. Then extract was filtered through Whatman No.1 filter paper. After that extract was evaporated using vacuum rotary evaporator for purpose of dryness and stored in glass vials. These crude solvent extract were diluted with 10% dimethyl sulphoxide (DMSO) which are to be used as negative control.

2.3 Antimicrobial activity:

2.3.1 Test Microorganisms

Bacterial strains were selected for the antimicrobial activities of these fruits study. The strains were used in Salmonella typhimurium, Staphylococcus aureus, Escherichia coli, Klebsiella pneumoniae, Bacillus cereus, All bacterial cultures were maintained on tryptic soy agar (HiMedia) and subcultured regularly. The fungal strain candida albicans was grown on Sabouraud dextrose agar (HiMedia). This culture was incubated at 35-37 ºC for 24 hours.

2.3.2 Well diffusion bioassay

Standardized inoculum suspension (0.1ml) of each bacterial strain was spread on Muller Hinton Agar plates with a sterile bent glass rod spreader. Then punch agar plate with a sterile cork borer of 4 mm size and then pour 100 µL of each sample with micropipette in the bore. Allow plates to stand for 30 min. Then incubate the plates at 37ºC for 24 h. After incubation measured the zone of inhibition in (mm).

2.4 Phytochemical analysis:

2.4.1 Test for carbohydrates (Fehling’s test): 1 ml of Fehling’s A and 1ml of Fehling’s B solution was added in 0.5 mg of extract and boil it in a water bath. The formation of yellow or red precipitate indicates the presence of reducing sugar.

2.4.2 Test for Saponins (Foam test): Diluted the methanolic extract with distilled water and shake well in a graduated cylinder for 15 min. The persistent foam to a length of 1cm indicates the presence of Saponins.

2.4.3 For steroids and sterols (Salkowski’s test): Dissolved 2 ml of methanolic extract in 2 ml of chloroform and 2ml of concentrated sulphuric acid along the sides of the test tube. The upper layer turns red and lower layer turns yellow with fluorescence, indicates the presence of the steroids and sterols compound in the extract.

2.4.4 Test for tannins (lead acetate test): To 2-3 ml of extract, was added in 0.5 ml of 1% lead acetate and the formation of white precipitate indicates the presence of tannins and phenolic compounds.

2.4.5 Test for amino acid (Ninhydrin Test): To a small amount of extract add a few drops of 5% Ninhydrin solution. Then heat the solution in a water bath for 10 mins. Yellow color appeared if amino acids are present.

2.4.6) Test for Terpenoids: Add 4 mg of extract treated with 0.5 ml of acetic anhydride and 0.5 ml of chloroform. Then add concentrated solution of sulphuric acid slowly and red violet color will observe for terpenoid.

2.4.7) Test for flavonoid: Crude extract mix with 2ml of 2% solution of NaOH. An intense yellow color formed which turn colorless on addition of few drops of diluted acid which indicated the presence of flavonoids.

2.4.8) Test for phenol: Crude extract was mixed with 2ml of 2% solution of FeCl₃.A green or black coloration indicates the presence of phenol.

2.5 Synthesis of silver nanoparticle and its characterization:

Starfruit was washed with distilled water and cut into small pieces. The aqueous fruit extract was prepared 5 g of fruit pieces with 100 ml distilled water
boiled for 15 min. The extract was cooled at room temperature and filtered through whatman filter paper No.1. The filtrate was collected and used for further experiment of synthesis of AgNPs. A stock solution of 4 Mmol L⁻¹ AgNO₃ will be prepared. 10 ml of fruit extract taken and 25 ml AgNO₃ stock solution and constant stirring at 40°C. The colourless fruit extract solution changes to reddish brown slowly indicating the formation of AgNPs. And to check the bactericidal activity against Escherichia coli, Pseudomonas aeroginosa by using agar well diffusion method. The synthesized AgNPs were characterized by using various analytical techniques. The reduction of the pure Ag⁺ ions was monitored by measuring the absorbance of the reaction medium with UV spectrophotometer.

3. RESULTS
3.1 Antimicrobial Activity:

Table No. 1: Antibacterial activities, indicated by diameter of inhibition zone of selected samples against the micro-organisms.

<table>
<thead>
<tr>
<th>Fruit</th>
<th>Bacillus cereus</th>
<th>klebsella pneumoniae</th>
<th>Escherichia coli</th>
<th>Staphylococcus aureus</th>
<th>Salmonella typhimurium</th>
<th>Candida albicans</th>
</tr>
</thead>
<tbody>
<tr>
<td>Averrhoa carambola</td>
<td>20mm</td>
<td>17mm</td>
<td>12mm</td>
<td>14mm</td>
<td>10mm</td>
<td>09mm</td>
</tr>
</tbody>
</table>

3. Phytochemicals 2

The edible parts of the fruits were analyzed for different phytochemicals (Table 2). The phenolics, flavonoids, tannin, steroids, carbohydrates, and proteins were present in Averrhoa carambola as well as terpenoids were absent in Averrhoa carambola as shown in Fig.No.1.

Table 2: Phytochemical qualitative evaluation of Averrhoa carambola

<table>
<thead>
<tr>
<th>Phytochemicals</th>
<th>Averrhoa carambola</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carbohydrates</td>
<td>Present</td>
</tr>
<tr>
<td>Saponins</td>
<td>Present</td>
</tr>
<tr>
<td>Steroids</td>
<td>Present</td>
</tr>
<tr>
<td>Tannins</td>
<td>Present</td>
</tr>
<tr>
<td>Amino acids</td>
<td>Present</td>
</tr>
<tr>
<td>Terpenoids</td>
<td>Absent</td>
</tr>
<tr>
<td>Flavanoids</td>
<td>Present</td>
</tr>
<tr>
<td>Phenols</td>
<td>Present</td>
</tr>
</tbody>
</table>
3.3 Synthesis of Nanoparticle And its characterization

Synthesis of AgNPs as shown in figure 2 (A). AgNPs are extensively used in the pharmaceutical industries and have inhibitory activities on various microorganisms. Biosynthesized AgNPs were analyzed for their antibacterial activity against organism *Escherichia coli* and *Pseudomonas aeruginosa* by agar well diffusion method. The antibacterial activity was assayed by measuring the diameter of zone of inhibition around the well. The images for antibacterial activity of biosynthesized AgNP’s against *Escherichia coli* and *Pseudomonas aeruginosa* were shown in figures 2 (b) and (c). The Table 3 shows summarized results of antibacterial activity.

![Figure 2: A) Silver Nanoparticles, B) Antibacterial activity of biosynthesized against (b) *Escherichia coli* And (c) *Pseudomonas aeruginosa.*](image)

### Table 3. Antibacterial activity of AgNPs.

<table>
<thead>
<tr>
<th>Test microorganism</th>
<th>Carambola Fruit Extract</th>
<th>AgNPs (Undiluted)</th>
<th>AgNPs (Diluted)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Escherichia coli</em></td>
<td>11 mm</td>
<td>14 mm</td>
<td>12 mm</td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em></td>
<td>0 mm</td>
<td>13 mm</td>
<td>11 mm</td>
</tr>
</tbody>
</table>

![Figure 3. UV-absorption spectra of AgNPs](image)
UV–spectral analysis of AgNPs the formation of AgNPs in an aqueous colloidal solution was investigated by using UV–spectrophotometer analysis as shown in Fig.3. AgNPs turned yellowish brown in the aqueous solution, which has been reported and check the absorbance with the help of UV spectrophotometer. It has been noticed that the absorption peak width gradually became narrower with time, and this suggests the narrow size distribution of newly formed AgNPs. Characterization of silver nanoparticles is done by UV-Spectrophotometer. The maximum absorption of silver nanoparticles was found in 450nm.

4. CONCLUSION
It could be concluded that Averrhoa carambola is an excellent plant due to its multifaceted medicinal properties like, antimicrobial, antioxidant activity and nutritional content. The tested extract of fruits was shown antimicrobial efficacy against most of microbes examined. Agar well diffusion assay was showed the fruit extracts have different degrees of bacterial and fungal growth inhibition depending on the strain. The edible parts of the fruits were analyzed for different phytochemicals like phenolics, flavonoids, alkaloids, Tannins, steroids, present in starfruits. Synthesis of AgNPs with the help of starfruit is the industrial application. Synthesize AgNPs was showed the better bactericidal activity and characterization of silver nanoparticles is done by using UV Spectrophotometer.

5. DISCUSSION
In this study discussing and determining the antimicrobial activity using different strains as well as study the synthesis and characterization of silver nanoparticles. Characterization of silver nanoparticles with the help of UV spectrophotometrically.

6. REFERENCES