Antiobesity, hypolipidemic, antioxidant and hepatoprotective effects of *Achyranthes aspera* seed saponins in high cholesterol fed albino rats

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Abstract

**Introduction:** Numerous herbal medicines have been recommended for the treatment of different diseases. *Achyranthes aspera*, Linn. (Family: Amaranthaceae), popularly known as Charchitta or Pitpapra, is commonly used by traditional healers for the treatment of fever, malaria, dysentery, asthma, arterial hypertension, pneumonia, and diabetes. The root extract is well reputed for its insect molting hormonal activity. This investigation was conducted to evaluate the effects of saponins from *Achyranthes aspera* seeds on the serum lipid profile of albino rats fed a high cholesterol diet.

**Material and methods:** Hypolipidemic, antioxidant and hepatoprotective activities of these saponins were tested as described previously. To determine the mechanism underlying the observed effects, serum antioxidant status was assessed according to ABTS (2,2'-azino-bis-3-ethylbenzo-thiazoline-6-sulfonic acid), superoxide dismutase and ferric ion reducing antioxidant power (FRAP) assays in saponin-treated hyperlipidemic animals. Liver enzyme levels were determined to reveal any possible hepatotoxicity.

**Results:** Four-week oral administration of *A. aspera* seed saponins produced a significant (*p* < 0.05) decrease of total cholesterol, total triglycerides and LDL-C and a significant increase of HDL-C level in hyperlipidemic rats. Treatment with *A. aspera* seed saponins also showed a significant (*p* < 0.01) improvement of serum antioxidant status in tested animals. No significant hepatotoxicity was produced by such treatment as the serum liver enzyme activity remained unaltered.

**Conclusions:** Saponins from *A. aspera* seeds possess antihyperlipidemic and antioxidant properties which might lead to improvement of serum lipid profile and blood antioxidant status. Our findings support the folkloric use of this indigenous plant in the treatment of hyperlipidemia. However, its exact mechanism of action remains to be elucidated.

**Key words:** hyperlipidemia, antioxidant, hepatoprotection, *Achyranthes aspera*, ABTS, high cholesterol diet.
Introduction

Hyperlipidemia has been defined as abnormally elevated lipids or lipoproteins in the blood. Hyperlipidemia is nowadays a major health problem faced by many societies, which concerns health professionals, since it constitutes one of the major risk factors for the development of cardiovascular disease [1]. It has been reported that atherosclerosis and coronary heart disease are mainly caused by hyperlipidemia [2, 3]. Over the past decades atherosclerosis-related diseases diminished but they still account for significant morbidity and mortality in many middle aged and elderly people [4]. In respect of hyperlipidemia, recent interest has been focused on strategies that could enhance the removal of reactive oxygen species (ROS), by using either natural antioxidants or drugs that have the capacity to enhance the endogenous antioxidant system [5]. The antioxidant defense system is driven by different antioxidant mechanisms, for which plants have different groups of compounds, namely preventive antioxidants (enzymes such as catalases or peroxidases), radical scavenging compounds (ascorbic acid or carotenoids), and enzymes repairing DNA material. In the group of antioxidant and free radical scavenging agents, plants synthesize different compounds, principally phenolic derivatives, such as flavonoids, phenylpropanoids, stilbenes and many others. These compounds are also potential antioxidant agents for humans, protecting us from the damage caused by reactive oxygen species (ROS) [6]. Due to the growing interest in free radical technology, the importance of naturally occurring antioxidants cannot be ignored, as the safety of synthetic antioxidants such as butylated hydroxytoluene (BHT), butylated hydroxyanisole (BHA), tertiary butylhydroquinone (TBHQ), gallic acid esters, etc. remains questionable. Moreover, these antioxidants have poor solubility and usually only show moderate antioxidant activity [7, 8].

Some preliminary studies have shown that A. aspera seed contains saponins (glycosides of oleanonic acid) A and B, which might exert an antiobesity effect in albino mice by reducing the excess accumulation of body fat and changing the serum lipid profile [9, 10].

In the light of previous research works the present investigation was carried out to more precisely evaluate the possible hypolipidemic and antiobesity activity of indigenous plant A. aspera seed saponins that have been used for centuries to treat various diseases. It is a very common weed of waste places and road sides in Pakistan.

Material and methods

Plant material

The ripe seeds of A. aspera locally known as Puthkanda were collected from various gardens and grazing areas of Islamabad between September and October, 2009. Seeds were authenticated and identified by the Department of Biological Sciences, University of Sargodha, Sargodha and the specimen was deposited in the departmental herbarium. The seeds were completely dried in the shade and were finally powdered with a Chinese herbal grinder. The powdered seeds were stored in closed cellophane bags at 4°C in a refrigerator until further analysis.

Experimental animals

Albino rats of both sexes, aged 3–4 months and weighing around 250–300 g, were used. The animals were kept in an animal room with an alternate light and dark cycle of 12 h (temperature 25 ±2°C). Standard diet and fresh water was supplied ad libitum. The study was conducted in accordance with Good Laboratory Practices (GLP) regulations of the WHO. The principles of laboratory animal care of the National Institute of Health (NIH) were followed in the study. The study protocol was approved by the institutional ethic committee.

Forty-eight rats were randomly divided into eight groups of six animals each (A–H: normal control, negative control, five groups treated with saponins and another one treated with atorvastatin – positive control) and given free access to a high cholesterol diet (HCD) ad libitum. The food and water intake by each group was monitored at least twice weekly. The HCD was made from preformed cholesterol powder, butter, soybean oil and liver extract. A description of each group is shown in Table I. Body weight (g) and nasoanal length (cm²) were measured at the start of the experiment, after 14 and 28 days, both in normal and HCD treated animals. Body mass index (BMI) was calculated by dividing the body weight (in g) by square of nasoanal length (in cm²).

Detection, extraction and isolation of saponins

Ten grams of powdered A. aspera seeds were extracted with 50 ml of hot water and filtered. Five ml of the above filtrate was added to a test tube (Pyrex, Germany) containing 5 ml of 1.8% NaCl solution; 5 ml of distilled water was added to another tube with 5 ml of 1.8% NaCl solution (blank tube). Next, 8 drops of blood were added to both tubes – hemolysis occurred in the test tube containing the extract, indicating the presence of saponins, while no hemolysis was seen in the blank tube.

Fifty grams of powdered A. aspera seeds were incubated with 1000 ml of 2 N hydrochloric acid for 2 h and filtered. Residues were neutralized by passing 4% ammonium hydroxide solution...
Table I. Effect of A. aspera seed saponins on body mass index and body weight (means ± SEM) in rats fed with HCD along with 2% aqueous gum tragacanth solution after 28 days

| Group | Description                                                                 | Weight [g]       | Body mass index |
|-------|-----------------------------------------------------------------------------|------------------|-----------------|
|       |                                                                             | Day 0 | Day 14 | Day 28 | Day 0 | Day 14 | Day 28 |
| A     | Untreated control fed orally normal diet with 2% gum tragacanth solution (0.5 ml/kg body weight) | 276.0 ±11.98 | 278.33 ±14.90 | 278 ±5.41 | 0.72 | 0.72 | 0.72 |
| B     | Negative control fed orally high cholesterol diet (HCD)                     | 276.16 ±6.70 | 302.83 ±5.63* | 321 ±5.46* | 0.73 | 0.80* | 0.83* |
| C     | Receiving orally 50 mg/kg body weight of A. aspera seed saponins suspended in 2% gum tragacanth aqueous solution along with HCD | 274.16 ±9.19 | 293.33 ±4.10* | 290.83 ±4.40** | 0.73 | 0.75* | 0.75** |
| D     | Receiving orally 75 mg/kg body weight of A. aspera seed saponins suspended in 2% gum tragacanth aqueous solution along with HCD | 268.83 ±8.29 | 282.16 ±4.48** | 287.17 ±7.15** | 0.71 | 0.74** | 0.74** |
| E     | Receiving orally 100 mg/kg body weight of A. aspera seed saponins suspended in 2% gum tragacanth aqueous solution along with HCD | 273.83 ±4.52 | 284.16 ±5.33** | 286 ±8.48** | 0.72 | 0.74** | 0.74** |
| F     | Receiving orally 125 mg/kg body weight of A. aspera seed saponins suspended in 2% gum tragacanth aqueous solution along with HCD | 275.16 ±8.6 | 279.83 ±10.63** | 276.17 ±6.2** | 0.73 | 0.71** | 0.71** |
| G     | Receiving orally 150 mg/kg body weight of A. aspera seed saponins suspended in 2% gum tragacanth aqueous solution along with HCD | 274.83 ±11.9 | 278.33 ±3.14** | 274 ±4.86** | 0.73 | 0.70** | 0.70** |
| H     | Treated orally with 8 mg/kg body weight of atorvastatin along with HCD      | 277.33 ±6.0 | 277.66 ±5.50** | 278 ±4.2** | 0.72 | 0.71** | 0.72** |

Data represented as mean ± SEM (n = 6). *p < 0.01 when compared with group A animals (normal control) receiving only normal diet with vehicle (2% gum tragacanth solution). **p < 0.05 when compared with group B animals (negative control) receiving only high cholesterol diet with vehicle (2% gum tragacanth solution).
through the filtrate, dried and filtered with a paper at 60°C for 16 h. The dry residues were next packed into a Soxhlet thimble and extracted with petroleum ether (Sigma Aldrich Chemicals, USA) at 40–60°C for 2 h. The solvent was reduced to 30 ml and placed in a cool place. Finally, saponins were precipitated with acetone (Sigma Aldrich Chemicals, USA). The exact identification of saponins was described elsewhere [11].

Preparation of plant drug suspensions and induction of hyperlipidemia

The saponins were triturated with a 2% aqueous gum tragacanth solution and the final volume was made up to 20 ml to obtain a suitable concentration of saponins per ml for oral administration. Then the plant drug suspension was administered orally to each animal for investigation of the antihyperlipidemic and antioxidant effects. As a control atorvastatin was also administered after suspension in 2% aqueous gum tragacanth solution. Hyperlipidemia was induced by oral administration of a high cholesterol diet with preformed cholesterol powder, butter and liver extract.

Collection of blood samples and estimation of blood lipid profile

Blood samples were collected from the animals one hour after administration of the drugs under mild ether anesthesia at the 14th and 28th day of the study. For the estimation of biochemical parameters blood samples were centrifuged at 3500 rpm for 20 min and the serum was stored at −20°C until further analysis.

The blood lipid profile included total cholesterol (TC), triglycerides (TG), high-density lipoprotein cholesterol (HDL-C) and low-density lipoprotein cholesterol (LDL-C). The above blood lipid parameters were determined with the help of reagent kits by Fluitest, Analyticon (Biotechnologies AG, Germany) (Stat fax 3300, Awareness Technology, USA). The LDL-C and very-low-density lipoprotein (VLDL) levels were calculated using the following formulae:

\[
LDL = TC - TG/5 - HDL, VLDL = TC - (HDL + LDL).
\]

Antioxidant assays

Ferric ion reducing antioxidant power (FRAP) assay

The oxidant in the FRAP assay was prepared by mixing 2,4,6-triprydyl-s-triazine (TPTZ), acetate buffer and ferric chloride monohydrate. The conglomerate was referred to as “FRAP reagent”. The final solution contained 1.67 mM of Fe(III) and 0.83 mM of TPTZ. To measure the FRAP value, 300 μl of freshly prepared FRAP reagent was warmed up at 37°C, a blank reading was taken at λ = 593 nm, and then 30 μl of tested sample and 30 μl of pure water were added. The absorbance reading was taken at λ = 593 nm [12, 13].

Determination of ABTS cation radical decolorization assay

ABTS, the oxidant, was produced by persulfate oxidation of 2,2’azinobis (3-ethylbenzothiazoline-6-sulfonic acid). This solution was diluted with phosphate buffer (pH = 7.4) until the absorbance reached 0.7 to 0.8 at λ = 734 nm. One ml of the resulting solution was mixed with the sample and the absorbance was read after mixing for 4 min at 30°C. The difference of the absorbance reading was plotted versus the antioxidant concentrations to give a straight line [13, 14].

Oxygen scavenging capacity assay (SOD)

The activity of SOD was assayed by using an assay mixture composed of 1.2 ml of 0.025 mmol/l sodium pyrophosphate buffer (pH = 8.3), 0.1 ml of phenazine methosulfate (186 mmol/l), appropriately diluted enzyme stock solution and water given to a total volume of 3 ml. After incubation at 30°C for 90 s, the reaction mixture was vigorously stirred and mixed with 4 ml of n-butanol. The color intensity of the chromatogram in the butanol layer was measured at λ = 540 nm against n-butanol [13, 14].

Determination of serum hepatic enzymes

The serum hepatic biochemical parameters such as aspartate aminotransferase (AST) and alanine aminotransferase (ALT) were estimated using an enzymatic test kit (Fluitest) and a chemistry analyzer (Stat fax 3300, Awareness Technology, USA).

Statistical analysis

The statistical tool GraphPad Prism 4 (GraphPad Software, USA) was used. The data were expressed as mean ± S.E.M (standard error of mean). Data were analyzed using one-way ANOVA, followed by Dennett’s multiple-comparison test. The IC50 values were calculated. The level of significance was set at p < 0.05.

Results

Effects of Achyranthes aspera seed saponins on body mass index (BMI) of high cholesterol diet (HCD) fed rats

Table I shows BMI (means ± SEM) at the starting day of HCD administration and after 14 days. In the untreated control group A (receiving only 2% aqueous gum tragacanth solution) a non-significant (p > 0.05) BMI change was observed. In con-
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In contrast, a significant increase of BMI ($p < 0.01$) was observed in group B receiving HCD for 2 weeks. In groups C, D, and E treated with lower doses of saponins from A. aspera, HCD also caused a significant ($p < 0.05$), albeit much smaller, increase of body weight. However, the BMI change was significant ($p < 0.05$) in groups F and G receiving high doses of saponins, as compared to group B, showing a dose-dependent decrease in BMI due to A. aspera seed saponins (Table I).

Regarding the results at day 28, no further increase of BMI was found in groups C to G treated with HCD along with the saponins. Remarkably, an additional highly significant ($p < 0.01$) increase of BMI was found in group B as compared with group A. The cumulative water and food intake was almost the same in all analyzed groups of animals and no change during the study was observed (data not shown).

**Effects of Achyranthes aspera seed saponins on lipid profile**

Administration of HCD to albino rats resulted in a significant ($p < 0.05$) increase of TC, TG, LDL-C and VLDL-C and a decrease of HDL-C levels in rats from group B, compared to the rats fed with a normal diet at the 14th and 28th day of our study. The LDL-C/HDL-C ratio was also found to be increased significantly ($p < 0.05$) (Figure 1). Dose-dependent protective effects of saponins isolated from A. aspera on blood serum lipid profile were noted after 14 days of treatment, as shown in Figure 1. Oral administration of A. aspera seed saponins significantly lowered the serum cholesterol level as compared to control animals only receiving HCD (Figure 1).

Administration of saponins of A. aspera for 28 consecutive days maintained the achieved improvement of lipid profile in a dose-dependent manner, compared to group B. The beneficial results of high saponin doses were comparable with the control standard drug (atorvastatin) on the 28th day of treatment (Figure 1).

**Effects of Achyranthes aspera seed saponins on serum antioxidant status**

Antioxidant activity of saponins of A. aspera seeds was judged by ABTS, FRAP and SOD assays. Administration of saponins of A. aspera seeds in different doses effectively acted as free radical scavengers, and demonstrated significant ($p < 0.01$) antioxidant activity (Figure 2). Administration of A. aspera seed saponins along with a high cholesterol diet caused a significant, dose-dependent improvement of the serum antioxidant status of treated rats. The FRAP content was reduced, whereas SOD and ABTS contents were significantly improved (Figure 2).

**Effects of saponins of Achyranthes aspera on hepatic functions**

To evaluate the hepatic status of rats treated with HCD and also any alteration due to oral administration of saponins from A. aspera for 28 days, AST and ALT were measured. Our results indicated mild hepatoprotective potential of saponins from A. aspera against HCD, as shown in Table I.

**Discussion**

All over the globe the incidence of hyperlipidemia increases day by day either due to genetic mutations or unhealthy lifestyle or secondarily due to other diseases [15, 16] and is becoming one of the major causes of death [17]. Pakistan is also facing the same threat [18]. Researchers have paid attention to natural sources to find new agents with hypolipidemic activity and having minimal side effects, as therapy with commercially available lipid-lowering drugs is often associated with significant side effects. Achyranthes aspera possesses potent antifungal, antiinflammation, antihyperlipidemic, antidiabetic, immunomodulatory, ant carcinogenic, diuretic, cardiotoxic, antiinflammatory, analgesic, and antibacterial activities [19]. The present study was conducted to evaluate the antiobesity, hypolipidemic and antioxidant effects of A. aspera seed saponins in hyperlipidemic albino rats.

Obesity is a major problem and can lead to many complications including hypertension, diabetes, myocardial infarction, etc. A vast range of natural products and medicinal plants, including crude extracts and isolated compounds from plants, can be used to induce weight loss and prevent diet-induced obesity. It has been reported that obesity can be improved by various mechanisms, i.e. by inhibition of lipase activity or delaying lipid absorption [20]. It has also been observed that dietary lipids are not directly absorbed from the intestine unless they have been subjected to the action of pancreatic lipasases. Inhibition of these digestive enzymes might be beneficial in the treatment of obesity. Although cumulative food intake was in our study similar in all groups, animals from group B showed a significant increase of body weight and BMI after consumption of HCD for a 4-week experimental period when compared to the normal diet group, duly supporting the idea that a high-fat diet may produce a marked increase in body weight and BMI compared with a normal diet [21–23]. It has been established that overnutrition from high calorie and fatty diets induces lipo- and glucoxicity, by increasing oxidative stress due to mitochondrial dysfunction with ROS overproduction, triggering the proinflammatory cascade, and potentiating tissue damage by
Figure 1. Mean total blood serum lipid parameters (mean mg/dl ± SEM, n = 6) on the 14th and 28th day after starting oral administration of 50, 75, 100, 125 and 150 mg of Achyranthes aspera seed saponins with high cholesterol diet (HCD) treated rats in comparison to normally, HCD and HCD + atorvastatin treated control rats (*p < 0.05 when compared with control receiving only normal diet with vehicle (2% gum tragacanth solution); **p < 0.05 when compared with animals receiving only high cholesterol diet with 2% gum tragacanth solution).
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**Figure 1.**

D

![Graph showing LDL-C levels](image)

Legend:
- Normal diet
- HCD
- HCD + 50 mg/kg A. aspera saponin
- HCD + 75 mg/kg A. aspera saponin
- HCD + 100 mg/kg A. aspera saponin
- HCD + 125 mg/kg A. aspera saponin
- HCD + 150 mg/kg A. aspera saponin
- HCD + atorvastatin

E

![Graph showing VLDL-C levels](image)

Legend:
- Normal diet
- HCD
- HCD + 50 mg/kg A. aspera saponin
- HCD + 75 mg/kg A. aspera saponin
- HCD + 100 mg/kg A. aspera saponin
- HCD + 125 mg/kg A. aspera saponin
- HCD + 150 mg/kg A. aspera saponin
- HCD + atorvastatin

F

![Graph showing LDL-C/HDL-C ratio](image)

Legend:
- Normal diet
- HCD
- HCD + 50 mg/kg A. aspera saponin
- HCD + 75 mg/kg A. aspera saponin
- HCD + 100 mg/kg A. aspera saponin
- HCD + 125 mg/kg A. aspera saponin
- HCD + 150 mg/kg A. aspera saponin
- HCD + atorvastatin

Figure 1. Cont.
Figure 2. Mean blood serum antioxidant profile (means ± SEM, n = 6) on the 28th day after oral administration of 50, 75, 100, 125 and 150 mg of A. aspera seed saponins with high cholesterol diet (HCD) in comparison to normally, HCD and HCD + atorvastatin treated control rats (*p < 0.05, when compared with normal control receiving only normal diet with vehicle (2% gum tragacanth solution), **p < 0.05 when compared with the control receiving only high cholesterol diet with 2% gum tragacanth solution)
Antioxidant and hepatoprotective effects of Achyranthes aspera seed saponins in high cholesterol fed albino rats

Lipid metabolism disorders have been reported as a widespread problem of modern civilization because they play a significant role in the development of generalized atherosclerosis. Increased serum concentrations of TC and TG have been reported as a meaningful abnormality as they are often associated with cardiovascular disorders and many other pathological syndromes. Both LDL and VLDL have a causative role in atherogenesis [29]. Treatment with saponins of A. aspera seeds significantly decreased the levels of TC, TG, LDL, VLDL, and LDL/HDL, whereas a significant increase was found in HDL-C.

Treatment with A. aspera seed saponins resulted in favorable changes of serum HDL-C, i.e. increased its level, which is currently recognized as a better atherosclerosis marker than each lipid parameter alone. It is possible that the hypolipidemic effects of A. aspera seeds may affect the reverse cholesterol transport of HDL. It has been established that HDL can be divided into two subclasses, HDL-2 and HDL-3; but still further studies are needed to determine the changes of these subclasses. Administering various doses of saponins of A. aspera seeds (50, 75, 100, 125, 150 mg/kg, p.o.) to hypercholesterolemic rats for 4 weeks decreased the susceptibility of LDL-C to oxidation.

A similar decrease of LDL-C also occurred on treatment with 8 mg/kg of atorvastatin, a standard therapy for hypercholesterolemia (Table II). Hypercholesterolemia and hypertriglyceridermia have been reported as major risk factors for atherosclerosis and related occlusive vascular diseases. Clinically relevant complications of atherosclerosis could be diminished and thus life prolonged when blood lipids are lowered by antihyperlipidemic/hypolipidemic drugs. Increased serum concentration of TC and TG is a major lipid abnormality syndrome as they are often associated with cardiovascular disorders and many other pathological syndromes. In the present study, a significant fall of TG levels was noted in tested animals consecutively treated with saponins of A. aspera seeds at different doses for 28 days. It may be suggested that this decrease of TG levels may be due to enhanced activity of lipases that hydrolyze TG under normal conditions or increased excretion of TG via feces. Whether the decrease of cholesterol level was due to reduction of resorption of endogenous cholesterol or an increase in the rate of secretion into the intestinal tract, or both, remains to be elucidated in the future. HDL-C has been reported to benefit the body in two ways: it removes cholesterol from the walls of arteries and returns it to the liver, and it helps to prevent oxidation of LDL-C. In fact, it appears to have antioxidant properties on its own. HDL-C then helps to keep arteries open and reduces the risk of heart attack.

Table II. Mean blood serum enzyme level (IU/dl ± SEM) on 28th day after starting oral administration of 50, 75, 100, 125 and 150 mg of A. aspera seed saponins given with HCD in comparison to normal, HCD and HCD + atorvastatin treated control rats

| Treatment group | Serum hepatic parameters |
|-----------------|--------------------------|
|                 | AST [IU/dl] | ALT [IU/dl] |
| A               | 60.57 ±2.99   | 18 ±1.98    |
| B               | 77.77 ±3.4*   | 24.14 ±3.18*|
| C               | 65.71 ±6.04*  | 23.14 ±3.07*|
| D               | 64 ±4.86*     | 20.57 ±3.86*|
| E               | 72.43 ±5.99   | 19.29 ±1.80*|
| F               | 70.86 ±7.55*  | 21.29 ±2.70*|
| G               | 68.29 ±6.18*  | 20.43 ±2.50*|
| H               | 70.71 ±6.54*  | 21.57 ±4.35*|

Data represented as mean ± SEM (n = 6). *p < 0.05, **p < 0.01 when compared with group A animals (normal control) receiving only normal diet with vehicle (2% gum tragacanth solution). *p < 0.05, **p < 0.01 when compared with group B animals (negative control) receiving only high cholesterol diet with vehicle (2% gum tragacanth solution).
antioxidative effect could not be obtained in vitro with all of these drugs, it seems that active drug metabolites, which are formed in vivo, could affect lipoprotein oxidizability [32, 33]. It has been reported that hypercholesterolemia leads to increased production of reactive oxygen species (ROS), which cause lipid peroxidation. Reactive oxygen species initiate a series of chain reactions in vivo and ultimately damage tissue and DNA. Superoxide dismutase and catalase have been reported as major enzymes that play an important role in the elimination of reactive oxygen species producing oxidative damage [34, 35]. Antioxidants are compounds capable of donating a single electron or hydrogen atom for reduction [36]. Antioxidant effects of saponins of *A. aspera* have been investigated in terms of ABTS, FRAP and SOD assay both in control and treated rats. Significant improvement was observed in the antioxidant status of the serum of rats treated with saponins of *A. aspera* seeds, i.e. by considerable improvement of FRAP and ABTS levels and increase of SOD. Higher FRAP values give higher antioxidant capacity because the FRAP value is based on reducing ferric ion, where antioxidants are the reducing agent. To conclude, *A. aspera* seed saponins appeared to ameliorate the oxidative damage, probably due to the presence of phenolic and flavonoids.

To evaluate the effect of consumption of HCD and saponins of *A. aspera* on the hepatic status of animals, serum AST and ALT were measured. It was found that saponins of *A. aspera* have no hepatotoxic effect but instead they possess hepatoprotective potential by normalizing the levels of serum AST and ALT in HCD-treated rats.

In conclusion, the results of the present investigation clearly indicate that saponins of *A. aspera* seeds possess hypolipidemic properties showing hypotriglycerideremic or hypcholesterolermic effects with undetectable side effects on liver and cardiac functions. Due to the significant effect on lipid profile, research is still ongoing to investigate the same hypolipidemic activity of different extracts from seed powder. This may lead to activity-guided fractionation of that particular extract in order to better know the phytochemical composition, isolation methods, and structure elucidation of some of the bioactive constituents followed by establishing the most probable mechanism of action of reducing serum lipids.

Based on the above results, we propose that *A. aspera* seed saponins, if properly formulated, may be used as potent hypolipidemic agents also showing beneficial antioxidant and hepatoprotective properties.

**Conflict of interest**

The authors declare no conflict of interest.

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