The Assessment of *Alchornea Cordifolia* Activity in Aflatoxicosis Reduction in Rat.

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Abstract:

**Introduction:** Aflatoxins are cytotoxic and serve as one of the key risk factors of hepatocellular carcinoma. Currently, plants and extract are widely used as potential scavenging substances for the detoxification of mycotoxins. Thus, this study aims to investigate the activity of the crude ethanolic leaves extract from *Alchornea cordifolia* in aflatoxicosis prevention.

**Material and methods:** The phytochemical screening was performed through qualitative analysis based on coloring and/or precipitation reactions. Groups of rats were treated daily with a mixture dose of aflatoxin B1 (AFB1) at 150 µg/kg and the crude extract of *Alchornea cordifolia* at doses of 50, 100, and 300 mg/kg for 21 days. The body weight, biochemical, and histological assessments were determined.

**Results:** The phytochemical screening revealed the presence of polyphenols, flavonoids, sterols and terpenoids, quinoid compounds, tannins catechic and alkaloids. AFB1 treatment caused a significant increase of transaminases, urea, and creatinine abundances but reduced the rates of albumin and total proteins. *Alchornea cordifolia* administration alleviated biochemical parameters and body weight gain compared with the AFB1 group (p<0.05). The histological lesions of organs (liver and kidney) caused by AFB1 were significantly improved after administration of the extract at a dose of 300 mg/kg.

**Conclusion:** This plant plays a beneficial role in AFB1-induced injury and may be used in the treatment of aflatoxicosis.

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Introduction

Aflatoxin B1 (AFB1) is one of the principal food contaminants and is mainly produced by mold strains such as *Aspergillus flavus* and *Aspergillus parasiticus*. This mycotoxin is one of the key risk factors of hepatocellular carcinoma (HCC) in human being in sub-Saharan Africa and in the Middle East [12, 34]. Many other reported harmful effects of aflatoxins such as growth delay, immune abnormalities and hepatic affections [13, 42]. These toxic and carcinogenic effects caused by aflatoxin B1 are due to the oxidization of this molecule in the liver by P450 cytochromes of the families of CYP1A2, CYP2A6 and CYP2A4. These enzymes convert AFB1 into its carcinogenic form: the AFB1-8,9-epoxyde which establishes covalent bonds with the DNA and serum albumin to respectively produce AFB1-N7-guanine and AFB1-Lysin adducts [11, 36]. These enzymes can oxidize AFB1 in several other metabolites such as Aflatoxin M1 (AFM1) which is regarded as possible carcinogen for human being [18]. Several studies showed the capacity of scavengers in the protection against carcinogenesis and other toxicities caused by aflatoxins administration in pre-pretreatment or simultaneously with the carcinogenic. Synthetic scavengers such as butylated hydroxyl toluene (BHT), butylated hydroxyl anisole (BHA) and the propyl gallate showed their capacity to inhibit carcinogenesis caused by AFB1. However, the toxicity of these synthetic scavengers is a stem for their use in aflatoxicosis prevention [7, 20, 41]. The purpose of this present study is to evaluate the scavenging potential of *Alchornea cordifolia* in the body protection against aflatoxicosis.

Material and Methods

Chemicals:

Aflatoxins B1 (AFB1) was bought at Fermentek Ltd (Jerusalem, Israel) and the Butylated hydroxyanisole (BHA) at Santa Cruz Biotechnology (Delaware, Canada). Kits of serum aspartate aminotransferase (AST), alanine aminotransferase (ALT), urea and creatinine were obtained from Randox Laboratories (Antrim, UK). Albumin and total protein measuring kits were purchased from QCA, (AMPOSTA, Spain). All the other reagents used, were of analytical quality.

Plant Material

The leaves of *Alchornea cordifolia* were collected in the suburbs of Abidjan in March 2016. They were identified at the National center of Floristic (CNF) of the University of Felix Houphouët Boigny (Abidjan, Cote d’Ivoire). The leaves were dried for two weeks at ambient temperature.

Preparation of the Crude Ethanolic Leaves Extract of *Alchornea Cordifolia*.

The dry leaves of *Alchornea cordifolia* were ground using a grinder and 100 grams of the powder were macerated in one liter of ethanol (95 %) and shaken using a magnetic stirrer for 72 hours at ambient temperature. Then, the alcoholic solution was filtered...
twice on absorbent cotton then on a Whatman filter paper. The filtrates obtained were put together and evaporated at reduced pressure at 45°C using a rotary evaporator. The mixture was then dried in an oven at 45°C for 72 hours. Finally, a brown black powder was obtained (13.33 g) representing the ethanolic leaves extract of *Alchornea cordifolia* (EELAc). Different concentrations of EELAc were prepared on the spot during experiment by dissolving the powder obtained in an olive oil.

**Phytochemical Screening**

The different chemical analyses were carried out through a phytochemical screening. It is a qualitative analysis based on coloring and/or precipitation reactions. This analysis was performed on the ethanolic crude leaves extract of *Alchornea cordifolia* according to the methodology described by Houghton [17]. Table 1 points out the various chemical groups investigated and the specific reagents used.

**Animal Material**

Experiments were carried on white albino male rats, *Rattus norvegicus* of Wistar strain. These adult rats were between 12 to 16 weeks of age with a mean body weight of 152 ±4.5 g. Animals were fed with pellets and water *ad libitum*. They were acclimatized in cages for two weeks for them to be accustomed to the experiment environment. All experiments were performed in accordance with the European guidelines 2010/63/EU related to animals care [8].

**Products Administration**

Six groups of six male rats were set out. AFB1 (150 μg/kg) and tested substances (*A. cordifolia*: 50, 100, 300 mg/kg and BHA; 50 mg/kg) were dissolved in olive oil and the mixture was administrated orally to animals at a rate of 0.5 ml for 100 g/kg Bw [28] according to the model of table 1. The administrations are done five days per week for twenty one days with resting periods (table 1). In this experiment, the BHA was used as a standard scavenger. Its effects were compared with those of *Alchornea cordifolia*.

**Sampling and Dosage of Required Parameters**

**Biochemical Parameters**

On completion of the experiment, blood samples were collected in rat orbital sinus after being anesthetized using ether diethyl. Blood samples were centrifuged at 4000 tours/min for 10 minutes. Collected serum was put in an aliquot and preserved at -20°C for analyses. AST, ALT, albumin, total proteins, urea and creatinine were dosed with a commercial kit according to the manufacturer recommendations. Parameters were evaluated using a biochemistry automate of the type A 15 bio-system (Barcelona, Spain).

![Table 1. Methods used for the phytochemical screening and administration of tested substances](image-url)
Spain).

**Histological Evaluation**

All the animals were sacrificed by an anesthesia (ether diethyl) at the end of the experiment, organs (liver, kidneys) were immediately taken from each animal and preserved in formalin at 10 %. Organ samples were then colored with hematoxylin and eosin and were observed through optical microscope.

**Statistical Analysis**

Statistical analysis and graphs were performed using the soft-ware Graph Pad Prism 5.01 (San Diego, CA, United States). The results are the arithmetic mean of individual values assigned to the standard error (mean ± SEM). The results were analyzed statistically by ANOVA 2 followed by multiple comparison tests with Tukey–Kramer method. The difference between groups was considered significant when a probability (p) was <0.05.

**Results**

**Phytochemical Screening**

From 100 g of *Alchornea cordifolia* leaves, we obtained 13.33 g of crude leaves extract of *Alchornea cordifolia*, representing a yield of 13.33 %. The phytochemical screening of the ethanolic leaves extract of *Alchornea cordifolia* showed the presence of polyphenols, the flavonoids, sterols and terpenoids, quinoid compounds, catechic tannins and alkaloids (Table 2).

**Effect of Aflatoxin B1 on Rats Treated with EELac**

Animals of group 1 treated with olive oil for 21 days, showed a significant (P < 0.05) body weight growth of 52.46 g at the end of experiment (Table 3). Rats from groups 2, 3 and 4 showed a body weight growth ranging from 10.5 g (group 2) to 25.3 g (group 4). These body weight growths, though positive, were significantly inferior to that of group 1 (Control). The body weight gain of rats from group 5 and 6 were respectively 45.16 and 47.6 g. The body weight growth were not significantly different between group 5 and 6 compared to group 1 (Control). The animals showed a positive body weight gain ranging from 10.5 to 52.46 g. The evolution of the body weight growth of group 5 is similar to those of groups 1 and 6 treated respectively with the olive oil (Control) and standard scavenger.

**Effect of the Ethanolic Extract of Alchornea Cordifolia on the Biochemical Parameters.**

**Effect of the Extract on Hepatic Enzymes.**

The treatment of rats with the various
substances caused a significant increase (P < 0.05) of transaminases rates in group 2 and 3 compared to the control (group 1). This increase was not dose dependent and varied from 262.4 ± 8.1 UI/L (group 2) to 257.5±9.2 UI/L (group 3) for ALT and from 274.16±7.3 UI/L (group 2) to 282.5±3.1 UI/L (group 3) for AST (Table 4). In contrast, there is no significant difference between these rates for group 4, 5 and 6 compared to the control (group 1). These rates were 115.4±7.7 UI/L (group 4), 100.22±2.3 UI/L (group 5) and 90.75±2.4 UI/L (group 6) for ALT and 203.2±4.9 UI/L (group 4), 180.13±5.5 UI/L (group 5) and 186.25±6.7 UI/L (group 6) for AST.

Effect of the Extract on Renal Function

Substances administration caused a significant increase of creatinine, urea rate and a reduction of albumin and total proteins rates for group 2, 3 and 4 compared to the control. These values increase from 7.96±1.3 mg/l (group 4) to 10.74 ±1.3 mg/l (group 3) for creatinine and from 0.58 ± 0.02 g/l (group 4) to 0.76 ± 0.05 g/l (group 2) for urea and decrease from 23 ± 1.1g/l (group 4) to 19.5± 2.5 g/l for albumin and from 41.82 ±3.4 g/l to 34.26 ± 2.1 (g/l) for total proteins. The evolution of these biochemical parameters is not statistically significant between groups 5 and 6 compared to the control (P < 0.05) (Table 4). The results of the biochemical parameters showed that the evolution of transaminases, urea, albumin and total proteins in group 5 was similar to that of group 6 which received the standard scavenger (BHA). There was no significant difference between the values of these parameters in groups 5 and 6 compared to the control group (1) at P < 0.05.

Results of the histopathological evaluations Hepatic tissues

The administration of AFB1 alone caused a sinusoid capillary dilation with hemorrhagic contents (figure 1 A). In contrast, the administration of the mixture AFB1+EELac at a dose of 300 mg/kg (group 5)
or AFB1+ BHA at a dose of 50 mg/kg (group 6) caused the disappearance of these histological abnormalities of the liver (figure 1 B, C).

Renal Tissues

The histopathological examinations of the kidney showed that AFB1 caused an atrophy of renal tubes (figure 2 D). However, the administration of the mixture AFB1+EELac at a dose of 300 mg/kg (group 5) or AFB1+ BHA at a dose of 50 mg/kg (group 6) caused the disappearance of these histological abnormalities observed in the renal tubes (figure 2 E, F).

Discussion

Extraction and Phytochemical Screening

Many methods were developed to fight aflatoxicosis. Several compounds such as food, scavengers, plants, pharmaceutical products are used in various experimental studies to modulate the damage entailed by AFB1 on some organs (liver, kidney, lungs) and macromolecules (DNA, proteins) [10, 14, 16]. Hepatocellular carcinoma induced by AFB1 can be modified in animals by chemo-protective substances such as phenolic antioxidants (butylated hydroxy anisole, ethoxyquine) [21, 43]. In this study, the effect of the crude ethanolic leaves extract of *Alchornea cordifolia* (EELac) was investigated on aflatoxicosis. The choice of *Alchornea cordifolia* was made according to an ethnobotanical survey which demonstrates that, this plant is widely used for the treatment of several pathologies linked to oxydative stress such as diabetes, high blood pressure and hemorrhoids. Moreover, this plant can be easily found in Côte.d'Ivoire. The ethanolic extraction of *Alchornea cordifolia* leaves gave a yield of 13.33 %. This rate was close to that reported by Okwu [32] which was 10.06 %. The phytochemical screening of *Alchornea cordifolia* leaves revealed the presence of several secondary metabolites such as phenolic compounds, flavonoids, sterols and terpenoids, quinoid compounds, tannins and alkaloids. These results are in agreement with those of several authors who observed the presence of these compounds in the leaves extract of this plant [24, 29, 33]. Studies also reported that flavonoids and polyphenols respectively account for 14.4% and 42.2 % of dry matter of *Alchornea cordifolia* leaves [24].

Body Weight Evolution

In order to evaluate the efficacy of *Alchornea cordifolia* against aflatoxicosis, we prepared increasing doses of 50, 100 and 300 mg/kg of body weight of the ethanolic leaves extract of this plant. The choice of these doses were carried out from preliminary studies. The BHA is a synthetic antioxidant authorized as food additive. We used it as a standard antioxidant in this study at a dose of 50 mg/kg according to the studies of Shatat [37]. Rats treated with AFB1 at a dose of 150 µg/kg for three weeks showed a weak increase in animals body weight (10.5 g) compared to the initial weight at D0. This growth though positive, is significantly lower than that of the animals of the control group which received the olive oil. This variation of body weight is due to the toxicity of aflatoxin B1 administered at a dose of 150 mg/kg. Groups 3, 4 and 5 which received the mixture containing aflatoxin at a dose of 150 mg/kg and the extract at doses ranging from 50 to 300 mg/kg also showed a positive body weight gain. The recorded body weight gain for these three groups vary from 15.91 g to 45.16 g. These body weight gain are dose-dependent. Therefore, the mixture having the highest concentration of EELac (300mg/ kg) showed a body weight gain of 45.16 g which is significantly different from the body weight gain of group 2 (treated only with aflatoxin B1) but close to the values of control groups (group 1 and 6). Thus, by increasing the extract concentration in EELac in the mixture, we gradually restore animals’ normal growth. These results are similar to those of several authors who reported that the acute or subacute forms of aflatoxicosis are dominated by anorexia, depression and body weight loss [15, 16]. These observed effects are due to the inhibition of proteins synthesis and DNA. Several authors reported
Figure 1. Effect of AFB1 on liver tissues
A photomicrograph of liver section from (A) Liver of rat treated with AFB1 alone (group 2) showing a sinusoid capillary dilation with hemorrhagic contents (arrow), (B) rat treated with AFB1 + EELac at 300 mg/kg (group 5) showed the normal hepatocytes (arrow) and portal area (star) without histological abnormalities, (C) rat treated with AFB1 + BHA at 50 mg/kg (group 6) showed the normal hepatocytes (arrow) and portal area (star) without histological abnormalities.

Figure 2. Effect of AFB1 on kidney tissues
A photomicrograph of kidney section from (D) kidney of rat treated with AFB1 alone (group 2) showed an atrophy of renal tubules (arrow), (B) Rat Treated with AFB1 + EELac at 300 mg/kg (group 5) showed a normal histology of renal tubules (arrow), (C) Rats Treated with AFB1 + BHA at 50 mg/kg (group 6) showed a normal histology of renal tubules and renal glomeruli (arrow).
that body weight decrease following an intake of AFB1 is linked to a reduction of food consumption causing a protein catabolism [38].

**Effect of AFB1 on Proteins, Hepatic Enzymes and renal Function in the Presence of EELac**

Aflatoxins are hepatotoxic for all vertebrates and induce a degeneration, hepatocytes necrosis and modify the liver function [35]. Administration of AFB1 at a dose of 150µg/kg for three weeks caused a significant increase of ALT and AST rates. This significant rise of transaminases rate is caused by the toxic effects of AFB1 in the hepatic tissue and the biliary system [1]. These results are in agreement with those reported in literature indicating that, liver is the principal target of aflatoxins [2, 9]. The results of the biochemical analyses are in accordance with the histological examinations which showed lesions in the hepatic tissues. The rise in the rate of transaminases is due to the sinusoid capillary dilation with hemorrhagic contents revealed by liver tissues through histological examinations. These cytoplasmic enzymes were released in plasma after the lesions of hepatic tissue lesions. Similar effects on liver tissues were reported by several studies in animals receiving aflatoxin B1 [2, 3, 19]. The administration of AFB1 caused a rise in the rates of urea and creatinine. The rise in the rate of these products is a sign of renal dysfunction due to an atrophy of renal tubes and glomeruli. AFB1 also caused a reduction of albumin and total proteins rates in serum, this is due to the reduction of proteins synthesis in the liver and to the production of RNA [1]. These results are in agreement with those reported in literature and clearly showed the toxic effects of AFB1 on proteins and on hepatic and renal tissues [1]. The rates of transaminases, urea, creatinine, albumin and total proteins gradually came to normalcy by the administration of a mixture of AFB1 and the extract of *Alchornea cordifolia*. These results are corroborated by those of Mohammed [26] who reported that *Alchornea cordifolia* acts on liver by restoring the rate in serum of ASAT and ALAT in diabetic rats. The effects of EEFac at a dose of 300 mg/kg are similar to those of BHA at a dose of 50 mg/kg. EEFac could protect the body organs and tissues against AFB1 toxicity following the same mechanism as BHA. Indeed, BHA is a standard phenolic antioxidant which acts effectively against the cancerogenicity of AFB1 by impeding its oxidative way of biotransformation in the liver [6]. Similar effects of EEFac to those of BHA could be essentially due to its phenolic compounds and the flavonoids revealed by the chemical screening.

**Conclusion**

Aflatoxins are a real issue of public health for African countries. The implication of this mycotoxin in the appearance of many pathologies such as hepatocellular carcinoma requires urgent and adequate prevention measurements. The pharmacological effect of the ethanolic leaves extract of *Alchornea cordifolia* in the reduction of aflatoxicosis was investigated. This study showed that, this plant has a protective effect on some body organs and tissues such as the liver and the kidneys against the toxicity of aflatoxin B1. These pharmacological properties of *Alchornea cordifolia* are due to flavonoids and to the phenolic compounds. The effect of this plant which is dose-dependent can be used in the prevention of some pathologies linked to the oxidative stress. In addition deep studies should be undertaken in order to evaluate the efficacy of this plant in the protection of other body organs on which aflatoxins have a toxic effect.

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