In vitro Antioxidant Potential of Different Parts of Oroxylum indicum: A Comparative Study

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The present study evaluated the in vitro antioxidant potential of different parts of Oroxylum indicum. 2,2-diphelyl 1-picrylhydrazyl (DPPH), nitric oxide, superoxide anion and hydroxyl radical scavenging potential and reductive ability assay of methanol extract of different parts i.e. root, root bark, stem, stem bark, leaves and fruits were performed. Leaves and bark extracts exhibits highest free radical scavenging activity than bark, stem and fruit extract. Leaves extract showed maximum reductive ability and found to contain maximum amount of polyphenolic compounds. The highest free radical activity may be due to presence of polyphenolic compounds.

Key words: Free radical scavenging, Oroxylum indicum, different parts, polyphenolic compounds

Antioxidants are now standing on the mainstay of the treatment and prevention of several diseases[1-3]. Current research is directed towards finding naturally occurring antioxidants particularly of plant origin. Oroxylum indicum Vent. (Bignoniaceae), a rare endangered and threatened medicinal plant widely used traditionally for treating several disorders[4]. The root-bark is used as an astringent and tonic and also in diarrhoea and dysentery. The stem bark is used in acute rheumatism. In the form of an infusion, it is used as a diaphoretic. The fruits are used as carminative and stomachic, while the seeds are used as purgative. The roots are used in dropsy and the leaves are reputed as an emollient. Tender fruits are described as carminative and stomachic[5]. The root of this plant is also one of the important ingredients in most commonly used ayurvedic formulations like dantyadyarista, brahma rasayana, dasamula, amartarista, dhanawantara ghrita, narayana taila[6]. The anti cancer potential of different parts of the plant has already been reported[7,8]. The present study describes a comparative evaluation of different parts of Oroxylum indicum for their in vitro antioxidant activity.

The different parts of the plant i.e. root, root bark, stem, stem bark, leaves and fruits were collected from the forest region of Orissa and identified at the Institute of Materials and Minerals Technology, Bhubaneswar, Orissa. All the plant materials were shade dried, powdered, sieved and successively extracted with petroleum ether and methanol to obtain the extracts. The each of these extracts was concentrated in a rotary evaporator under reduced pressure, giving individual extracts. Ten milligrams of methanol extract of different parts was dissolved in methanol (1 ml) and solution was serially diluted for antioxidant studies.

Total polyphenolic compounds of different extracts were performed according to the method of Slinkard and Singleton[9]. DPPH radical scavenging activity[10], nitric oxide scavenging assay[11], superoxide
scavenging assay\(^\text{[12]}\), hydroxyl radical scavenging assay\(^\text{[13]}\) and reducing power assay\(^\text{[14]}\) were performed as per the standard procedure.

Total polyphenolic content as pyrocatechol equivalent was determined from the standard curve equation of pyrocatechol and presented in Table 1. Root bark and leaves extract were found to contain maximum amount of polyphenolic compounds. The results of different radical scavenging activity are shown on Table 2. Leaves extract also showed maximum reducing power when tested in reducing power assay.

**TABLE 1: TOTAL POLYPHENOLIC CONTENT OF DIFFERENT PARTS OF OROXYLUM INDICUM**

<table>
<thead>
<tr>
<th>Plant part extract</th>
<th>Phenolics as pyrocatechol equivalents (µg/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>RE</td>
<td>35.65±3.25</td>
</tr>
<tr>
<td>RBE</td>
<td>105.56±4.78</td>
</tr>
<tr>
<td>SE</td>
<td>21.03±2.51</td>
</tr>
<tr>
<td>SBE</td>
<td>80.36±5.03</td>
</tr>
<tr>
<td>LE</td>
<td>124.7±4.36</td>
</tr>
<tr>
<td>FE</td>
<td>21.03±1.15</td>
</tr>
</tbody>
</table>

Re is root extract, RBE is root bark extract, SE is stem extract, SBE is stem bark extract, LE is leaves extract and FE is fruit extract. Values are mean±standard deviation of n=3 determinations.

**TABLE 2: IC\(_{50}\) VALUE OF EXTRACT OF DIFFERENT PARTS OF O. INDICUM AND STANDARD ANTIOXIDANTS**

<table>
<thead>
<tr>
<th>Scavenging method</th>
<th>RE</th>
<th>RBE</th>
<th>SE</th>
<th>SBE</th>
<th>LE</th>
<th>FE</th>
<th>Standard</th>
</tr>
</thead>
<tbody>
<tr>
<td>DPPH radical</td>
<td>158.23</td>
<td>112.21</td>
<td>189.36</td>
<td>149.59</td>
<td>106.4</td>
<td>159.46</td>
<td>34.6 (Rutin)</td>
</tr>
<tr>
<td>Nitric oxide radical</td>
<td>102.56</td>
<td>89.36</td>
<td>88.15</td>
<td>139.90</td>
<td>72.05</td>
<td>182.25</td>
<td>32.4 (Curcumin)</td>
</tr>
<tr>
<td>Superoxide radical</td>
<td>455.36</td>
<td>154.96</td>
<td>681.22</td>
<td>137.30</td>
<td>170.87</td>
<td>399.86</td>
<td>17.84 (Curcumin)</td>
</tr>
<tr>
<td>Hydroxy radical</td>
<td>179.35</td>
<td>47.01</td>
<td>309.27</td>
<td>33.30</td>
<td>46.52</td>
<td>76.39</td>
<td>25.85 (Catechin)</td>
</tr>
</tbody>
</table>

As per the Ayurvedic aspect and reported literature, the root and root bark possess maximum therapeutic potentiality. Since, the plant is an endangered species availability of roots as well as root bark and stem bark is difficult. As per the antioxidant aspect, leaves also can be used instead of root, root bark and stem bark.

**REFERENCES**

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