



Research Article

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Cardioprotective Property of Extracts of *Dialium guineense* Stem Bark in Rats Exposed to CCl_4

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Abstract

The aim of this study was to investigate the cardioprotective property of extracts of *Dialium guineense* stem bark in rats exposed to carbon tetrachloride (CCl_4). A total of twenty-five (25) adult male Wistar rats, which weighed between 170 and 190 g (mean weight = 180 ± 10 g) were randomly assigned to five groups (5 rats per group): normal control, CCl_4 control, silymarin, aqueous extract and ethanol extract groups. Cardiac injury was induced with CCl_4 (single oral dose of 1.0 mL/kg body weight, bwt). Rats in the silymarin group were administered 100 mg silymarin/kg bwt (standard cardioprotective drug), while those in the two treatment groups received 1000 mg/kg bwt of aqueous or ethanol extract for a period of 28 days. Activity of aspartate aminotransferase (AST), level of nitric oxide (NO) as well as atherogenic index of plasma (AIP) were measured. Rat heart was also subjected to histopathological examination. The results showed that the activity of AST as well as levels of low-density lipoprotein cholesterol (LDL-C), AIP, atherogenic coefficient (AC) and cardiac risk ratio (CRR) were significantly higher in CCl_4 control group than in normal control group, but they were markedly reduced after treatment with extracts of *D. guineense* stem bark ($p < 0.05$). However, the extracts significantly promoted the release of NO from the coronary endothelium of rats exposed to CCl_4 ($p < 0.05$). Carbon tetrachloride (CCl_4) markedly disrupted the normal architecture of rat's hearts inducing fibrosis, necrosis, and haemorrhage. However, treatment with aqueous and ethanol extracts of the medicinal plant led to marked regeneration of cardiomyocytes. The toxic cardiac injury induced by CCl_4 was significantly blocked by the plant extracts.

Key words: Aspartate aminotransferase, Cardiac function, *Dialium guineense*, Nitric oxide, Tissue histology

Introduction

Cardiac injury induced by drugs is becoming an increasingly important concern, since it constitutes serious risk to human health [1]. Cardiac dysfunction caused by cardiotoxic agents such as CCl_4 may lead to heart failure, myocardial ischemia, arrhythmias, hypertension, myocarditis, pericarditis, and thromboembolism [2]. Oxidative and nitrate stress, as well as protein adduct formation are established mechanisms of chemical-induced cardiotoxicity. The latter leads to cardiomyocyte inflammation, altered calcium homeostasis, programmed cell death (apoptosis), swelling of cardiomyocytes, nuclear splitting, vacuolization, and alteration in signaling pathways. These events result in diseases of the heart muscle including heart failure, which could lead to death [3].

Medicinal plants are sources of natural antioxidants which are used for the treatment of different ailments. These plants produce important therapeutic effects [4]. One of such plants is *Dialium guineense*. It is a tall, tropical, fruit-bearing tree, belonging to the

Leguminosae family, and has small, typically grape-sized edible fruits with brown hard inedible shells. In Africa, it grows in dense forests along the southern edge of the Sahel [5]. The plant grows naturally in West African countries, Central African Republic, and Sudan [6,7]. Different parts of the medicinal plant are used against several diseases [5]. Extracts of the plant have been shown to be reservoirs of important phytochemicals [8-10]. At present little or nothing is known about the cardioprotective capacity of extracts of *Dialium guineense* stem bark. The aim of this study was to investigate the cardioprotective property of extracts of *D. guineense* stem bark in rats exposed to CCl_4 .

Materials and Methods

Chemicals

Analytical grade chemicals and reagents were used in this study, and they were purchased from Sigma-Aldrich Ltd. (USA).



Collection of Plant Material

The stem barks of *D. guineense* were procured from Auchi, Edo State, Nigeria. Identification and authentication of the plant took place at the herbarium of the Department of Plant Biology and Biotechnology, University of Benin, Benin City, Nigeria (No. UBHD330).

Plant Preparation and Extraction

The stem barks of *D. guineense* were washed and shade-dried at room temperature. The plant material was then pulverized using mechanical blender. Extraction of the stem bark was carried out as described in literature [11,12].

Experimental Rats

A total of twenty-five (25) adult male Wistar rats, which weighed between 170 and 190 g (mean weight = 180±10 g) were purchased from Anatomy Department, University of Benin, Benin City, Nigeria. The rats were housed in metal cages under standard laboratory conditions, and they were allowed free access to rat feed and clean drinking water. Prior to commencement of the study, the rats were acclimatized to the laboratory environment for a period of seven days. The study protocol was approved by the University of Benin Faculty of Life Sciences Ethical Committee on Animal Use.

Experimental Design

The rats were assigned to five groups (5 rats per group): normal control, CCl₄ control, silymarin, aqueous extract and ethanol extract groups. Cardiac injury was induced with CCl₄ (single oral dose of 1.0 mL/kg bwt). Rats in the silymarin group were administered 100 mg silymarin/kg bwt (standard cardioprotective drug), while those in the two treatment groups received 1000 mg/kg bwt of aqueous or ethanol extract for a period of 28 days.

Blood and Tissue Sample Collection and Preparation

At the end of the treatment period, the rats were euthanized. Blood samples were collected from the anesthetized rats via cardiac puncture in heparinized sample bottles. Rat hearts were also excised and used to prepare 20 % tissue homogenate. The blood and homogenate were centrifuged at 2000 rpm for 10 min to obtain supernatants which were used for biochemical analysis.

Biochemical Analyses

Activity of AST, and levels of NO and LDL-C were measured [13-15]. Atherogenic index of plasma (AIP), AC and CRR were also determined [16-17].

Histological Examination of the Tissues

Portions of the heart were serially sectioned and fixed in 10 % formalin for 48 h. The specimen was then dehydrated using varied

concentrations of ethanol and cleared in three changes of xylene before embedment in paraffin. Serial sections (4 µm thick) were made and stained with haematoxylin and eosin (H&E) according to standard method. Histological assessment was performed under light microscopy using an image analyzer (Image Proplus, version 3.0).

Statistical Analysis

Count data are expressed as mean±SEM (n=5). The statistical analysis was performed using SPSS (version 20). Groups were compared using Duncan multiple range test. Statistical significance was assumed at p<0.05.

Results

Effect of Extracts of *D. guineense* Stem Bark on Relative Organ Weight

There were no significant differences in relative organ weight among the groups (p > 0.05) (Table 1).

Group	Relative Organ Weight x 10 ⁻²
Normal Control	3.34 ± 0.54
CCl ₄ Control	3.02 ± 0.10
Silymarin	3.17 ± 0.16
Aqueous Extract	3.52 ± 0.24
Ethanol Extract	2.96 ± 0.14

Note*: Data are relative organ weights and are expressed as mean ± SEM (n=5).

Effect of Extracts of *D. guineense* Stem Bark on Cardiac Function

As shown in Figures 1-3, the activity of AST as well as levels of LDL-C, AIP, AC and CRR were significantly higher in CCl₄ control group than in normal control group, but they were markedly reduced after treatment with extracts of *D. guineense* stem bark (p < 0.05). However, the extracts significantly promoted the release of NO from the coronary endothelium of rat's hearts exposed to CCl₄ (p < 0.05) (Figures 1 -4).

Histology of normal control rat heart revealed bundles of myocardial fibres, interstitial space, and coronary artery (normal cardiac architecture), while that of CCl₄ control showed congested coronary artery, mild myocyte hypertrophy and fibrosis. Interstitial inflammatory cells infiltrate, necrosis and haemorrhage were visible. Histopathological examinations of silymarin and the two treatment groups revealed markedly reduced interstitial oedema and significant reduction in cellular debris. There was also mild collagen deposition.

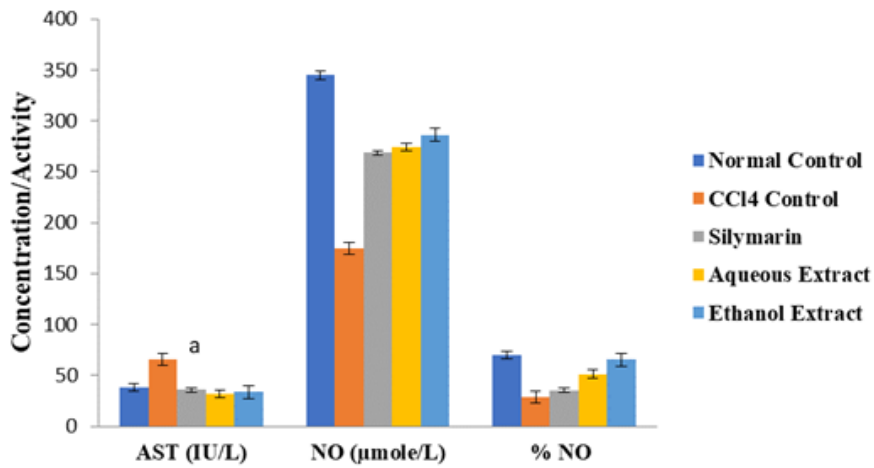


Figure 1: Effect of Extracts of *D. guineense* Stem Bark on Markers of Cardiac Function.
 Note*: Data are markers of cardiac function and are expressed as mean ± SEM. ^ap < 0.05, when compared with CCl₄ control group.

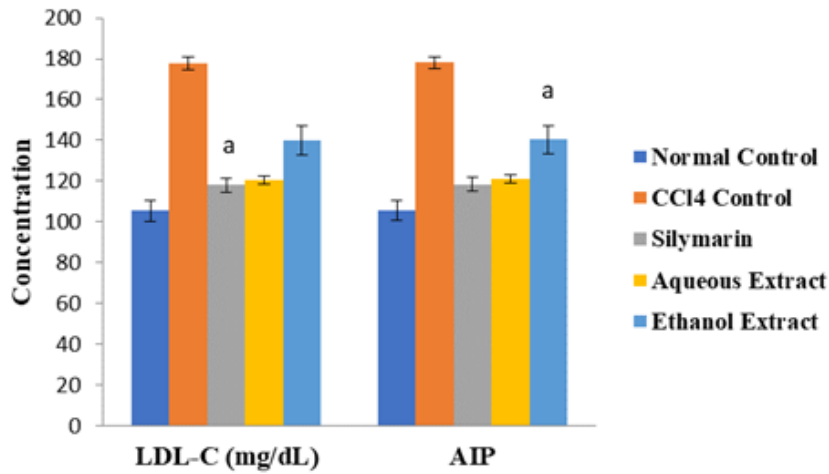


Figure 2: Comparison of the Levels of LDL-C and AIP Among the Groups.
 Note*: ^ap < 0.05, when compared with CCl₄ control group.

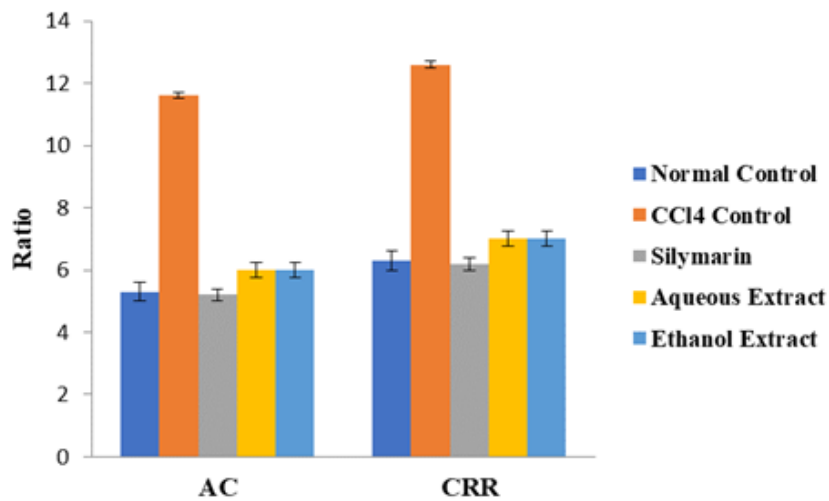


Figure 3: Comparison of AC and CRR Among the Groups.
 Note*: ^ap < 0.05, when compared with CCl₄ control group.

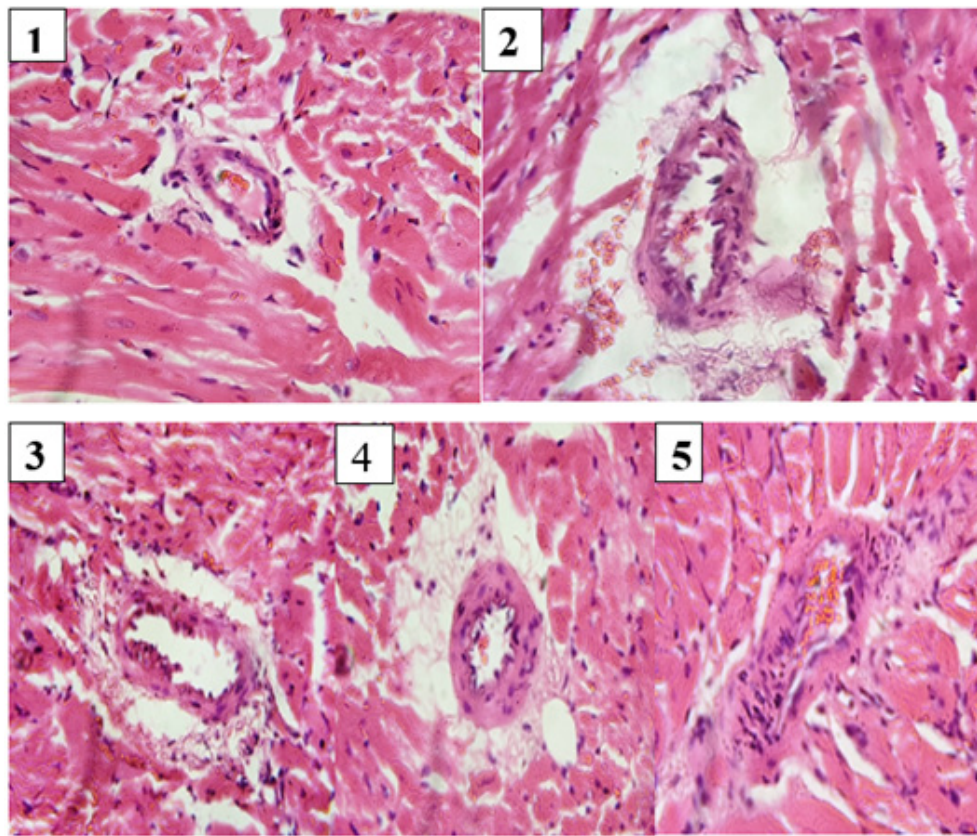


Figure 4: Effect of Extracts of *D. guineense* Stem Bark on Histology of Rats Hearts.

Discussion

Carbon tetrachloride (CCl_4) is a well-known substance for induction of tissue injury via generation of free radicals. The compound is highly toxic to organs/tissues such as liver, kidneys, heart, lung, testis, brain and blood [18]. It is biotransformed by hepatic microsomal cytochrome P450 to trichloromethyl radical, which initiates lipid peroxidation [19,20]. Free radicals' formation, a rate limiting process in tissue peroxidative damage, is the generally accepted mechanism of CCl_4 -induced cardiotoxicity [21]. This leads to loss of myocardial structural integrity and depressed cardiac function [21]. The aim of this study was to investigate the cardioprotective property of extracts of *D. guineense* stem bark in rats exposed to CCl_4 . The results showed that CCl_4 administration significantly increased lipid profile (LDL-C and AIP) and these results agree with those of previous reports [22]. Oxidative stress impairs the cell's ability to remove free cholesterol as high-density lipoprotein cholesterol, thus increasing the amounts of cellular free cholesterol. Antioxidants have been shown to inhibit all these mechanisms [23]. A high LDL-C level is associated with increased risk for heart disease and stroke. Low-density lipoprotein (LDL) transports cholesterol to the arteries, and when its level is elevated, this cholesterol accumulates in the blood vessel walls and contribute to the formation of plaque [24]. Treatment with

aqueous and ethanol extracts of *D. guineense* stem bark significantly decreased all the parameters of cardiovascular disease risk factors, measured in this study. The effects were comparable to those of the standard cardioprotective drug, silymarin. The protective property of the extracts may be due to the abundant phytochemicals which act as antioxidants [8,10&25]. Previous studies have elucidated the mechanistic association of antioxidant effect with hyperlipidemic cardiomyopathy [26,27].

Histological examination of the myocardium, valves and cardiac blood vessels is often as important as the gross examination. The diagnostic features and categories of heart disease are many and varied, possibly more than any other organ. In this study, histopathological findings of heart tissues further established the results regarding the effects of extracts of the medicinal plant on anatomical (heart mass index) and biochemical parameters. Heart sections of normal control rats revealed normal structures of heart, while rats exposed to CCl_4 showed congestion, degeneration of myocardial tissue, and necrosis as a confirmation of cardiotoxicity. Reversals of such damage were observed in silymarin- and extracts-treated rats. It is likely the extracts possess antiplatelet activity. Platelets, platelet products and thrombosis play important roles in the occurrence of acute occlusive vascular events, including myocardial infarction (MI) and ischemic stroke, since the disruption

of platelet- and fibrin-rich atherosclerotic plaque may be followed by aggressive platelet deposition and, ultimately the development of a thrombus that can precipitate an acute occlusive event. Decreased platelet aggregation could be a plausible mechanism for the cardioprotective effects of the extracts [28].

Conclusion

The results of this study indicate that aqueous and ethanol extracts of *D. guineense* stem bark possess cardioprotective property against CCl_4 -induced cardiotoxicity. Further study aimed at the isolation, identification, and characterization of the active component(s) of the extracts and elucidation of their possible molecular mechanism of action is recommended.

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