

**EFFECT OF HYDROALCOHOLIC EXTRACTS OF *SIDA CORDIFOLIA L.*  
LEAVES ON LIPID PROFILE IN RATS**

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**Summary**

*Sida cordifolia* L (Malvaceae) is important medicinal plant of Ayurvedic system of medicine and used as home remedy as cardioprotective agent without any scientific background. In this investigation an attempt was made to study the effect of hydroalcoholic extract of *Sida cordifolia* L (HESC) on serum lipid profile, which is closely associated with many cardiovascular diseases. Albino rats were administered HESC (100 mg/kg and HESC 500 mg/kg) once daily orally for 30 days. Accordingly, serum total cholesterol (TC), triglyceride (TG), low-density lipoprotein cholesterol (LDL-C) and high-density lipoprotein cholesterol (HDL-C) were among the parameters investigated. We also evaluated the activities of serum aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels as a measure of hepatic cell damages. It was found in our study that the administration of both doses of HESC to animals fed with high fat diet (HFD) lowered their serum TC, TG and LDL-C significantly, while raised the serum HDL-C. Treatment with HESC reduced serum ALT and AST levels compared to HFD groups. Further, it was also observed that the high dose of HESC significantly decreases serum TC, TG and LDL-C and increases HDL-C in animals fed with normal fat diet (NFD). Based on these data, it is concluded that the chronic administration of HESC in high dose up to 500 mg/kg has definite cardioprotective potential and these results constitute a valid scientific basis for consuming *Sida cordifolia* L for medicinal application.

**Key words:** Hydroalcoholic extract of *Sida cordifolia* L; high fat diet; lipid profile; normal fat diet.

## **Introduction**

Hyperlipidaemia is considered a predisposing factor involved in the development of cardiovascular disease (1, 2). The search for new drugs capable of reducing and regulating serum cholesterol and triglyceride levels has gained momentum over the years, resulting in numerous reports on significant activities of natural agents (3). Plant products are frequently considered to be less toxic and free from side effects than synthetic agents. These properties have led to the discovery of new therapeutic agents including antioxidants, hypoglycemics, and hypolipidemics. It is well established that diet rich in vegetables and fruits can reduce cardiovascular diseases (4-6). In view of the increasing use of herbal remedies by the general public and subsequent interest by the physicians, it is imperative to promote credible research for validation of their claimed activities based on modern scientific techniques. In India *Sida cordifolia* L. (Family: Malvaceae) is popularly known as “Country mallow” and “Bala” found along roadsides throughout the tropical and subtropical plains of India and Sri Lanka. The leaves of the plant are reported to possess analgesic, anti-inflammatory (7), anticancer (8), diuretic, laxative, hypoglycemic (9) and hepatoprotective (10) activities. Further, studies showed that aqueous fraction of hydroalcoholic extract of leaves induce vasorelaxation (11), hypotension and bradycardia (12). It is also considered as an excellent anti-oxidant (13). The aim of the present study was to evaluate the influence of the HESC on serum lipid levels in rats fed high-fat diets and normal fat diet. The lipid profile was taken as the major marker of hypercholesterolaemia. Accordingly, serum total cholesterol (TC), triglyceride (TG), low-density lipoprotein cholesterol (LDL-C) and high-density lipoprotein cholesterol (HDL-C) were among the parameters investigated. In addition, to have a measure of the effects of the plant extract on liver function of the treated rats, the serum aspartate aminotransferase (AST), alanine aminotransferase (ALT) and alkaline phosphatase (ALP) activities were estimated relative to plant-untreated control rats.

## **Methods**

### *Plant Collection and Extraction*

The leaves of *Sida cordifolia* L. (Voucher Specimen No.-RRCBI 3276) were purchased from medicinal garden of foundation for revitalization of local health traditions, Attur, via Yelahanka, (Bangalore, India) and authenticated by Regional Research Institute (Ay.), Bangalore, India. The leaves were manually separated and dried at 70 °C for 72 h; then ground to a granulated powder with a grinder. The powdered leaves were extracted in a soxhlet extractor with petroleum ether (60 °C

for 8 h) and defatted leaves powder was re-extracted in a soxhlet apparatus for 72 h with methanol at 60 °C. The % yield of leaves extract was calculated.

#### *Experimental animals*

Laboratory bred female Wistar albino rats weighing between 200-250 g were housed at 25° ± 5°C in a well-ventilated animal house under 12:12 hour light and dark cycle. Institutional Animal Ethics Committee approved the experimental protocol; animals were maintained under standard conditions in an animal house approved by Committee for the Purpose of Control and Supervision on Experiments on Animals (CPCSEA).

#### *Acute toxicity studies*

The acute toxicity study was carried out according to the limit test described by the OPPTS guidelines (<http://www.epa.gov/oppts/home/guideline.htm>). Test dose of 2 g/kg and 5 g/kg were given to mice. Both doses were found to be safe. Hence, 1/10<sup>th</sup> and 1/50<sup>th</sup> of the maximum safe dose corresponding to 100 mg/kg and 500 mg/kg orally were selected as low and high dose respectively.

#### *Experimental Protocol*

The animals were either fed normal fat diet (NFD) or high fat diet (HFD). The NFD comprised of standard rat chow (Amrut Laboratory Animal feed, Maharashtra, India) containing protein 22.10%, oil 4.13%, fibre 3.15%, ash 5.15%, sand (silica) 1.12% w/w), whereas, HFD was composed of standard rat chow - 68%, Dalda (saturated fat) - 30% and cholesterol - 2% (14).

#### *Measurement of body weight and diet intake*

Diet intake was measured daily and the body weights were recorded before starting the plant treatments and also after each week of the plant treatments.

#### *Treatment*

The rats were divided into six groups, each containing six animals: the first three group of rats was fed NFD (Group I-III); first group was considered as NFD control, whereas, the second and third groups were treated with low and high dose of HESC (HESC-100 & HESC-500 mg/kg) respectively. The second three groups were fed HFD (Group IV-VI); first among them was kept as HFD control, while, second and third were administered HESC-100 and HESC-500 mg/kg. The administration of HESC was done orally at same time once daily for 30 days. On

14<sup>th</sup> and 21<sup>st</sup> days, the overnight fasted rats were anesthetized with diethyl ether and their blood samples were collected by retro orbital vein and centrifuged at 3000 rpm, 4° C for 15 min. After 30 days of treatment, the animals were fasted overnight and then sacrificed under diethyl ether anesthesia. Blood samples were taken from the animals of all groups. Serums were separated from the blood samples and were stored at -70° C pending biochemical analyses. The triglycerides (TG), total cholesterol (TC) and HDL-cholesterol (HDL) were estimated using autoanalyzer (15-17). The LDL-cholesterol was calculated by using the formula: LDL in mg/dl=Total cholesterol- (HDL-Cholesterol-1/5Triglycerides) (18).

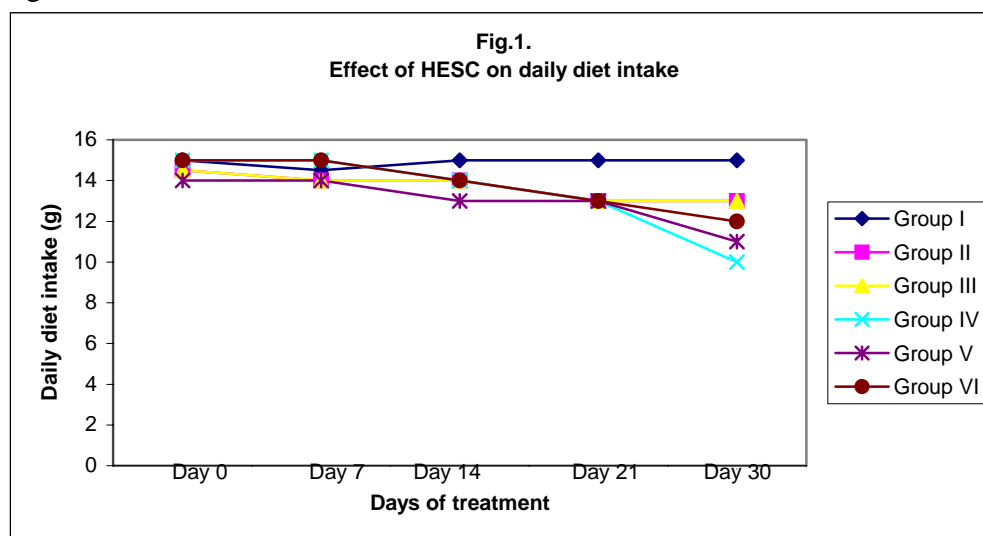
#### Statistical analysis

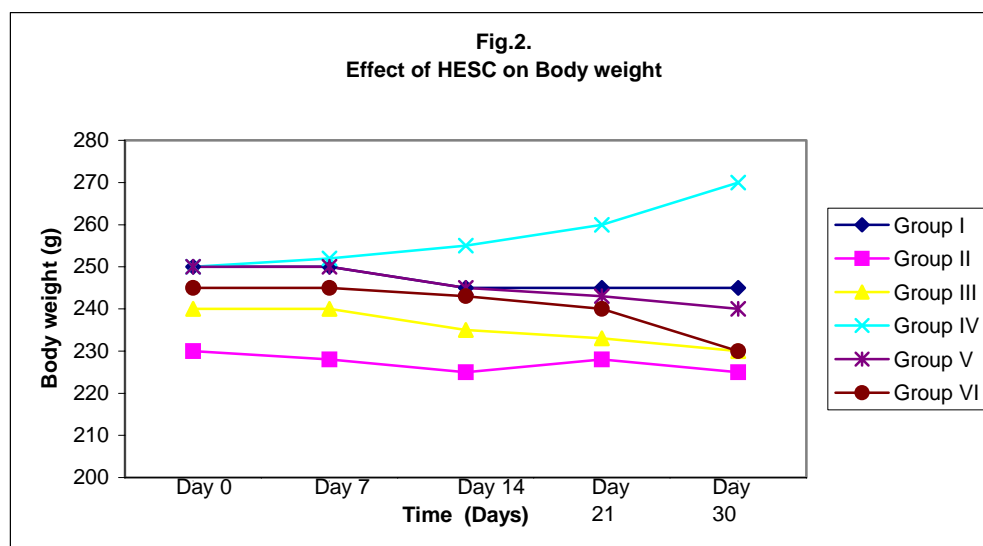
Results are expressed as mean  $\pm$  SEM. Statistical significance was assessed using One-way Analysis of variance (ANOVA) followed by Tukey multiple comparison tests.  $p < 0.05$  was considered significant.

### Results

#### Effects of HESC on body weights

The daily diet intake of the rats decreased, in groups II, III, IV, V and VI, from the 21<sup>st</sup> day of the experiment (Fig. 1). High-fat diet apparently caused an increase in the body weight of rats despite their lower food intake. However, in Group III and IV the body weights of the plant treated rats decreased after day 21, as shown in Fig. 2.

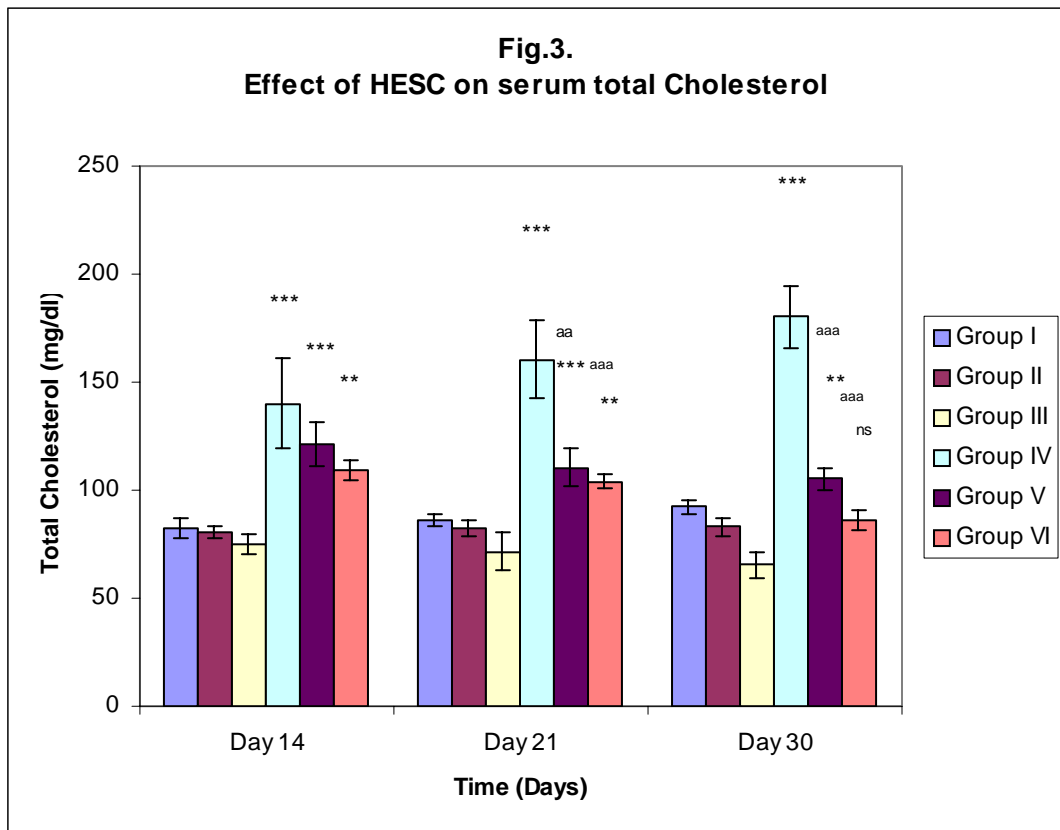


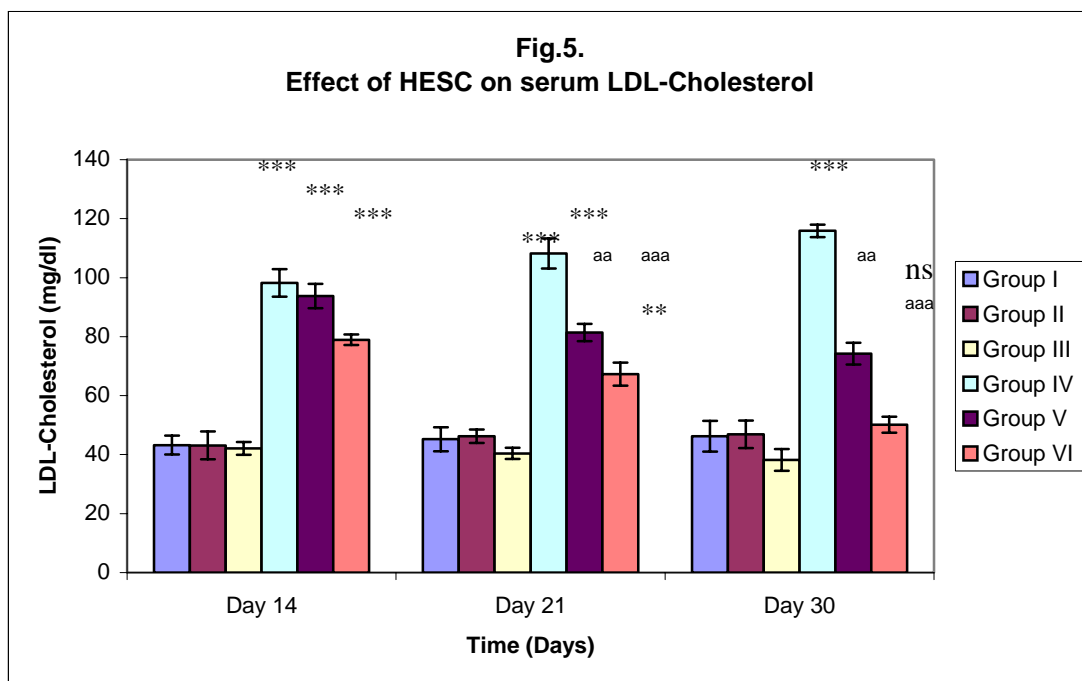
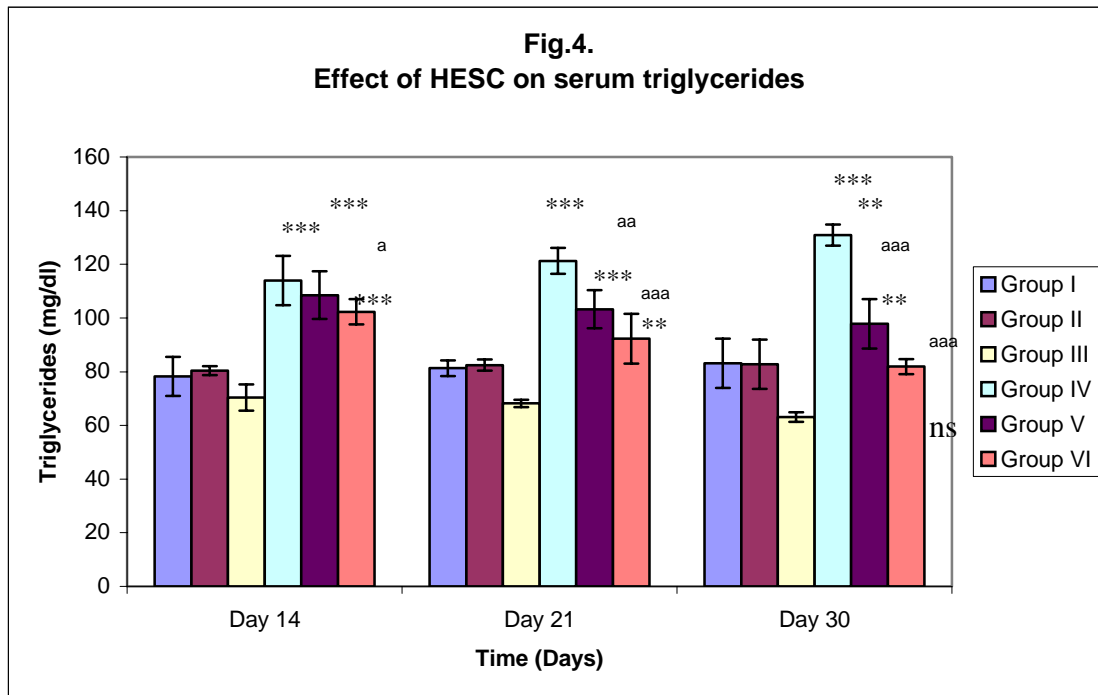


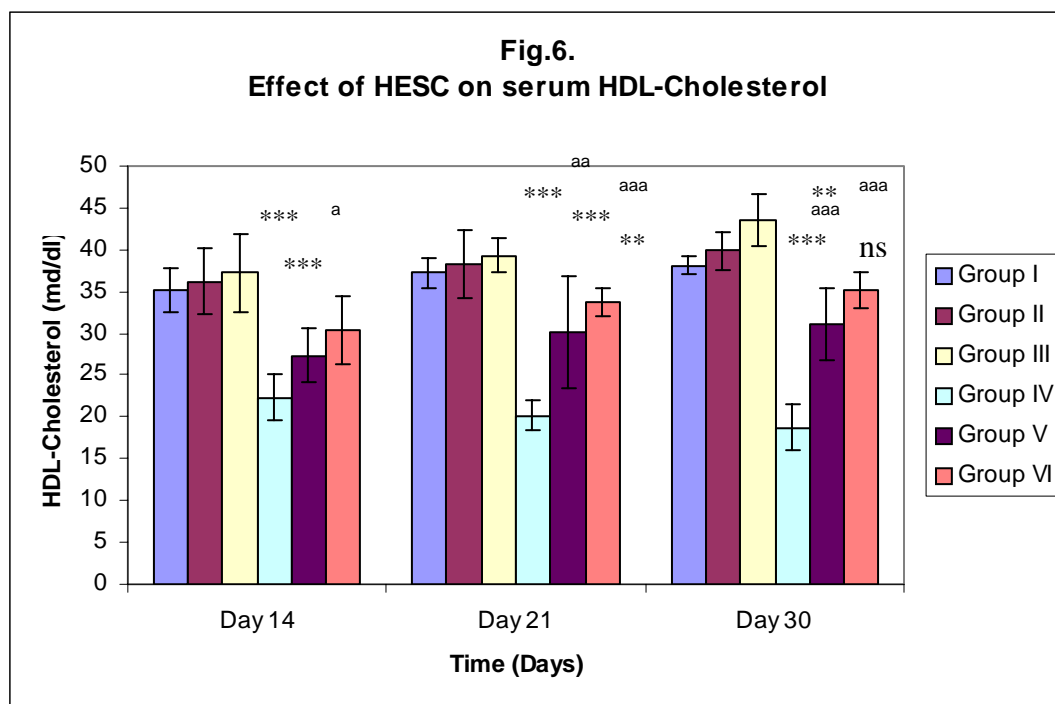
Values are expressed as mean  $\pm$  SEM for six rats in each group; Group I, NFD; Group II, NFD+HESC-100; Group III, NFD+HESC-500; Group IV, HFD; Group V, HFD+HESC-100; Group VI, HFD+HESC-500.

#### *Effect of HESC on serum lipid profile*

The general effect of HESC on serum lipid levels was examined in rats fed NFD and HFD. In NFD groups, there was no significant difference in serum lipid levels of rats. Feeding the animals a high-fat diet resulted in a significant increase in the serum total cholesterol (TC), triglycerides (TG), and low-density lipoprotein-cholesterol (LDL-C) levels. However, the level of serum high-density lipoprotein-cholesterol (HDL-C) decreased (Fig. 3-6) compared to NFD control group. Treatment of HFD rats with HESC at a dose of 100 and 500 mg/kg body weight for 21 days shows lowering of lipid level. After 30 days of continuous treatment with plant extract, almost the same observations were made demonstrating the hypolipidaemic effect of high dose of HESC (serum TC, TG, LDL-C and HDL-C were same like NFD control). There was significant reduction in lipid levels of TC, TG and LDL-C in HESC groups when compared to HFD control group. On the other hand, there was significant elevation in HDL-C level in HESC treated groups fed HFD.







Values are expressed as mean  $\pm$  SEM for six rats in each group; \*\*\* Significantly different from NFD control; <sup>aaa</sup> Significantly different from HFD control in HFD-HESC groups  $P < 0.001$ ; Group I, NFD; Group II, NFD+HESC-100; Group III, NFD+HESC-500; Group IV, HFD; Group V, HFD+HESC-100; Group VI, HFD+HESC-500; <sup>a</sup> \*  $p < 0.05$ ; <sup>aa</sup> \*\*  $p < 0.01$ ; <sup>aaa</sup> \*\*\*  $p < 0.001$ ; ns, not significant when compared to NFD control.

#### *Effect of high-fat diet on hepatic enzymes in serum*

The effects of HESC on serum aminotransferases (AST, ALT) are shown in table. Animals fed high-fat diet exhibited an elevation in serum AST (GOT) and ALT (GPT) compared to NFD control group. Treatment with plant extract reduced activities of AST and ALT compared to animals of NFD control, after 21 days. After 30 days of continuous treatment with plant extract, almost the same observations were made.



Table. Effect of HESC on hepatic enzymes in serum

Treatment	ALT (GPT)		AST (GOT)	
	Day 21	Day 30	Day 21	Day 30
Group I	17.80±3.9	16.93±2.1	24.90±3.0	23.30±2.5
Group II	18.90±2.7	17.80±3.3	23.70±2.1	23.20±1.9
Group III	16.30±1.9	16.10±2.2	22.20±1.7	20.21±1.6
Group IV	38.20±3.7 <sup>***</sup>	41.21±1.2 <sup>***</sup>	40.20±3.9 <sup>***</sup>	42.21±1.4 <sup>***</sup>
Group V	30.70±2.3 <sup>**aa</sup>	28.90±1.3 <sup>*aa</sup>	31.30±1.9 <sup>**aa</sup>	30.10±1.8 <sup>*aa</sup>
Group VI	22.30±2.27 <sup>aaa</sup>	18.90±2.9 <sup>aaa</sup>	27.20±1.3 <sup>aaa</sup>	25.70±2.0 <sup>aaa</sup>

Values are expressed as mean  $\pm$  SEM for six rats in each group; <sup>\*\*\*</sup> Significantly different from NFD control; <sup>aaa</sup> Significantly different from HFD control in HFD-HESC groups  $P < 0.001$ ; Group I, NFD; Group II, NFD+HESC-100; Group III, NFD+HESC-500; Group IV, HFD; Group V, HFD+HESC-100; Group VI, HFD+HESC-500; <sup>a</sup> \*  $p < 0.05$ ; <sup>aa</sup> \*\*  $p < 0.01$ ; <sup>aaa</sup> \*\*\*  $p < 0.001$ ; ns, not significant when compared to NFD control.

### Discussion

The main causative factor for atherothrombotic diseases is the disturbances occurring in lipid metabolism. Despite the presence of different hypolipidaemic drugs in the market, their therapeutic application is usually associated with severe side effects (19). Hence, efforts are being made to find safer and more efficient anti-hyperlipidaemic drugs. In that respect, medicinal plants have been considered as promising resources for the discovery of new drugs. In the present study, the hyperlipidaemic activity of HESC was evaluated among the hypercholesterolaemic rats. Based on our data, the body weight of rats of Group IV increased relative to NFD control group. This is probably due to supplementation with cholesterol and cholic acid. Treatment with HESC affectively prevented the increase in body weight to a significant extent.

Cholesterol is an essential structural element of the biological membranes. In addition, it is the precursor of many compounds such as the starting materials for the synthesis of bile acids, steroid hormones, and vitamin D among others. Despite this knowledge, high concentration of serum cholesterol increases the risk of developing CHD (20). The present study demonstrated that rats fed a HFD showed

a higher concentration of serum TC compared to rats fed a NFD, while oral administration of HESC reduced the high level of TC. It has been reported that low plasma TG levels are associated with lower risks of CHD (21). In our study, the administration of HESC to hyperlipidaemic rats significantly lowered their serum TG levels. This reduction might be related to decreased triglyceride absorption as well as increased excretion of TG via feces (22). It is widely accepted that elevation of plasma LDL levels are major risk factors for CHD (23). Direct correlation between LDL and atherosclerosis and also the reversibility of the related pathological events by lowering the serum level of LDL has already been reported by many research groups (23,24). Our data indicated that, the high concentration of LDL-C in hypercholesterolaemic rats was significantly reduced by oral administration of HESC. Therefore, HESC might constitute a good candidate for the treatment of atherosclerosis by lowering serum LDL-C level. Another risk factor for developing atherosclerosis is the reduced serum level of HDL-C. This effect of HDL is largely attributed to its central function in the reverse cholesterol transport, a process whereby excess cell cholesterol is taken up and processed by HDL particles for further delivery to the liver for metabolism (25). Therefore, it is logical that an increase in HDL level can contribute to lower risk of atherosclerosis (26). Our results clearly showed that HESC is capable of increasing the serum level of good cholesterol (i.e. HDL-C) in the treated rats relative to HFD rats. Our data clearly indicated that HESC is capable of potentially decreasing this risk factor after 21 days of drug treatment. Phytochemicals, especially the phenolics found in fruits and vegetables, have been proposed as the major bioactive compounds providing the health benefits associated with diets rich in plant-foods. Flavonoids are a class of secondary plant phenolics found ubiquitously in fruits and vegetables as well as food products which act as pharmacological active compounds in many medicinal plants (27,28). Many of the biological action of flavonoids have been attributed to their powerful hypolipidaemic properties (29,30). It is reported that flavonoids intake decreased LDL-C and increased HDL-C in hypercholesterolaemic individuals that may hasten removal of cholesterol from peripheral tissue to liver for catabolism and excretion (31). Independent studies have confirmed the presence of antioxidant phenolic compounds, mainly flavonoids HESC (10). Regarding the high total phenolics and flavonoids of HESC, the results strongly suggest that the hypolipidaemic activity of this medicinal plant could be attributed to the presence of the valuable polyphenolic compounds. It might be possible that these active constituents reverse the adverse effects associated with hyperlipidaemia. (30). The decrease in cholesterol balance, which indicates the total change in body pools of cholesterol, may be due to the compensatory mechanisms, such as a decrease in resorption of endogenous cholesterol or an increase in the rate of secretion into intestinal tract or both. The effect(s) of HESC on liver function was evaluated by

measuring the serum transaminase activities of ALT and AST. These enzymes leak into the circulation when hepatocytes are damaged (32). It is believed that high serum cholesterol level can cause liver damages (33). Our results show that high-fat diet has caused significant increase in serum ALT and AST levels. However, rats treated with HESC had lower serum ALT and AST levels compared to rats of Group NFD control.

Thus on the basis of the result of the present research, it is concluded that the high dose of HESC is a powerful agent for combating elevated levels of lipids during CHD and other cardiac manifestation. However, further studies should to carry out to understand the influence of HESC during myocardial infarction.

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