Evaluation of anti-cholesteremic and anti-lipidemic activity of seed extract of *Achyranthes aspera* in diet induced hyperlipidemia model in rats

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**ABSTRACT**

The present study was carried out to evaluate the anti-cholesteremic and Anti-lipidemic activity of methanolic seed extract of Achyranthes aspera using Diet-induced hyperlipidemia model in Wistar albino rats. Albino Wistar rats of either sex weighing in between 200-350 gms were divided into 6 groups containing 6 animals in each group. All the groups received cholesterol (400mg/kg b.w. in 5ml coconut oil p.o) except saline control for 2 weeks and were tested orally at a dose of 150, 300 and 600 mg/kg body weight. The extract (300 and 600 mg/kg) showed significant (P<0.05) reduction in total serum cholesterol and lipid levels compared to vehicle control group. The serum Total cholesterol, Triglyceride, HDL, VLDL and LDL levels were analyzed. This present study proved that methanolic seed extract of Achyranthes aspera shows significant Anticholesteremic and Antilipidemic activity against the Diet induced hyperlipidemia model by reducing the serum TC, TG, VLDL, LDL levels and increasing HDL level.

**Keywords:** Achyranthes aspera, Antilipidemic, Anticholesteremic TC, TG, HDL, VLDL, LDL.

**INTRODUCTION**

Hyperlipidemia has been ranked as one of the greatest risk factors contributing to the prevalence and severity of coronary heart diseases. Coronary heart disease, stroke, atherosclerosis and hyperlipidemia are the primary cause of death. Hyperlipidemia is the presence of high levels of cholesterol in the blood. It is not a disease but a metabolic derangement that can be secondary to many diseases and can contribute to many forms of disease, most notably cardiovascular disease. The treatment of hyperlipidemia depends on the patient’s cholesterol profile. Many antihyperlipidemic agents like statin, fibrates, niacin, bile acids, ezitimibe etc reduce cholesterol level with different condition. Hyperlipidemia characterized by elevated serum total cholesterol, low density, very low density lipoprotein and decrease high density lipoprotein are the risk factor for coronary heart diseases. Hyperlipidemia associated lipid disorders are considered to cause the atherosclerotic cardiovascular disease. Hyperlipidemia is classified into a primary and a secondary type, which indicates the complexities associated with disease. for that reason selected the *Achyranthes aspera* (family: Amaranthaceae) is an important medicinal herb found as a weed throughout India. Though almost all of its parts are used in traditional systems of medicines, seeds, roots and shoots are the most important parts which are used medicinally. Traditionally which possesses activities like antiperiodic, diuretic, purgative, laxative, antiasthmatic, hepatoprotective, anti-allergic and various other important medicinal properties. It is seen from the literature that Achyranthes aspera is a very important plant for its large number of medicinal properties as well as medicinally important chemicals like ecadytone, achyranthine, betaine, pentatrairotane, 6-pentatrairotanone, hexatrairotane and tritriairotane. The plant shows many pharmacological activities like spermicidal, anti-allergic, cardiovascular, nephroprotective, antiparassitic, hypoglycemic, analgesic and antipyretic. Many traditional uses are also reported like antiperiodic, purgative and laxative, in various types of
gastric disorders and in body pain which are being studied till today and further research has to be done4,5,6. Thus, Achyranthes aspera is quite promising as a multipurpose medicinal agent so further clinical trials should be performed to prove its efficacy. The crushed plant is used in pneumonia and infusion of the root is used as mild astringent in bowel complaints. Decoction of powdered leaves with honey or sugar candy is useful in early stages of diarrhoea and dysentery. For the last few decades or so, extensive research work has been done to prove its biological activities and pharmacology of its extracts. Saponins, oleonolic acid, dihydroxy ketones, alkaloids, long chain compounds and many other chemical constituents have been isolated. There are some reports on Anticholesteremic and Antilipidemic activity of this plant. Achyranthes aspera Lann. plant is claimed to possess cholesterol-reducing effect and is used to treat patients with heart disease and obesity7.

**EXPERIMENTAL SECTION**

**Collection of plant materials**

The seed of Achyranthes aspera were collected in June 2013 from different localities of Warangal district, Andhrapradesh state. The plant materials authenticated by botanist Dr. V. Raju, Professor & head of Department of botany, Kakatiya University, Warangal. The seeds were washed with distilled water and dried in shade, pulverized by mechanical grinder to get course powder and stored in an airtight container.

**Preparation of Plants Extracts**

The powdered material obtained was then subjected to successive extraction by Hot Percolation Method using methanol solvent in a soxhelet extractor. The extract obtained were evaporated at 45°C to get a semisolid mass. The percentage yield of Methanolic extract was found to be 45.50% w/w and the methanolic extract was used for further studies8.

**Acute toxicity test LD50 (9-12)**

Adult male and female wistar albino rats (6 – 8 weeks old) were weighed between 150 – 180 gm. The animals were given standard rat pellets and tap water and *libitum*. The acute toxic study was used to determine a safe dose for the stem bark extract. Eighteen rats (9 males and 9 females) were assigned equally each into 3 groups labelled as vehicle (CMC, 0.25% w/v, 5 ml/kg); 2000 and 5000 mg/kg of Achyranthes aspera seeds extract preparation, respectively. The animals were fasted overnight (water but not food) prior to dosing. Food was withheld for a further 3 to 4 h after dosing. The animals were observed for 30 min and 2, 4, 8, 24 and 48 h after the administration for the onset of clinical or toxicological symptoms. Mortality, if any was observed over a period of 2 weeks. The acute toxicity LD50 was calculated as the geometric mean of the dose that resulted in 100% lethality and that which caused no lethality at all.

**Experimental Animals**

Wistar albino rats of either sex weighing around 200-350gm were taken from inbreeds colony animals, which were housed in polypropylene cages under standard laboratory conditions (light period 7.00 A.M. to 7.00 P.M., 21±2 °C, and relative humidity 55%). The animals were given standard rat pellets and tap water *ad libitum*, but they were deprived of food 36 h before the experiments. The rats were acclimatized to laboratory condition for 7 days before commencement of experiment. All procedures involving laboratory animal use were in accordance to the Institute Animal Ethics Committee regulations approved by the Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA).

**Animal treatment(13-15)**

A high fat / cholesterol supplement to normal diet was administered separately incorporated into diet pellets for 1-2 weeks (like 400 mg/kg cholesterol in 5ml coconut oil). The high fat diet was prepared by mixing calculated amounts of 2% cholesterol, 1% cholic acid and 1 mL coconut oil. The Parachute coconut oil was chosen because of its high saturated fat content which aggravates the atherogenic profile in the rats (Hassarajani et al., 2007) with slight modification. Hyperlipidemic control group was orally administered with vehicle (carboxymethyl cellulose, CMC, 0.25% w/v, 5ml/kg). The reference group received oral doses of 4 mg/kg simvastatin in CMC (5 ml/kg) as positive controls. Experimental groups were orally administered with 150, 300 and 600 mg/kg of methanolic seeds extract of Achyranthes aspera in distilled water (5 ml/kg), respectively. The biochemical parameters (HDL, LDL, VLDL, TC and TG) have been investigated in serum after 2 weeks during the treatment with Simvastatin and alcoholic extract of the plant. Blood was withdrawn using the heparanised capillaries from the retro-orbital sinus in the overnight fasted animals. The serum was obtained after centrifuging the blood, which was used to estimate the concentration of biochemical parameters using the semi auto analyser and relevant lipid profile kits. Total cholesterol, triglyceride level, and were estimated from serum by CHOD-PAP method and GPO-PAP method and HDL by the precipitation method using phosphotungstate magnesium acetate reagent. LDL and VLDL-cholesterol were calculated following
the method by Johnson et al. (1997). Thus obtained data’s were tabulated and checked for the statistical tools like one way analysis of variance (ANOVA). Finally Dunnet’s test was applied to find out difference between groups.

**Statistical analysis**

The values are represented as mean ± S.E.M, and statistical significance between treated and control groups was analyzed using of One way ANOVA, followed by Dunnett’s test where P<0.05 was considered statistically significant.

**RESULTS**

**Acute toxicity study**

An acute toxicity study was carried out in which the animals were treated with the seeds extracts at a dose of 2000 and 5000 mg/kg of *Achyranthes aspera* and were kept under observation for 14 days. All the animals remain alive and did not manifest any significant visible signs of toxicity at these doses. There were no abnormal signs, behavioural changes, body weight changes, or macroscopic finding at any time during the observation period.

**Diet induced hyperlipidemia**

There was a significant increase in the serum cholesterol, triglyceride, LDL, VLDL and decrease in the levels of HDL in the high fat diet fed animals when compared to normal fed rats. Treatment with methanolic extract, at three different doses, decreased the serum level of cholesterol, triglyceride, LDL and VLDL and increased the serum HDL levels as compared to hyperlipidemic control group. (Table-1),

**DISCUSSION**

The present study was undertaken to evaluate the Anticholesteremic and Antilipidemic activity of methanolic extract of seeds of *Achyranthes aspera*. The study was conducted by using Diet induced hyperlipidemia model. The parameters used for the assessment of Anticholesteremic and Antilipidemic activity are Total Cholesterol, Triglycerides, High density Lipoproteins, Very Low Density Lipoproteins, and Low Density Lipoproteins. Flavonoids may augment the activity of lecithin acyl transferase (LCAT), which regulates blood lipids. LCAT plays a key role in the incorporation of free cholesterol into HDL (this may increase HDL) and transferring it back to VLDL and LDL which are taken back later in liver cells. Saluja et al. (1978) reported isolating β-sitosterol. from the hybrid variety of *Achyranthes aspera* is a plant sterol with a structure similar to that of cholesterol, except for the substitution of an ethyl group at C24 of its side chain [16]. It is believed to lower cholesterol by lowering plasma concentrations of LDL (Kane and Malloy, 1982)(17) Plant sterol reduces the absorption of cholesterol and thus increases the fecal excretion of steroids that results in decrease of body lipids. We can suggest that it may be possible to use plant seeds extracts as remedy to prevent hyperlipidemia. However, further investigations are required to elucidate their exact mechanism of action of antihyperlipidemic activity.

**Table 1: Effect of *Achyranthes aspera* seed extract on various parameters in Diet induced Hyperlipidemia model**

<table>
<thead>
<tr>
<th>Treatment Groups</th>
<th>Cholesterol (mg/dl)</th>
<th>Triglyceride (mg/dl)</th>
<th>HDL (mg/dl)</th>
<th>VLDL (mg/dl)</th>
<th>LDL (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>82.17±3.78</td>
<td>125.5±4.11</td>
<td>44.50±2.44</td>
<td>25.10±0.82</td>
<td>12.50±2.12</td>
</tr>
<tr>
<td>Control(HFD)</td>
<td>190.3±4.24</td>
<td>197.8±6.22</td>
<td>39.17±3.0</td>
<td>39.57±1.24</td>
<td>111.6±5.08</td>
</tr>
<tr>
<td>Methanolic Extract 150mg/kg</td>
<td>183.8±3.87</td>
<td>188.8±3.27</td>
<td>41.67±1.33</td>
<td>37.77±0.65</td>
<td>104.5±4.44</td>
</tr>
<tr>
<td>Methanolic Extract 300mg/kg</td>
<td>143.2±4.16**</td>
<td>165.8±4.16**</td>
<td>47.83±2.12**</td>
<td>33.17±0.83*</td>
<td>62.17±2.82*</td>
</tr>
<tr>
<td>Methanolic Extract 600mg/kg</td>
<td>113.2±4.51***</td>
<td>140.8±4.83 **</td>
<td>51.67±2.55***</td>
<td>28.17±1.16***</td>
<td>33.33±6.58**</td>
</tr>
<tr>
<td>Simvastatin (standard)</td>
<td>96.50±3.24***</td>
<td>128.0±3.14***</td>
<td>57.67±3.14***</td>
<td>25.60±2.62***</td>
<td>13.23±4.23***</td>
</tr>
</tbody>
</table>

Values are expressed as (mean±SEM) P***<0.001, P**<0.01, P*0.05 (When compared with hyperlipidemic control) P<0.001 (When compared with control). Values are expressed in mean±SEM

**CONCLUSION**

The present study was under taken to assess the Anticholesteremic and Antilipidemic activity of methanolic extracts of seeds of *Achyranthes aspera*. The methanolic extracts of seeds of *Achyranthes aspera* showed marked decrease in the serum Total Cholesterol, Triglyceride, VLDL, LDL levels and increase in HDL level in diet induced hyperlipidemia model in a dose dependent manner. Methanolic extracts of seeds of *Achyranthes aspera* at 600 mg/kg reduced the lipids and cholesterol significantly, thus showing the Anticholesteremic and Antilipidemic mechanism involved. So, methanolic extract at 600 mg/kg exhibited almost equipotent effect as that of simvastatin.
and these results offer pharmacological evidence and support on the folkloric use of Achyranthes aspera seeds as an anticholesteremic and Antilipidemic agent. (Table-1), It leads to the conclusion that the seeds of Achyranthes aspera can be utilized for its Anticholesteremic and Antilipidemic activity and in future, this work has been extended by including more hyperlipidemic models for meaningful and tangible conclusion. Toxicological studies can also be carried out to know about the toxic and non-toxic nature of the drug.

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REFERENCES

[8] A. Banerji et al. (1971) isolated ecdysterone from the whole plant [32]. K.S.Laddha (2005) et al. reported extraction, isolation and purification of 20-hydroxyecdysone from Achyranthes aspera and its characterization by DSC, UV, IR, CD, 1H and 13C NMR, MS and quantification by HPLC.