

Effects of *Acacia nilotica* Leaf Extract on Adrenaline-induced Hyperlipidemia and Cardiac Remodeling in Rats

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Abstract

Hyperlipidemia is one of the major causes of disability in western as well as in developing countries. The present study was designed to investigate the effect of the ethanolic leaf extract of a Bangladeshi medicinal plant *Acacia nilotica* (AN) on adrenaline-induced hyperlipidemia and cardiac remodeling in rats. The plant extract and commercial lipid lowering drug atorvastatine were administered in adrenaline-induced hyperlipidemic rats (AIHRs) through intraperitoneal route for one week at a dose of 30 mg and 0.04 mg per 150 gm body weight of rat, respectively. The AN extract showed a significant ($P < 0.01$) decrease in lipid profile, and reducing the left ventricular hypertrophy (heart weight/body weight), heart weight, cardiac myocytes size as well as kidney, pancreas and liver weight of AIHRs. The observation was also made for consecutive four weeks (long term treatment) to confirm the results obtained in one week model. The results indicated that the leaf extract of *A. nilotica* in comparison with atorvastatine has got profound hypolipidemic and anticardiac remodeling activities.

Key words: Hyperlipidemia, Fabaceae, *Acacia nilotica*, Atorvastatine, Cardiac remodeling.

Introduction

Atherosclerosis and coronary heart disease are the major health problem in developed and modern societies and leading causes of death in diabetic patients (Stamler *et al.*, 1993; Wilson, 2001). Coronary vessel disease and atherosclerosis being primary reasons for the increased incidence of cardiovascular dysfunction (Lteif *et al.*, 2003). The American heart association estimates that 65% of patients with diabetes die due to some form of cardiovascular disease (CVD), hence, good control of blood sugar levels can reduce the risk of CVD (Bhopal and Rafnsson, 2007; Richard and Mary, 2007). Cardiac remodeling, defined as the alteration in the dimensions, mass, shape and function of the heart, is generally accepted as a determinant of the clinical course of heart failure (HF). Although patients with major remodeling demonstrate progressive worsening of cardiac function, slowing or reversing remodeling has only recently become a goal of HF therapy (Opie *et al.*, 2006).

The use of traditional medicine has played an important role in nearly every culture on earth, including Asia, Africa, Europe and America. Traditional medicine is based on plants contain chemicals that can promote health

and alleviate illness (Grabely and Thiericke, 1999). Herbal treatments are the most popular form of traditional medicine, and are highly lucrative in the international marketplace due to availability, low cost and less side effect. Some traditional medicinal plants have already been reported to work against cardiovascular diseases (Wong, 2011) as well as high blood cholesterol (Aattar, 2006).

Acacia nilotica, locally known as babla, belongs to the family fabaceae. The plant is widely distributed in the Indo-Bangla subcontinent and cultivated throughout the tropical belt. Different parts of this plant are reported to be used for the treatment of human ailments. The whole plant has antiplatelet aggregatory activity (Shah *et al.*, 1995). Methanol extract of leaf of *A. nilotica* possesses hypotensive and hypospasmodic activities (Gilani *et al.*, 1999). The leaves are also reported to be a good hypocholesterolemic agent (Ghasi *et al.*, 2000). An aqueous extract of the seed of *A. nilotica* was shown to possess sustained dose-related contractile activity on isolated guinea-pig ileum (Amos *et al.*, 1999). On the basis of the reported pharmacological activity, the plant was selected for hypolipidemic and anticardiac

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remodeling investigation and the aim of our present study is to investigate the effects of *A. nilotica* leaf extract on adrenaline induced hyperlipidemia and cardiac remodeling in rats.

Materials and Methods

Plant materials: Fresh leaves of the plant *A. nilotica* were collected from the Botanical Garden of the University of Rajshahi and authenticated at the Department of Botany, University of Rajshahi, Rajshahi, Bangladesh.

Preparation of crude leaf extract: The leaves of AN were dried completely under the mild sun and ground with an electric grinder into coarse powder. The powdered material was then extracted with 95% ethanol and room temperature for 7 days with occasional shaking and stirring. The extract was filtered through cotton filter and the filtrate was then concentrated with a rotary evaporator under reduced temperature and pressure to get brownish mass.

Animal studies: All protocols for the animal experiments were reviewed and approved by the animal care and use committee of Institute of Biological Science, University of Rajshahi. Long-Evans male rats weighing about 110-130 mg, aged 2 months, were purchased from animal's house of International Centre for Diarrhoeal Disease Research, Bangladesh (ICDDR,B). Prior to commencement of the experiments, all the rats were acclimatized to the new environmental condition for a period of one week. During the experimental period, the rats were kept in a well ventilated animal house at room temperature and were supplied standard pellets from ICDDR,B and fresh drinking water. All the rats were kept in cages and maintained with natural 12 hour light and dark cycle. Adrenaline was injected intra-peritoneally (i.p.) in rats. AIHRs were treated with i.p. injection of AN at various doses for dose selection. After completing the dose selection, AIHRs were treated with the most effective observed dose of AN (30 mg /150 gm b.w.) and adrenaline (0.04 mg/150 gm BW) for both 1 and 4 weeks.

Measurement of lipid profile: After completing the treatment with ethanolic extract of AN leaves and atorvastatin, the rats were first anesthetized with sodium phenobarbital. Then abdominal skin was cut, and thoracic artery was opened. Finally, 3-5 ml of blood was collected

directly from the thoracic artery by syringes. At last, the blood was centrifuged at 4000 rpm for 10 minutes, and the serum was obtained. Serum lipid profiles, such as TC, TG, LDL-C and HDL-C, were assessed using diagnostic kits (Human, Germany).

Histopathological analysis: Histological studies were performed for the determination of LV weight and cardiomyocyte cross sectional area. In brief, LV weight was measured, and the ratio of LV weight to body weight was calculated to determine an index of LV hypertrophy. Four sections were obtained from each heart, and mounted on slides and stained with hematoxylin and eosin. To evaluate the extent of cardiomyocyte hypertrophy, cross-sectional images of cardiomyocyte were scanned at $\times 400$ magnifications. Approximately 20 cross-sections of cardiomyocyte were analyzed in each heart. Average values for each heart were used for analysis. All images were taken using an Olympus BX51TF (Olympus Corporation, Tokyo, Japan), and all measurements were determined using Scion Image software (Scion Corporation, Frederick, MD).

Statistical analysis: The results were expressed as mean \pm SEM. We used a one-way analysis of variance (ANOVA), followed by Dunnett's post-hoc test or students paired or unpaired *t*-test where appropriate. The statistical method applied in each analysis was described in each figure. Results were considered to be significant when *p* values were less than 0.05 ($p < 0.05$).

Results and Discussion

Effects of AN leaf extract on TC, TG, LDL-cholesterol, and HDL-cholesterol level in adrenaline induced hyperlipidemic rats (AIHRs): AIHRs were treated with ethanolic extract of AN at 30 mg/150 gm b.w. for one week and it was observed that AN leaf extract significantly reduced total cholesterol level (1.805 mmol/l) compared to atorvastatin (1.34 mmol/l). AN leaf extract also reduced triglyceride level (2.022 mmol/l) as well as LDL-cholesterol level (1.95 mmol/l) and raises HDL-cholesterol level (1.50 mmol/l) in adrenaline induced hyperlipidemic rats (Figure 1A-1D).

Four weeks treatment with AN extract also significantly reduced total cholesterol level (1.34 mmol/l), triglyceride level (1.98 mmol/l), and LDL-cholesterol level (1.99 mmol/l) and raised HDL-cholesterol level (1.48 mmol/l) in adrenaline induced hyperlipidemic rats (Figure 2A-2D).

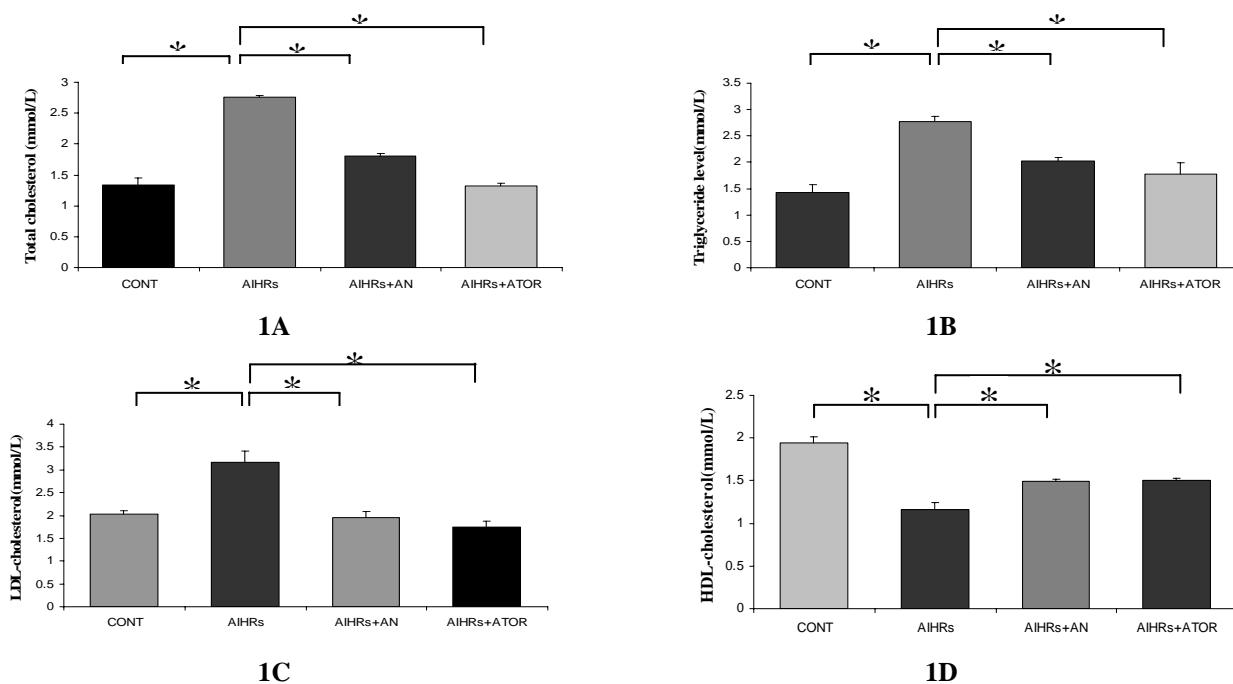


Figure 1. Effects of AN leaf extract on lipid profile in AIHRs for one week. Figures (1A-1D) represent the estimation of TC, TG, LDL, and HDL-C levels, respectively, after i.p administration (30 mg/150 gm b.w.) of AN in AIHRs for one week. All values are presented as mean \pm SEM, n=4 in each case. * P <0.05 vs control. ** P <0.01 vs AIHRs. ATOR = Atorvastatine, AN = *Acacia nilotica* leaf extract

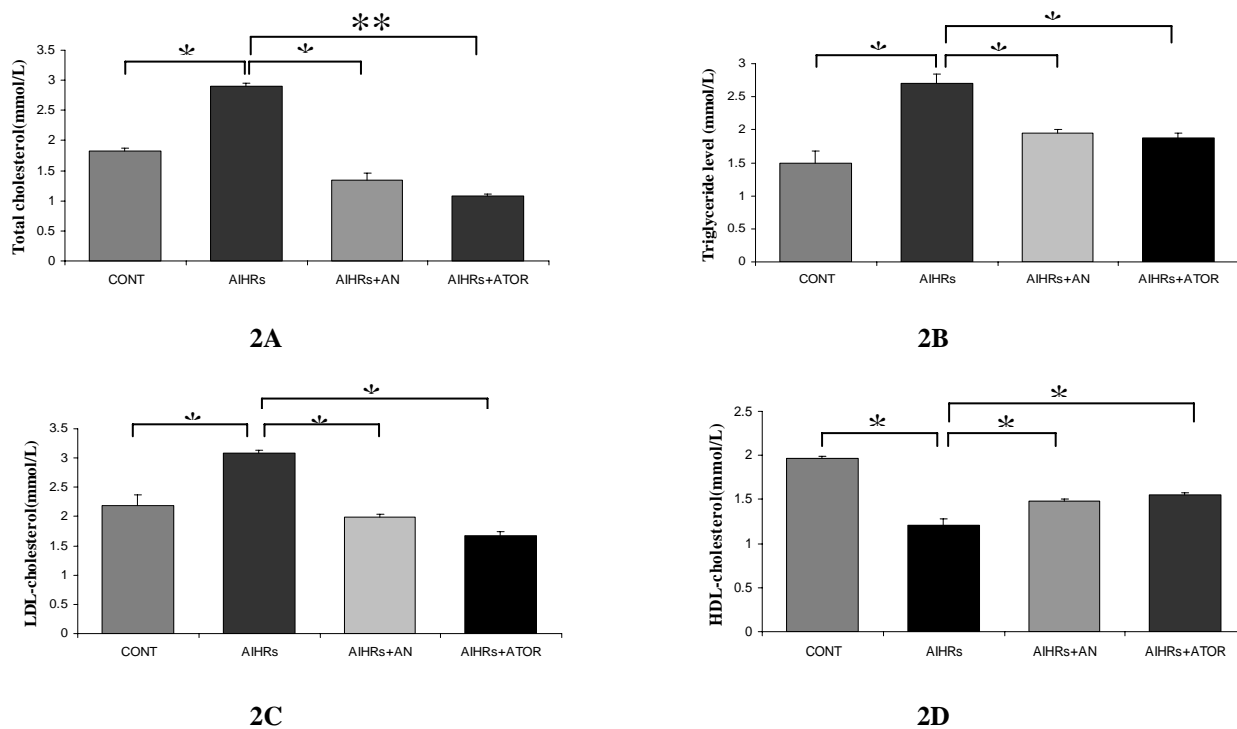


Figure 2. Effects of AN leaf extract on lipid profile in AIHRs for four weeks. Figures (2A-2D) represent the estimation of TC, TG, LDL, and HDL-C levels, respectively, after i.p administration (30 mg/150 gm b.w.) of AN in AIHRs for two weeks. All values are presented as mean \pm SEM, n=4 in each case. * P <0.05 vs control. ** P <0.01 vs AIHRs. ATOR = Atorvastatine, AN = *Acacia nilotica* leaf extract

Effects of Acacia nilotica (AN) leaf extract on left ventricular hypertrophy, heart weight, and cardiac myocytes in AIHRs: AN leaf extract as well as atorvastatine significantly reduced left ventricular hypertrophy (0.004 and 0.0038 for one week vs 0.0033 and 0.0028 for four weeks protocol, respectively) in adrenaline induced hyperlipidemic rats (Figure 3A-3B).

Induction of adrenaline in rats for one week significantly increased heart weight (0.7gm) in comparison with control rats (0.3gm). Treatment with AN extract and atorvastatine reduced heart weight (0.5 and 0.4 gm, respectively) in AIHRs. For four weeks protocol similar results were obtained where adrenaline increased heart weight (0.9gm) in comparison to control rats

(0.4gm). Treatment with AN leaves extract and atorvastatine reduced heart weight (0.6gm and 0.5 gm, respectively).

Figures 3C and 3D demonstrate the effects of AN leaves extract on cardiac myocytes size in AIHRs. Administration of adrenaline in rats for one week as well as four weeks increased cardiac myocytes cross sectional area (639.75 and 643.75 μm^2 for one and four weeks, respectively) in comparison with control rats (417.5 and 408.75 μm^2 , respectively). Treatment with AN leaves extract significantly reduced cardiac myocytes cross sectional area (430 and 431.25 μm^2 for one and four weeks, respectively) compared to atorvastatine (425 and 418.75 μm^2 for one and four weeks, respectively).

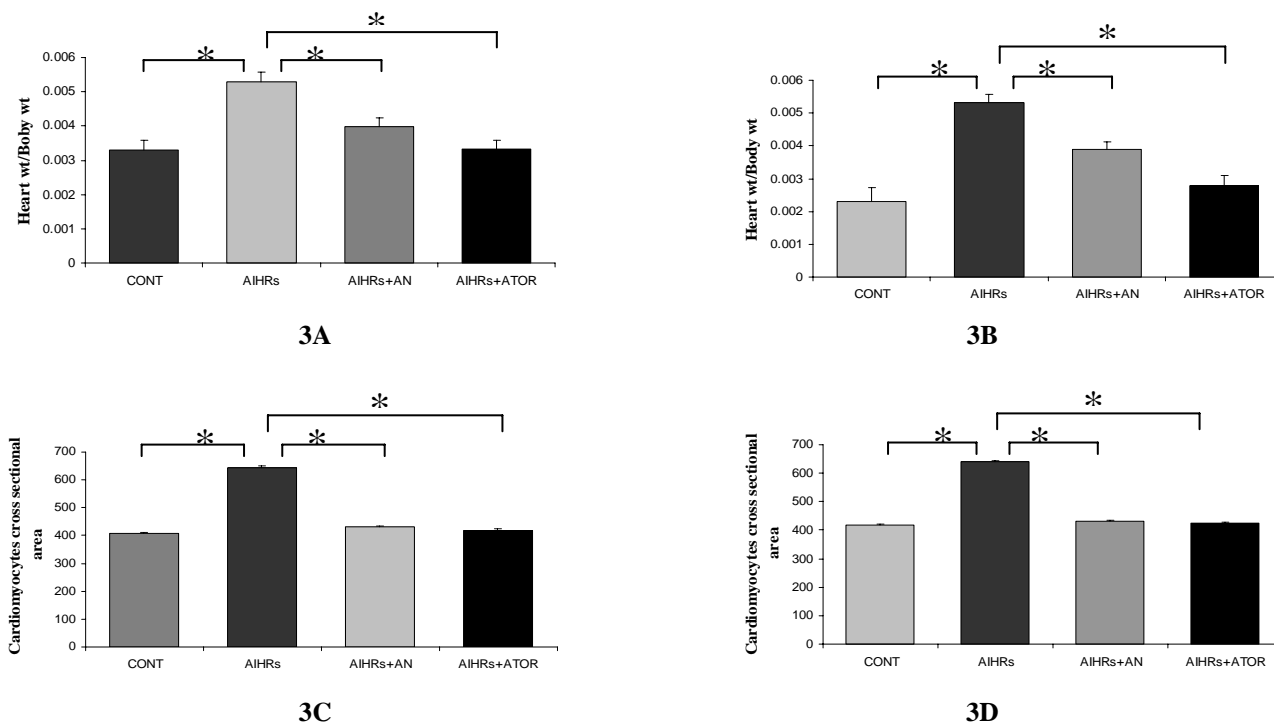


Figure 3. Effects of AN leaf extract on left ventricular hypertrophy and cardiac myocytes in AIHRs for one and four weeks. Figure 3A and 3B represent the effect on left ventricular hypertrophy, whereas 3C and 3D represent the effect on cardiac myocytes in AIHRs for one and 4 weeks, respectively. The data are shown as mean \pm SEM, n=4 in each case * $P < 0.05$ vs control. ** $P < 0.01$ vs AIHRs. ATOR = Atorvastatine, AN = *Acacia nilotica* leaves extract

Effects of AN leaf extract on kidney, pancreas and liver weight (in gm) on AIHRs for one and four weeks: Figures 4A and 4B demonstrate the effects of AN leaves extract on kidney, pancreas and liver weight on AIHRs. Infusion of adrenaline in rats for one week significantly increased kidney, pancreas and liver weight (0.7, 0.6, and 5.2 gm, respectively) in comparison with control (0.4, 0.4 and 3.1 gm, respectively). Treatment with AN leaf extract

significantly reduced kidney, pancreas and liver weight (0.6, 0.5 and 4.8 gm, respectively) in AIHRs and the result is comparable to atorvastatine (0.5, 0.4, and 4.2 gm, respectively). Similar results were observed for four weeks study (Figure 4B).

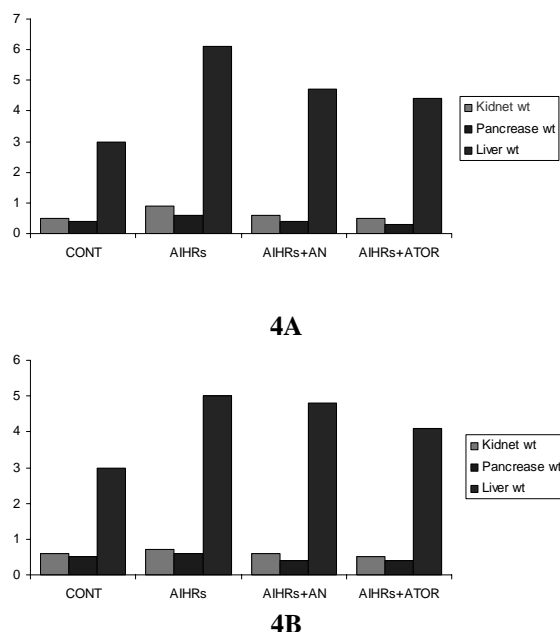


Figure 4. Effects of AN leaf extract on kidney, pancreas and liver weight in gm on adrenaline induced hyperlipidemic rats (AIHRs). 4A and 4B are the effects for one week and four weeks, respectively. ATOR = Atorvastatine

Discussion

Our study explored that *A. nilotica* leaf extract reduced high lipid profile and cardiac remodeling in adrenaline induced hyperlipidemic rats as shown in figure 1 to 4. The present study revealed that *A. nilotica* extract at a dose 30 mg per 150gm body weight of rats has got profound lipid lowering activity and reducing left ventricular hypertrophy, heart weight, cardiac myocytes size when compared with the control rats and AIHRs. Asian herbal medicine reported that most potent constituent of *A. nilotica* leaf is reserpine which is recommended for the treatment of hyperlipidemia (Jigam *et al.*, 2010) and works by decreasing heart rate and relaxing the blood vessels so that blood can flow more easily through the body. Present study with *A. nilotica* leaf extract also evaluates the hypolipidemic effect *in vivo* on adrenaline induced hyperlipidemic rats. This relevant experimental study proved that the crude leaf extract has antihyperlipidemic properties and reduce cardiac change. Thus intake of *A. nilotica* leaf extract as medicine might have potential benefit in the treatment of hyperlipidemia and cardiac remodeling.

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