Anti-oxidative Activity of Nepalese Traditional Anti-diabetic Medicines

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Abstract
Chronic overproduction of free radicals in human body has a role in numerous disease states including diabetes. Proper use of antioxidants may help in the prevention and/or treatment of diabetes. For the management of diabetes, a combination of herbal drugs is being practiced traditionally in Nepal. These herbal medicines might have a good correlation to anti-oxidative properties. Anti-oxidative property was evaluated with the methanolic extracts of nineteen common traditional anti-diabetic medicines by 1,1-diphenyl-2-picrylhydrazyl (DPPH) radical scavenging method. Among the evaluated herbal medicines, Phyllanthus emblica, Punica granatum, Syzygium cumini, Terminalia bellirica and Terminalia chebula showed potent radical scavenging activity. The activities of P. emblica, T. bellirica and T. chebula were more potent than that of ascorbic acid. Some of the reported popular hypoglycemic herbal drugs did not show strong activity. Therefore strong anti-oxidative herbal drugs in combination with hypoglycemic drugs might have better results in the management of diabetes.

Key words: antioxidant, diabetes, DPPH, free radicals, traditional medicines

Introduction
Reactive oxygen species (ROS) such as superoxide ions (O$_2^-$), hydroxyl (HO$^-$) and nitric oxide (NO$^-$) radicals along with non-free radical species such as hydrogen peroxide (H$_2$O$_2$) and nitrous acid (HNO$_2$) are involved in normal cell regulation process (Droge 2002). Overproduction of ROS weaken the natural antioxidant system in body, first resulting in oxidative stress, and then leading to oxidative injury and finally to numerous disease states including diabetes (Robertson 2004), cardiovascular diseases (Asplund 2002), retinal ischemia, cancer, neurodegenerative disorders such as Alzheimer’s disease (Delagarza 2003), Parkinson’s disease (Packer et al. 2000), and aging processes (Beckman & Ames 1996).

The number of diabetic population is increasing due to food habit changes, aging, urbanization, increasing prevalence of obesity and physical inactivity. The prevalence of diabetes for all age groups worldwide was estimated to be 2.8% in 2000 and 4.4% in 2030. The total number of people with diabetes is projected to rise from 171 million in 2000 to 366 million in 2030 (Wild et al. 2004). There are considerable evidences that hyperglycemia results in generation of ROS, ultimately leading to increased oxidative stress in varieties of tissues. In the absence of an appropriate compensatory response from indigenous antioxidant network, the system becomes overwhelmed (redox imbalance) leading to activation of stress-sensitive intracellular signaling pathways (Packer et al. 2000). One of the major consequences is the production of gene products that cause cellular damage and are ultimately responsible for late complication of diabetes like insulin resistance, retinopathy, neuropathy and cardiovascular problems like atherosclerosis (Vincent
In animal model, Streptozotocin (STZ) induced diabetes, free radical mediated toxicity particularly to pancreatic \( \beta \) cells and nitric oxide free radicals, which played a central role (Kaneto et al. 1995, Turk et al. 1993). Antioxidants like vitamin C, E and \( \beta \)-lipoic acid have been found to be beneficial in preventing diabetes and diabetic complications (Packer et al. 2000). Various traditional medicines (Basnet et al. 1995) and plant species (Bnouham et al. 2006) have been screened for their anti-diabetic activity in animal models, however, their beneficial effect may be due to potent anti-oxidative activity of plant polyphenols or some other mechanisms remained to be explored.

Traditionally, various herbal remedies have been used for the treatment of diabetes in Nepal (Manandhar 2002). Therefore, it is important to evaluate their anti-oxidative activities and no such comprehensive study has been reported on Nepalese traditional anti-diabetic medicines. Previously, we have reported the antioxidant activity of common natural medicines (Devkota et al. 2006) and wild edible fruits (Chalise et al. 2010) from Nepal. In continuation, we have evaluated the anti-oxidative activity of Nepalese traditional anti-diabetic medicines by 1,1-diphenyl-2-picrylhydrazyl (DPPH) radical scavenging method.

**Methodology**

**Chemicals**

1,1-Diphenyl-2-picrylhydrazyl (DPPH) was purchased from Wako Pure Chemicals Co. Ltd., Osaka, Japan. HPLC grade methanol (Merck Limited, India) was used for the extraction.

<table>
<thead>
<tr>
<th>S.N.</th>
<th>Scientific Name</th>
<th>Family</th>
<th>Local Name</th>
<th>Parts Used</th>
<th>DPPH radical scavenged (%)a</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Aegle marmelos Corr.</td>
<td>Rutaceae</td>
<td>Bel</td>
<td>Fruit</td>
<td>34.4±4.8</td>
</tr>
<tr>
<td>2</td>
<td>Asparagus racemosus Wild.</td>
<td>Asparagaceae</td>
<td>Kurilo</td>
<td>Root</td>
<td>22.2±0.5</td>
</tr>
<tr>
<td>3</td>
<td>Azadirachta indica A. Juss.</td>
<td>Meliaceae</td>
<td>Neem</td>
<td>Leaf</td>
<td>10.0±3.8</td>
</tr>
<tr>
<td>4</td>
<td>Berberis aristata de Candolle</td>
<td>Berberidaceae</td>
<td>Chutro</td>
<td>Root</td>
<td>15.1±0.1</td>
</tr>
<tr>
<td>5</td>
<td>Coriandrum sativum L.</td>
<td>Umbelliferae</td>
<td>Dhaniya</td>
<td>Leaf</td>
<td>7.0±2.7</td>
</tr>
<tr>
<td>6</td>
<td>Curcuma longa L.</td>
<td>Zingiberaceae</td>
<td>Besar</td>
<td>Rhizome</td>
<td>44.6±0.5</td>
</tr>
<tr>
<td>7</td>
<td>Momordica charantia L.</td>
<td>Cucurbitaceae</td>
<td>Karela</td>
<td>Fruit peel</td>
<td>15.7±3.1</td>
</tr>
<tr>
<td>8</td>
<td>Nardostachys grandiflora DC.</td>
<td>Valerianaceae</td>
<td>Jatamansi</td>
<td>Rhizome</td>
<td>10.7±3.1</td>
</tr>
<tr>
<td>9</td>
<td>Ocimum tenuiflorum L.</td>
<td>Labiatae</td>
<td>Tulsi</td>
<td>Aerial part</td>
<td>35.2±2.8</td>
</tr>
<tr>
<td>10</td>
<td>Phyllanthus emblica L.</td>
<td>Euphorbiaceae</td>
<td>Amala</td>
<td>Fruit</td>
<td>93.3±0.2</td>
</tr>
<tr>
<td>11</td>
<td>Pierochira scrophuliflora</td>
<td>Scrophulariaceae</td>
<td>Kutki</td>
<td>Rhizome</td>
<td>16.1±3.2</td>
</tr>
<tr>
<td>12</td>
<td>Punica granatum L.</td>
<td>Punicaceae</td>
<td>Anar</td>
<td>Fruit peel</td>
<td>89.9±0.5</td>
</tr>
<tr>
<td>13</td>
<td>Swertia chirata L.</td>
<td>Gentianaceae</td>
<td>Chiraito</td>
<td>Whole plant</td>
<td>11.0±4.3</td>
</tr>
<tr>
<td>14</td>
<td>Syzygium cumini Skeels</td>
<td>Myrtaceae</td>
<td>Jarnun</td>
<td>Fruit</td>
<td>88.6±0.4</td>
</tr>
<tr>
<td>15</td>
<td>Terminalia bellirica Guertn</td>
<td>Combretaceae</td>
<td>Barro</td>
<td>Fruit</td>
<td>93.6±0.2</td>
</tr>
<tr>
<td>16</td>
<td>Terminalia chebula Retz</td>
<td>Combretaceae</td>
<td>Harro</td>
<td>Fruit</td>
<td>91.9±2.2</td>
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<tr>
<td>17</td>
<td>Tinospora sinensis Merrill</td>
<td>Menispermaceae</td>
<td>Gurjo</td>
<td>Stem</td>
<td>13.7±4.3</td>
</tr>
<tr>
<td>18</td>
<td>Trigonella foenum-graceum L.</td>
<td>Papilionaceae</td>
<td>Methi</td>
<td>Seed</td>
<td>5.2±3.2</td>
</tr>
<tr>
<td>19</td>
<td>Urtica dioica L.</td>
<td>Urticaceae</td>
<td>Sisnu</td>
<td>Young Shoots</td>
<td>34.1±4.6</td>
</tr>
<tr>
<td>20</td>
<td>Ascorbic acid</td>
<td></td>
<td></td>
<td></td>
<td>92.6±0.5</td>
</tr>
</tbody>
</table>

a Results are expressed as mean±SD (n= 3).
DPPH assay. Ascorbic acid was the product of Qualigens Fine Chemicals, India. All the chemicals used were of analytical grades.

**Collection of crude drugs**
Nineteen selected Nepalese traditional anti-diabetic herbal medicines were collected from the Pokhara valley, Nepal. They were identified by comparing literature (Manandhar, 2002) and authentic samples. Voucher specimens were preserved in the museum of materia medica, the school of pharmaceutical and biomedical sciences, Pokhara University, Nepal. The list of the natural medicines included in the present study is given as in the Table 1.

**Preparation of extracts**
Dried and ground sample of each of the natural medicine (50 g) was percolated with methanol (300 ml) at room temperature for 24 hours. After filtration, the extracts were concentrated to thick viscous mass under rotary evaporator and a portion of the extract was then dried completely in the desiccators at room temperature.

**Determination of DPPH radical scavenging activity**
DPPH radical scavenging activity was measured as described in our previous reports ((Devkota et al. 2006, Chalise et al. 2010). Briefly, 1 ml of methanolic solution of each extract at various concentrations (10, 50 and 100 µg/ml) was mixed with 1 ml of methanolic solution of DPPH (60 µm). The reaction mixture was shaken vigorously and left for 30 minutes at room temperature in dark. The anti-oxidative activity of extracts corresponding to the scavenging of DPPH radicals was measured at 520 nm with UV spectrophotometer using following formula:

\[
\text{Radical scavenging activity (\%) = 100} \times \frac{(A-B)}{A}
\]

where, A is the control absorbance of DPPH radicals without samples and B is the absorbance after reacting with samples. Ascorbic acid was used as the positive control. The result is expressed as mean of three experiments.

Those natural medicines which showed potent radical scavenging activity were again analyzed at the concentration of 1, 2, 5, 10, 15 and 20 µg/ml. From these data a curve was plotted and effective concentration (EC<sub>50</sub>) value was calculated which is defined as the concentration (µg/ml) of the extract required for 50% reduction of the DPPH radical absorbance.

**Results and Discussion**
Methanolic extracts of nineteen selected Nepalese traditional anti-diabetic medicines were screened for their anti-oxidative activity by DPPH radical scavenging method. All of these extracts showed certain activity at concentration dependant manner (Table 1). Among them, the extracts of *Phyllanthus emblica* (fruit), *Punica granatum* (fruit peel), *Syzygium cumini* (fruit), *Terminalia bellirica* (fruit) and *Terminalia chebula* (fruit) have shown more than 89% free radical scavenging activities at 10 µg/ml concentration level (Table 1). These five extracts along with ascorbic acid were further analyzed at the lower concentrations of 1, 2, 5, 10, 15 and 20 µg/ml for the calculation of the EC<sub>50</sub> values.

The EC<sub>50</sub> values for *P. emblica*, *T. chebula* and *T. bellirica* were found as 3.1, 3.3 and 3.7 µg/ml, respectively, which was lower than that of ascorbic acid (3.9 µg/ml) (Fig. 1). Phenolic compounds present in these plant species may have role in the antioxidant activity. Previous studies have reported that the traditional Ayurvedic formulation *Triphala* (equi-ratio of *P. emblica*, *T. chebula* and *T. bellirica* fruits powder) were found to inhibit lipid peroxide formation and to scavenge hydroxyl and superoxide radical *in vitro* (Sabu & Kuttan 2002).

![Fig. 1. EC<sub>50</sub> (g/ml) value of the potent traditional medicines](image)

**Azardichta indica, Momordica charantia** and *Trigonella foenum-graceum* are very common and considered as effective traditional anti-diabetic drug

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in Nepal. The hypoglycemic activities of the individual drug are also widely studied. However, the antioxidant properties of these drugs were found to be very low (Table 1). In the meantime, it is very common practice that diabetic people in Nepal take A. indica, M. charantia and/or T. foenum-graceum together with Triphala. The herbal practitioner and patients often claim on the improvement of glycemic condition by taking combined prescription, however we did not find any systematic study so far.

This study suggested that radical scavenging activity may be one of the pathways to show further beneficial effects of these traditional medicines in the treatment of diabetes in combination with modern drugs. Further studies on the activity guided isolation of the potent antioxidant constituents from these medicines and evaluation of other mechanism for anti-diabetic activity may help in better understanding of diabetes treatment.

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References