PHYTOCHEMICAL AND ANTIBACTERIAL ACTIVITY OF DIVERSE SOLVENT EXTRACT OF LEAF (Plumbago zeylanica).

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ABSTRACT
The present investigation is focused to assess the Phytochemical study and antibacterial activities of the leaf extract of Plumbago zeylanica. The leaf extracts were prepared by using various solvents with an increasing polarity such as hexane, chloroform, ethyl acetate and Methanol. The filtered extract was assessed for its activity against multifarious bacteria such as Staphylococcus aureus, Klebsiella pneumoniae, Bacillus cereus, Salmonella typhi and Pseudomonas aeruginosa by performing well diffusion method. Out of the four solvents trialled, ethyl acetate extract alone showed Zone of inhibition and confirmed significant activity against all the tested pathogens.

Keywords: P. zeylanica, Phyto extract, Antibacterial activity, Well diffusion method.

INTRODUCTION
Indiscriminate use of antibiotics has led to an increased percentage of many defenses against a human harmful microorganism in recent times. Specific plants have restricted Heritance along with universal significance. Our country was blessed with a wealthy origin of medicinal plants. They form the main ingredients of the Indian system of medicine. Various forms of drug modified elements have been isolated from natural resources, especially plants. These kinds of the modern drug have been used to cure diseases in all places of the world. One such plant is Plumbago zeylanica belonging to family Plumbaginaceae. Its flowers are used as Digestant, leaves are used to treat scabies and dysentery. The Root is pungent and has been used as Abortifacient, good appetizing agent, expectorant, tonic, useful in rheumatoid arthritis, detoxifier, skin infection, and disease of the spleen.² Antiviral, Antifungal, Antibacterial, Antipneumococcal, Leishmaniasis, Trypanophobia, and Anticarcinomic activity of different parts of P. zeylanica has been reported.³ Plumbago zeylanica comprises of important chemical composites for instance alkaloids, Napthaquinones, steroids, tannins, Glycosides, Triterpenoids, Flavonoids, Coumarins, Carbohydrates, Saponins, phenolic compounds, fats, fixed oil and proteins.⁴ A Polyphenolic compound from this plant possess Apoptotic, antibacterial, antioxidant and anticancer properties.⁵ Unfortunately, new lead molecules have not been synthesized so far. Hence, there is a need for in-depth research to know the mode of action, Bioactivity of various phytochemicals present and its medicinal values. The present study focused on Plumbago zeylanica leaf extract using four different solvents based on polarity and its Phytochemical analysis and antibacterial activity.

EXPERIMENTAL
Solvent Extraction of Plant
The leaf of P. zeylanica was purchased from the homeopathic store. It was washed and shade dried for 15 days, then ground into powder. 100g of powder was mixed with 200ml of solvents as per polarity (Hexane, Chloroform, Ethyl acetate and Methanol). The Blending of the mixture was done by keeping the flask on the orbital shaker at room temperature for 48 hours. It was filtered by Whatman No-1 filterpaper and the filtrate was collected. Solvent from the filtrate was evaporated by rotary evaporator at low
temperature under medium pressure condition. The dried crude extract was stored in the airtight container for further experiment and analysis.

**Phytochemical Screening**

The standard protocol was used to screen the presence of phytochemical constituents like carbohydrate, alkaloids, glycosides, saponins, phytosterols, the phenolic compound, flavonoids, amino acids, and tannins present in the plant extract. The observation of the phytochemical screening results was summarized in the Table-1.

**Test for Alkaloids**

1ml of plant leaf extract diluted, mixed with 5ml Hydrochloric acid and was filtered. 1ml of Mayer’s reagent added to the mixture to observe the change yellow precipitate formation.

**Test for Carbohydrates**

1ml Plant leaf extract was added with 2ml of Fehling solution A and B and then boiled for 5 minutes. The red deposit showed the presence of reducing sugars.

**Test for Protein**

Two drops of Ninhydrin solution added with plant extract. This reaction produces a purple colour indicates the presence of amino acids.

**Test for Flavonoids**

4ml of plant leaf extract was allowed to mix with 1.5ml of fifty per cent methanol solution. This mixture was heated along with metal magnesium then add 5-6 drops of concentrated HCl were added and orange colour was observed for Flavones and Red color for Flavonoids.

**Test for Saponins**

2ml of leaf extract of the plant was added to 5 ml of distilled water and kept on a shaker for 20 minutes. Determination of the foam indicated presence of Saponine.

**Test for Steroid**

0.5ml of plant leaf extract mixed well with acetic anhydrous. When this mixture cooled in an ice bath, add concentrated sulfuric acid. The Reddish brown colour identified the presence of steroid.

**Test for Terpenoids**

5ml of plant leaf extract was treated in 2ml of chloroform and 3 ml of concentrated H2so4 added dropwise. The reddish-brown layer formed at the interface, thus shows the presence of Terpenoids.

**Test for Tannins**

1ml of Plant leaf extract was added with 1ml ferric chloride. A greenish-brown precipitate showed the presence of tannins.

**Test for Phenol**

1ml of Extract treated with 2ml of distilled water and add a few drops of 10% aqueous Ferric chloride. The presence of green or blue colour showed the existence of phenols.

**Antibacterial Effect of P. Zeylanica**

Antibacterial activity of leaf extracts was tested by the good diffusion method up against *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Bacillus cereus*, *Salmonella typhi*, and *Pseudomonas aeruginosa*. Muller Hinton Agar was used for testing. Lawn culture of the bacteria was obtained on MHA using a sterile cotton swab. Wells was punctured using a cork borer. Leaf extract was added (25 µl/µg, 50 µl/µg, 75
µl/µg, and 100 µl/µg) in the well along with the control. Plates were incubated at 37°C overnight. Plates were observed in the zone of inhibition around the well. The zones were measured on Millimetre scale. 

**RESULTS AND DISCUSSION**

**Phytochemical constituents of the leaf extract of** *P. Zeylanica*

The result of preliminary phytochemical screening carried out by solvent extraction using Hexane, chloroform, ethyl acetate and methanol leaf extracts of *p. Zeylanica* is given in the Table-1. Carbohydrates are in the Hexane extract. Flavonoids are present in the Hexane, ethyl acetate and methanol extract. Alkaloids, Terpenoids and phenolic compounds are found in the chloroform and ethyl acetate extract. Saponins are in the methanol extract. At last tannins were found in the chloroform, ethyl acetate and methanol extract.

Polyphenols are a large and a contrasting form of compounds, many of which exist a naturally inaccurate range of herbs and food molecules. Flavonoids are one of the best polyphenols. Many of which have properties such as antioxidant, antimutagenic, Anticarcinogenic and anti-inflammatory effects. These kinds of polyphenols potentially inhibit the disease and control the strength of the genome.

**Antibacterial Activity**

Extracts were tested for antibacterial activity by the good diffusion method. Among the four solvents used for extraction, only ethyl acetate extracts showed a zone of inhibition against bacteria tested (*Staphylococcus aureus, Klebsiella pneumoniae, Bacillus cereus, Salmonella typhi, and Pseudomonas aeruginosa*). As the concentration of the extracts increases the zone of inhibition also increased. Zone sizes were shown in the Table-2 and Figs.-1 to 5.

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Phytocomponents Tests</th>
<th>Hexane</th>
<th>Chloroform</th>
<th>Ethyl Eacetate</th>
<th>Methanol</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Alkaloids Mayer’s test</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>2.</td>
<td>Carbohydrates Molisch test</td>
<td>_</td>
<td>+</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>3.</td>
<td>Proteins Ninhydrin test</td>
<td>_</td>
<td>+</td>
<td>-</td>
<td>-</td>
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<tr>
<td>4.</td>
<td>Flavonoids Alkaline test</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>5.</td>
<td>Saponins Froth forming test</td>
<td>_</td>
<td>+</td>
<td>-</td>
<td>+</td>
</tr>
<tr>
<td>6.</td>
<td>Steroid Steroid test</td>
<td>_</td>
<td>+</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>7.</td>
<td>Terpinoids Terpinoids test</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>-</td>
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<tr>
<td>8.</td>
<td>Tannins Fecl 3 test</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>9.</td>
<td>Phenolic compound Lead acetate test</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

(-) Absent, (+) Present.

**Table-2: Antibacterial Activity of** *Plumbago zeylanica* **against** *Salmonella typhi, Bacillus cereus, Pseudomonas aeruginosa, Staphylococcus aureus* and *Klebsiella pneumoniae.*

<table>
<thead>
<tr>
<th>S. No.</th>
<th><em>Plumbago zeylanica</em> Organic Solvent Extract</th>
<th>Microorganism</th>
<th>25µl/µg</th>
<th>50 µl/ µg</th>
<th>75 µl/ µg /</th>
<th>100 µl/ µg</th>
<th>Control Ethyl Acetate µl/ µg</th>
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<tbody>
<tr>
<td></td>
<td><strong>Zone of Inhibition in mm</strong></td>
<td><strong>Zone of Inhibition in mm</strong></td>
<td><strong>Zone of Inhibition in mm</strong></td>
<td><strong>Zone of Inhibition in mm</strong></td>
<td><strong>Zone of Inhibition in mm</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1.</td>
<td>Ethyl Acetate Extract</td>
<td><em>Salmonella typhi</em></td>
<td>10±0.1</td>
<td>12±0.5</td>
<td>13±0.5</td>
<td>12±0.73</td>
<td>No Zone of Inhibition</td>
</tr>
<tr>
<td></td>
<td><em>Bacillus cereus</em></td>
<td></td>
<td>9±0.6</td>
<td>10±0.5</td>
<td>12±0.7</td>
<td>15±0.46</td>
<td>No Zone of Inhibition</td>
</tr>
<tr>
<td>Phytoconstituents</td>
<td>Pseudomonas aeruginosa</td>
<td>Staphylococcus aureus</td>
<td>Klebsiella pneumoniae</td>
<td></td>
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<tr>
<td></td>
<td>6±0.6</td>
<td>8±0.5</td>
<td>6±0.43</td>
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<tr>
<td></td>
<td>7±0.56</td>
<td>10±0.5</td>
<td>9±0.66</td>
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<tr>
<td></td>
<td>16±0.5</td>
<td>14±0.53</td>
<td>10±0.46</td>
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<tr>
<td></td>
<td>17±0.5</td>
<td>16±0.63</td>
<td>15±0.76</td>
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</tbody>
</table>

**CONCLUSION**

Phytoconstituents analysis of leaf extract of *P. zeylanica* confirmed the presence of carbohydrates, alkaloids, glycosides, saponins, phytosterols, phenolic compounds, flavonoids, amino acids, and tannins. Among the four solvent extracts tested for antibacterial activity, Ethyl acetate showed the varying zone of inhibition.
inhibition depending on the concentration of the extract. Hence this study concludes that the ethyl acetate extract can be subjected to further study.

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REFERENCES

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