Impact of repeated administration of Cannabis sativa on some biochemical parameters in albino rats

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Abstract: Cannabis sativa is an illicit drug with proven anti-inflammatory, analgesic, antipyretic and anti-diarrhoea activities. This present study is aimed at ascertaining the impact of Cannabis sativa on some biochemical indices of albino rats. Twenty (20) male wistar albino rats ranging from 180-200g were acclimatized to laboratory conditions for 7 days, following which they were randomly assigned into 4 groups i, ii, iii and iv of five animals each based on average body weight. Groups (ii-iv) were administered 0.5 ml aqueous extract of Cannabis sativa via oral route corresponding to 1, 2 and 3 mg/kg/body weight, while group i (control) received 0.5 ml of distilled water orally. Rats in all groups were sacrificed 24 hours after the experimental periods of 30 days after administration of the extract. The extract significantly (p<0.05) increased serum Albumin, Globulin, Total protein, Conjugated bilirubin while Total bilirubin, unconjugated bilirubin concentration showed a significant (p<0.05) increase only in group ii and iv when compared with the control. Likewise group ii and iii showed a significant (p<0.05) reduction in total cholesterol. The serum AST and ALT showed a significant (p<0.05) increase with the control. The alterations by the results are manifestation of mild hepatorenal toxicity and anti-hypercholesterolemic effect.

Keywords: Analgesic, anti-hypercholesterolemic, anti-inflammatory, Cannabis sativa hepatorenal,

I. Introduction
Cannabis sativa is a dioecious (having male and female flowers in separate plants), green, leafy plant with characteristic opposite, usually seven-fingered, lance-shaped leaves; on dry, sandy, slightly alkaline soil it can grow to more than seven meters in height. Glandular hairs develop, usually on the female flower, which secrete a resin. The female plants are more important than the male plants for commercial purposes: their fibres are thicker, they form the nutritious seeds, and they contain the psychoactive principle tetrahydrocannabinol (THC) [1].

Over the years, Cannabis remains the most widely used illicit drug worldwide due to its affordability and availability [2]. Besides, cannabis is a major controversial drug as there are numerous conflicting and controversial reports concerning its psychological and physiological effects [3]. Many reports have linked cannabis smoking to the development of psychosis [4]. Certain studies have also suggested that cannabis smoking is only a form of self-medication in people with psychotic symptoms rather than a causative factor in development of psychosis [5]. Recently, there seems to be an increase in the number of reports lending support to cannabis psychosis. Further, a review of the evidence surrounding the acute impact on memory concluded that cannabinoids impair all aspects of short-term memory, especially short-term episodic and working memory [6]. Apart from the psychological effects, various reports have shown that cannabis smoking has significant physical effects on the body [7,8]. Cannabis has been found to increase heart rate by 20-50%. This is the most immediate effect and occurs within a few minutes after cannabis intake. After cannabis usage, a sudden change of posture from lying down to standing up may produce orthostatic hypotension, a feeling of ‘light-headedness’ and faintness that is often the earliest indication of intoxication in naive users [9]. Other physical effects of cannabis include reddening of the eyes due to congestion of the conjunctiva blood vessels, lowering of the body temperature, dry mouth, reduced intraocular pressure and relaxation of the muscles [10]. Cannabinoids have also been shown to affect the immune system [11]. Clinical studies and survey data have found that cannabis increases appetite [12]. The Δ⁴-tetrahydrocannabinol (THC) which is the main constituent of cannabis has been shown to have effect on both the action and the release of insulin [13]. This may explain why cannabis has been employed to self-medicate in diabetes. Cannabis has also been reported to have medicinal use in treating depression [14]. Cannabis sativa have also been reported to be used for treatment of specific human ailments such as allergies, burns, cuts and wounds, inflammation, leprosy, leucoderma, scabies, smallpox and sexually
transmitted diseases [15]. The different preparation of *Cannabis sativa* has been used in Asian traditional medicine for treatment of variety of diseases including: inflammation, nausea, headache, hematochesia, diarrhea, and alopecia [16,17,18] have demonstrated a potent anti-inflammatory, analgesic, antipyretic and antidiarrhea activities.

In the developing countries, like Nigeria, data available is scarce as well as reliable information of actual and potential health consequences, which could give input into health analysis of national developmental strategies. Actual evidence of chronic health effect of cannabis sativa can be provided in the laboratory using experimental animals models in which well controlled doses are administered over a period of time. Therefore this research is focused on the impact of cannabis sativa on some biochemical parameters in albino rats.

II. Materials And Methods

SAMPLES

The plant materials was collected from a local farm in Akpet and authenticated at the Botanical unit of the Biological Science Department, University of Calabar, Nigeria. The crude extract was prepared according to the method described by [19]. The leaves were oven dried at 45°C and hundred grams of the dried fruit materials was pulverized and soaked for 72 hours in 800mls of distilled water. It was then filtered with Whatman No 1 filter paper and the residue discarded. The filtrate was subsequently evaporated to dryness in an aerated oven at 45°C. The resulting slurry was stored in closed cab bottles until used.

Animals and treatment protocols:

Twenty (20) wistar albino rats weighing 180-200g were used for this work. The animals were obtained from the animal holding unit of the Department of Human Physiology, University of Calabar, Calabar, Cross River State. The animals were housed in plastic cages and were allowed acclimatization period of Seven (7) days in a well-ventilated room with a temperature and pressure of 29±2 °C and70% respectively. The wistar albino rats were maintained with rat chow (vital feeds Ltd) and water *ad libitum*. The animals were exposed to 12hours light -dark cycle and handled according to standard protocols. At the end of acclimatization period, they were divided into four groups i, ii , iii and iv of five (5) rats each. Group i served as control while ii - iv were the test group .The control group was treated with 0.5mls of normal saline while B and C were treated with 0.5 mls corresponding to 1, 2, and3 mg/kg body weight of the drug(*Cannabis sativa*) respectively and were administered orally for Thirty days (30) days. The wistar albino rats were sacrificed 24 hours after the last administration in accordance with the guide lines of the European Convention for the protection of vertebrate animals and other scientific purposes-ETS-123[20].

Preparation of Serum.

The animals were anaesthetized in a jar containing cotton wool soaked in diethyl-ether and chloroform in ratio:1:1. When the animal became unconscious, they were brought out quickly of the jar, the abdominal region was opened along the linear Alba and diaphragm cut with scalpel blade to expose the organs and blood was collected into a sterile sample container by cardiac puncture. Blood was collected into a clean, dry centrifuge tube and allowed to clot for 30 min before centrifuging at 300rpm x 10min using Uniscope Laboratory Centrifuge. The serum was thereafter aspirated into clean, dry, sample bottles using Pasteur pipette and was kept or stored in sample bottles and used within 12hours of preparation as described by [21]. Later it was transferred into specimen bottles before being used for biochemical analysis.

Statistical Analysis

Statistically analysed data used was presented as mean ± SD of five (5) determinations. Statement analysis was carved out using one way analysis of variance (ANOVA). Differences were statistically significant at P<0.05 [22].

III. Results

The results depict the effect of *Cannabis sativa* on some biochemical parameters indices of Wistar albino rats. Fig 1-2 showed the effect of *Cannabis sativa* on serum AST and ALT of Wistar albino rats, though, group ii showed no significant difference when compared with the control but both group iii and iv showed a significant (P<0.05) increase when compared with the control. More so, the total bilirubin in group ii and group iv showed a significant (P<0.05) increase when compared with the control. While group iii was significantly (P<0.05) reduced when compared with the control (Fig 3).The conjugate bilirubin showed a significant (P<0.05) decrease in group ii when compared with the control while group iii and iv showed a significant (P<0.05) increase when compared with the control (Fig 4). The effect of *Cannabis sativa* on serum unconjugated bilirubin on Wistar albino rats showed a significant (P<0.05) increase in group ii and iii but significant (P<0.05) decrease

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when compared with control (Fig 5). The drug also reveals a significant (P<0.05) increase when compared with the control in animals in group ii but a significant (P<0.05) decrease following the administration of the drug in both group iii and iv respectively(Fig vi). On the other hand, Total protein showed a significant increase in all the experimental groups as seen in (Fig 7). Similar patterns was also shown in albumin and globulin(Fig 8-9). Fig 10 showed a significant decrease in albumin /globulin ratio in all the experimental groups.

Fig 1: Effect of Cannabis sativa on Serum AST of Wistar Albino Rats

Fig 2: Effect of Cannabis sativa on Serum ALT of Wistar Albino Rats
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Fig 3: Effect of Cannabis sativa on Serum Total Bilirubin of Wistar Albino Rats

Fig 4: Effect of Cannabis sativa on Serum Conjugated Bilirubin of Wistar Albino Rats

Fig 5: Effect of Cannabis sativa on Serum Unconjugated Bilirubin of Wistar Albino Rats
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**Fig 6:** Effect of *Canabis sativa* on Serum Total Cholesterol of Wistar Albino Rats

**Fig 7:** Effect of *Canabis sativa* on Serum Total Protein of Wistar Albino Rats

**Fig 8:** Effect of *Canabis sativa* on Serum Albumin of Wistar Albino Rats
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IV. Discussion

The Liver has an incontrovertible influence on several functions of many organs in the body. It is prone to xenobiotic induced injury due to its central role in xenobiotic metabolism and its portal location within the system [23]. The liver plays important role in metabolism, detoxification and biotransformation. Therefore, an alteration in the biomarkers of the liver function indices might be used to monitor the level of injury or damage by the plant extract before biopsy [24].

The observed significant (p<0.05) increase the Albumin, Globulin, Total protein and Total bilirubin concentration implied that the extract produced an increase in protein synthesis and (or) mobilization. Moreover, the observed increase in Globulin level may indicate the efficiency of the plant extract to produce antibody [25] or due to the presence of bioactive constituent like flavonoids and alkaloids.

Albumin is the protein with the highest concentration in the plasma. It transports many molecules in the blood. It prevents the fluid in the blood from leaking out the tissue [26]. Albumin is a constituent of the total protein produced in the liver. Albumin levels are decreased in chronic liver disease such as cirrhosis or nephrotic syndrome. Therefore, the observed increase in serum albumin is an indication that the Cannabis sativa may promote good functioning of the liver or might possess a hepatoprotective role and may help calcium in the blood stream to regulate the movement of water blood stream into body tissue [27]. This is also supported by the result displayed by the albumin /globulin ratio and significant increase seen in total protein concentration. However, the significant increase in the serum level of both total bilirubins, unconjugated and conjugated.
biliburin is an indication that the drug might induce injury to the hepatic tissue or caused conjugated hepatobiliary injury on the wistar albino rats [28].

Serum Alanine Amino Transferase (ALT) is known to increase when there is liver disease and it has been used as a tool for measuring hepatic necrosis [29,26]. Hence, the observed increase in serum ALT suggests that extract may not be safe to the hepatic tissue at 1, 2 and 3 mg/kg body weight. Aspartate Amino Transaminase (AST) is predominantly localized within the cells of the gills, kidney, muscle and liver parenchyma cells. The observed increase in serum AST might connote acute liver damage or liver cytolsis. ALP is a marker enzyme for the plasma membrane and endoplasmic reticulum [30] or cell lining of the biliary ducts of the liver, placental tissue and bone. The significant decrease in total cholesterol in group iii and iv following the administration of the drug suggest that the drug exhibits an antilipidaemic or antihypercholesterolemic or cholesterol lowering effect.

V. Conclusion

It may therefore be too hasty to conclude that the drug was responsible for liver or hepatorenal toxicity as observed in the serum enzymes. More studies are highly imperative so as to be able to understand fully the various effects and mechanisms of cannabis on smokers. Though, it appears that the drug can enhanced protein synthesis, mobilization and antihypercholesterolemic or cholesterol lowering effect. Further studies can also reveal other possible pharmacologically active substance(s) that may have a good medicinal value, present in the drug.

References


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