Lipid Lowering Activity of *Anthocephalus indicus* Root in Hyperlipidemic Rats

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The lipid lowering activity of *Anthocephalus indicus* (family Rubiaceae; Hindi name Kadamba) root extract has been studied in triton WR-1339 induced hyperlipidemia in rats. In this model, feeding with root extract (500 mg kg⁻¹ b.w.) lowered plasma lipids and reactivated post-heparin lipolytic activity in hyperlipidemic rats. Furthermore, the root extract (50–500 μM) inhibited the generation of superoxide anions and hydroxyl radicals, in both enzymic and non-enzymic systems, *in vitro*. The results of the present study demonstrated both lipid lowering and antioxidant activities in root extract of *A. indicus*, which could help prevention of hyperlipidemia and related diseases.

**Keywords:** *Anthocephalus indicus* – hypolipidemic agent – oxygen free radical scavengers – plant antioxidant – Rubiaceae – triton model of hyperlipidemia

Introduction

Disorders of lipid metabolism, hyperlipidemia, hypertension and obesity are associated with increased oxidative stress and overproduction of oxygen free radicals (1). An excess of superoxide anions (O₂⁻) are further converted into other reactive oxygen species and among them hydroxyl radical (OH⁻) is more damaging to lipids and lipoproteins (2). Moreover, hyperlipidemia following oxidative stress may cause oxidative modifications in low-density lipoproteins, which play an important role in the initiation and progression of atherosclerosis and related cardiovascular diseases (3). Furthermore, there have been reports that the lipid lowering drugs: fibrates, statins and bile acid sequestrants used for the treatment of hyperlipidemia associated disorders do not possess antioxidant property and they are also not free from toxic side effects (4). Therefore, on the basis of above stated facts there is an urgent need to have a drug having the dual property of lowering lipid level and antioxidant activities together and for this natural products are the best option.

World ethnobotanical information reported that a number of herbal medicines from plants and vegetables are used for controlling hyperlipidemia and related complications in patients (5). *Anthocephalus indicus*: the *cadamba Miq* (family Rubiaceae, Hindi name; Kadamba) is one such Ayurvedic remedy that has been mentioned in many Indian medical literatures for the treatment of fever, anemia, uterine complaints, menorrhagia, blood and skin diseases, diarrhea, colitis, stomatitis, dysentery and in improvement of semen quality (6,7). *Anthocephalus indicus* grows through out India, especially at low levels in wet places. In traditional system of remedies warm aqueous extract of *A. indicus* leaves have been used to alleviate the pain, swelling and for cleansing and better wound healing. Recently, *A. indicus* has been reported to posses...
antimicrobial, wound healing, antioxidant, antimalarial and hepatoprotective activity (8–10). The fruit juice of the plant augments the quantity of breast milk of lactating mothers and also works as a lactodepurant (11). Root extract of this plant is salutary in urinary ailments like dysurea, calculi and glycosuria (11). There are reports that heartwood, leaves, flower and seeds of A. indicus contain typical alkaloid; Cadambine and its derivatives, some complex polysaccharides and other common constituents (12). However, pharmacological effects of these compounds are not well known. Also much work has not so far been done to explore other biological activities and/or the isolation of active ingredients from the root. Therefore, the present study was designed to investigate the lipid lowering activity of A. indicus root in hyperlipidemic rats. Furthermore, to explain the positive role in protection against oxidative degradation of body lipids, the antioxidant potential of natural product was also assessed in vitro.

Methods

Preparation of Root Extract

Anthocephalus indica roots were collected from Lucknow and identified taxonomically by the Department of Pharmacology, Era’s Lucknow Medical College, Lucknow. A voucher specimen was also submitted. Roots were dried under shade and made into fine powder using laboratory mill. Powder (100 g) was extracted thrice with 500 ml portions of 95% ethyl alcohol in a laboratory perculator at room temperature. Time allowed for each extraction was 4 h. The root extract obtained after third extraction was colorless. All the extracts were mixed with 500 ml portions of 95% ethyl alcohol in a laboratory perculator. Time allowed for each extraction was 4 h. The root extract obtained after third extraction was colorless. All the extracts were mixed with 500 ml portions of 95% ethyl alcohol in a laboratory perculator at room temperature. Time allowed for each extraction was 4 h. The root extract obtained after third extraction was colorless. All the extracts were mixed

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Lipid Lowering Activity in Triton Induced Hyperlipidemia

Animal study was performed with the approval of Animal Care Committee of Division of Laboratory Animals; Central Drug Research Institute, Lucknow, India and confirmed to the guide for the Care and Use of Laboratory Animals (CDRI, Lucknow). Male adult rats of Charles Foster strain (200–225 g) bred in the animal house of the Institute were used. The animals were divided into four groups. A group of six animals in a cage were kept in controlled conditions, temperature 25–26°C, relative humidity 60–80% and 12/12 h light/dark cycle (light

Table 1. Ingredient and nutrient composition of the control basal diet fed to rats

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Amount (g kg⁻¹)</th>
</tr>
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<tbody>
<tr>
<td>Casein</td>
<td>21</td>
</tr>
<tr>
<td>Corn starch</td>
<td>440</td>
</tr>
<tr>
<td>Sucrose</td>
<td>100</td>
</tr>
<tr>
<td>Maltose dextrin</td>
<td>100</td>
</tr>
<tr>
<td>Cellulose</td>
<td>50</td>
</tr>
<tr>
<td>Soybean oil</td>
<td>50</td>
</tr>
<tr>
<td>Vitamins mix</td>
<td>10</td>
</tr>
<tr>
<td>Mineral</td>
<td>35</td>
</tr>
</tbody>
</table>

Other ingredients include choline bitartrate (2 g kg⁻¹) and L-ascorbic acid (50 mg kg⁻¹). Proximate analysis 21% crude protein, 5% crude fat, 4% crude fiber, 8% ash.

from 08:00 a.m. to 08:00 p.m.) and provided with standard pellet diet (Lipton India, Ltd) and water ad libitum. Details about dietary ingredients and nutrients of pellet diet are given in Table 1. Hyperlipidemia in rats was induced by a single injection of triton WR-1339 (400 mg kg⁻¹, b.w.) and injected 1 ml kg⁻¹ b.w. to each rat intraperitoneally (13). The root extract as well as gemfibrozil were used in the experiment.

Blood Collection and Biochemical Analysis in Plasma

At the end of experiment, rats were fasted for 18 h and then injected with heparin solution prepared in normal saline (10 mg ml⁻¹) at the dose of 1 ml kg⁻¹, b.w. through tail vain. After 15 min of treatment the animals were anaesthetized with sodium pentothal solution (50 mg kg⁻¹, i.p.) prepared in normal saline. Blood was withdrawn from retro-orbital sinus using glass capillary in EDTA coated tubes (3 mg ml⁻¹ blood). The blood was centrifuged and the post-heparin plasma was separated. Plasma was diluted with normal saline in the ratio 1:3 v/v and used for the analysis of lipids and proteins.

Total cholesterol (TC), phospholipids (PL), triglyceride (TG), protein and post-heparin lipolytic activity (PHLA) were determined according to the methods reported earlier (16).
Estimation of Antioxidant Activity of *A. indicus* Root Extract

**Generation of Superoxide Anions**

Superoxide anions (O$_2^-$) were formed in an enzymic system composed of xanthine, xanthine oxidase in the absence or presence of *A. indicus* root extract at different concentrations of 50, 100, 250 and 500 μM. The amount of uric acid formed was assayed spectrophotometrically at 295 nm (17). Another reaction mixture contained xanthine, xanthine oxidase and nitroblue tetrazolium (NBT) with or without addition of root extract (50–500 μM). After incubation the amount of formazone formed, as a result of reduction of NBT by O$_2^-$, was measured at 560 nm on spectrophotometer (17). The system employed for non-enzymic generation of O$_2^-$ comprised of phenazine methosulphate, NADH and NBT in the absence or presence of root extract (50–500 μM). After reaction in the absence or presence of root extract (50–500 μM), the reaction mixture was assayed for malondialdehyde formed (20).

**Generation of Hydroxyl Radicals**

Root extract of *A. indicus* (50–500 μM) was tested against the formation of hydroxyl radicals (OH$^-$) in an enzymic system composed of hypoxanthine, FeSO$_4$.7H$_2$O (Fe$^{+2}$), sodium ascorbate and xanthine oxidase, assayed for 3,4-dihydroxybenzoate formed by OH$^-$ mediated hydroxylation of salicylate (19). In another set of experiment OH$^-$ were generated non-enzymically by Fe$^{+2}$, sodium ascorbate, H$_2$O$_2$ and deoxyribose. After reaction in the absence or presence of root extract (50–500 μM), the reaction mixture was assayed for malondialdehyde formed (20).

**Statistical Analysis**

One way analysis of variance (ANOVA-New man’s student $t$-test) were performed by comparison of values for triton treated group with control, triton and drug treated groups with triton (21). All hypothesis testing were two-tailed. $P < 0.05$ was considered statistically significant and the results were expressed as mean ± SD. The statistical analysis was carried out by the Graph Pad INSTAT 3.0 software. Similarly, the generations of oxygen free radicals with different concentrations of *A. indicus* root extract were compared with that of their formation without extract. The values were tested for significance at a $P < 0.05$.

**Results**

**Effect of *A. indicus* Root on Triton Induced Hyperlipidemia**

The acute administration of triton WR-1339 caused marked increase ($P < 0.001$) in plasma levels of TC (260%), PL (103%), Tg (124%) and protein (81%) following inhibition of PHLA by 24%. Treatment with *A. indicus* root extract caused significant reversal ($P < 0.01$) in these levels of TC, PL, Tg and protein by 22–23% following reactivation of PHLA by 40% (Figs 1, 2, 3). The lipid lowering activity of gemfibrozil (29–37%) was comparatively more than that of *A. indicus*.

**Effect of *A. indicus* on Generation of O$_2^-$ Anions**

Figure 4 shows that enzymic oxidation of xanthine to uric acid by O$_2^-$ (i) as well as the generation of O$_2^-$ anions in xanthine-xanthine oxidase systems, as measured
by reduction of NBT to formazone (ii) was inhibited to varying extents by root extract (50–500 μM) in a concentration dependent manner (P < 0.05–0.001) and this effect was maximum by 47% and 43%, respectively, at 500 μM of extract. The root extract also trapped the O$_2^-$ anions generated by non-enzymic system of NADH-phenazine methosulphate as measured by reduction of NBT in the reaction mixture. The effect was dose dependent (P < 0.05–0.001) and 59% inhibition was observed at 500 μM of test substance.

**Effect of A. indicus on Generation of OH$^-$/C0 Radicals**

The data in Fig. 5 shows that A. indicus root extract inhibited the formation of OH$^-$ by enzymic system of hypoxanthine-xanthine oxidase and Fe$^{+2}$. Addition of extract (50–500 μM) inhibited the OH$^-$ mediated formation of 2,3-dihydroxybenzoate in concentration dependent manner, the significant effects were observed with (100–500 μM) showing 38% inhibition (P < 0.001) at 500 μM of extract. Furthermore, this preparation, when added with the reaction mixture containing Fe$^{+2}$-Na ascorbate-H$_2$O$_2$ employed for non-enzymic generation of OH$^-$, inhibited OH$^-$ mediated fragmentation of deoxyribose (P < 0.05–0.001) into MDA and this was maximum by 44% at 500 μM concentration of root extract.

**Discussion**

The present study was planned, as there have been reports that various hypolipidemic drugs available in market, like statins, fibrates and bile acid sequestrants have numerous side effects. It has been reported that statins, the inhibitors of cholesterol biosynthesis, regress hypercholesterolemia and also possess antioxidant properties. In our earlier studies we also observed (our unpublished data) that lovastatin at the dose of 20 mg kg$^{-1}$ lowered the blood level of TC, PL and Tg, by 39, 32 and 31%, respectively, in Triton-rat model. Furthermore, we also observed that lovastatin possess potent antioxidant activity as at test concentrations of 5, 10 and 20 μM, it inhibited the generation of OH$^-$ in vitro by 62, 80 and 98%, respectively. However, all the statins,
including lovastatin, have many adverse effects. Comp-actin and mevinolin cause nausea, skin rashes, diarrhea, rise in CPK and transaminases and increases risk of development of premature nuclear cataract. Clinical studies have contraindicated the use of lovastatin in case of pregnant and lactating women (22). Similarly, the bile acid sequestrants, cholestyramine and cholestipol, are reported to cause gastrointestinal problems (23), and 3β-isomers of dihydrocadambine and isodihydrocadambine (28,29). It has been reported that its stem bark contains cadambagic acid along with quinovic acid and β-sitosterol (30). Moreover, a complex polysaccha-
ride from flowers and seeds of A. indicus has been isolated (31). The above-mentioned compounds of A. indicus may be responsible for exerting beneficial effects. However, not much work has so far been done to investigate the biological activities of these compounds. Chlorogenic acid isolated from A. indicus (Cadamba) has been reported to be a potent hepatoprotective agent (10). There are reports that the plant metabolites also possess antimalarial activity (9). Recently, alcoholic and aqueous extracts of A. indicus are reported to have significant antimicrobial, wound healing and antioxidant activities (8). It is suggested that the antimicrobial activity may be due to the presence of terpenes (32). It is suggested that A. indicus root may have typical saponins, terpenes, glycosides and alkaloids which may be responsible for lipid lowering and antioxidant activities, as observed by us, however, this needs further confirmation. In conclusion, it may be stated that the results of the present study demonstrated new properties of A. indicus root as a potent lipid lowering and antioxidant agent and these beneficial activities may contribute to its cardio protective and antiatherosclerotic role. Moreover, further studies on drug metabolism and to assess the biological activity, in vivo and in vitro, of A. indicus root and its various fractions are under progress to substantiate the present findings.

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Received July 10, 2007; accepted December 17, 2007