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Hepatoprotective Activity of Extract of *Erythroxylum monogynum* in Albino Rats

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ABSTRACT

Development in the science is helping the people to have the knowledge of unknown things. In that attempt, finding the plants as the best therapeutic option for the health problems and making them familiar to the people. The present experiment was conducted for 7 days to evaluate the hepatoprotective activity of Erythroxylum monogynum in CCl₄ (1ml/kg) induced rats. The hydroalcoholic extract of plant was prepared by maceration technique. The 5 group of rats were maintained as Control, CCl₄ induced, CCl₄+ Liver tonic, CCl₄+extract 150mg/kg and CCl₄+extract 200mg/kg. On the 8th day blood was collected by retro orbital puncture for the study of serum parameters like SGOT (Serum Glutamate Oxaloacetate Transaminase), SGPT (Serum Glutamate Pyruvate Transaminase) and bilirubin. Then the liver separated and processed for the histological studies. The decreased levels of SGOT, SGPT, total bilirubin in the treated rats were an indication of the hepatoprotective activity of extract. The regeneration of hepatocytes was also evidence for hepatoprotective activity of extracts.

Keywords: Hepatoprotective activity, SGOT, SGPT, Liver tonic, Necrosis.

INTRODUCTION

Though the Science is enormously developed certain diseases and syndromes are not controlled. Some of the diseases like cancer, HIV, malaria etc. are still in the mind of Scientists to which they are unable to find out solution to cure completely. Though, the development is enormous in various fields the poor people unable to get the treatment facility. That's why the people preferring the local doctors. Sometimes the treatment of those practitioners may workout or not. The ancient practitioners for medicine depending on the plants can solve the various diseases. Among them there are several plants that have the capacity to combat against the liver diseases.

Liver is vital organ, which is maintaining of metabolic reactions in the human body. But unnecessary food habits, consuming of impure drinks may bring problems in functioning of the liver. Consuming more number of drugs also cause the liver damage, intake of alcohols, junk food etc are also questioning the functional ability of the liver by damaging the liver architecture. Damage of liver can be assessed by the elevated levels of serum enzymes like SGOT, SGPT and bilirubin¹. Though the allopathic drugs are available they are not fulfilling the solution. But the medicinally important plants are capable to target these types of problems. In that attempt many plants are proved that they possess the hepatoprotective activity². Plants having the capability of fighting against free radicals generated in the body leads to protect of vital organs like liver.

The present work is to find out the free radical scavenging activity and liver protection activity of leaf extracts of *Erythroxylum monogynum* against CCl₄ induced hepatic damage.

MATERIALS AND METHODS

Plant material

Erythroxylum monogynum plants were brought from the forest area of the Gudur Village, Warangal district. The plant was identified and authenticated by Prof. V.S. Raju, Department of Botany, Kakatiya

University, Warangal. The plants materials are generally practiced by the village tribal people for various ailments. They are getting relief of inflammations, skin diseases, malaria, and stomach ache³.

Preparation of plant extract

The leaves were dried under shade and the powder was prepared from dried leaves. 50g of powder was mixed in 250 ml of hydroalcoholic solvent (70% of ethanol, 30% of distilled water) and allowed for 24 hrs with the random shaking. Then the filtrate-I was collected and the marc dissolved in 250 ml of hydroalcoholic solvent for 24 hrs and collected the filtrate-II. Then the filtrates (I&II) were distilled to get extracts and stored in refrigerator prior to treatment.

Animal models

Albino rats (Wistar strain) weighing 150 to 200gr were brought from Mahaveer Enterprizers, Hyderabad. The rats were housed and acclimatized to standard laboratory conditions (25° C, 50-60 % of humidity) with the approved protocol of Institutional Animal Ethical Committee (IAEC/03/ UCPS/KU/10). The animals were fed with standard diet (Hypro feed for animals, Pune) and water ad libitum.

Toxic study of the extracts

To study the toxicity of extracts the doses 150,200,250,300 mg/ kg of extracts were administered to the rats (4 groups – 6 animals in each group) and put under observation for 7 days⁴. There was no toxic effect observed to the rats up to 300 mg/kg and the 150 and 200 mg / kg were selected for the experiment.

Experimental design for hepatoprotective activity

The animals were divided into 5 groups of 6 in each

Group-1: Control- Treated with dist. water for 7 days

Group-2: CCl₄ (Carbon tetra chloride) was given (i.p., 1ml/ kg) with 1:1 dilution of coconut oil on the 5th day⁵.

Group-3: Administered with liver tonic (LIV – 52 Syrup, with the dose 5ml/kg) daily for 7 days and on 5th day the CCl₄ (1 ml / kg) was given through i.p.

Group - 4: Treated with extract 150mg/ kg for 7 days and the CCl₄ induced on 5th day⁵.

Group -5: Treated with extract 200 mg/kg for 7 days and the CCl₄ induced on 5th day⁵.

On the 8th day, blood samples were collected through plexus from all rats and they were sacrificed. The collected blood was centrifuged and separated serum samples were used for estimation of SGOT, SGPT and bilirubin (through commercially available kits) tests for the study of the toxic effect of CCl₄ and also the therapeutic effect of the plant extracts. The livers were fixed in the fixative (Bouin's fluid) and processed further for histological studies. The results were analyzed by one way ANOVA followed by Dunnet multiple comparison test with the significant level at p<0.05.

RESULTS

The results were observed that the serum parameters like SGPT values were increased in the CCl₄ induced rats. The elevated SGOT, bilirubin values were also indicating the damage of the liver in the CCl₄ induced rats. The values of SGOT, SGPT and bilirubin were significantly decreased in the group CCl₄ + Liver tonic. The decreased levels of SGPT, SGOT and bilirubin levels were also seen in the CCl₄+150mg/kg and CCl₄+200mg/kg extract treated groups.

DISCUSSION

CCl₄ damages the liver by its metabolite CCl₃· free radical, with which the damage of cellular membranes occur through the lipid peroxidation⁶. The serum parameters like Serum Glutamate Oxaloacetate Transaminase (SGOT) or Aspartate aminotransferase (AST), Serum Glutamate Pyruvate Transaminase (SGPT) or Alaline aminotransferase (ALT), including the bilirubin content also elevated because of their release into the blood in the CCl₄ induced hepatotoxic rats⁷ (Table-1, Chart- 1). Whereas, the extract treated rats' serum parameters revealed the significant decrease in the SGOT, SGPT and bilirubin levels compare to the CCl₄ induced rats (Table-1, Chart - 1 and 2). These enzymatic values were also decreased in the liver tonic treated rats. The hepatoprotection of the drug depends on the reduced effects of toxic levels of the CCl₄ in the damaged liver⁸. The results that decreased levels of SGOT, SGPT and bilirubin

in the extract treated rats against CCl₄ were observed similar to the results of the hepatoprotective activity of poly herbal drug against CCl₄ damaged liver^{9,10}.

The histological sections were also revealed that necrosis of hepatocytes occurred in the CCl₄ induced hepatotoxic group (group- 2) (figure-2). The group – 4 and group - 5 were showed the rearrangement of damaged cells. The reformation of hepatocytes was observed in histology of group – 5 (figure- 4, 5) rats. The histology can be easily comparable with the CCl₄+ liver tonic group rats (figure- 3).

The dose 200mg/kg showed that serum enzymes were decreased more than to the dose of 150mg/kg in the extracts treated rats. The hepatoprotective activity is increased according to the dose.

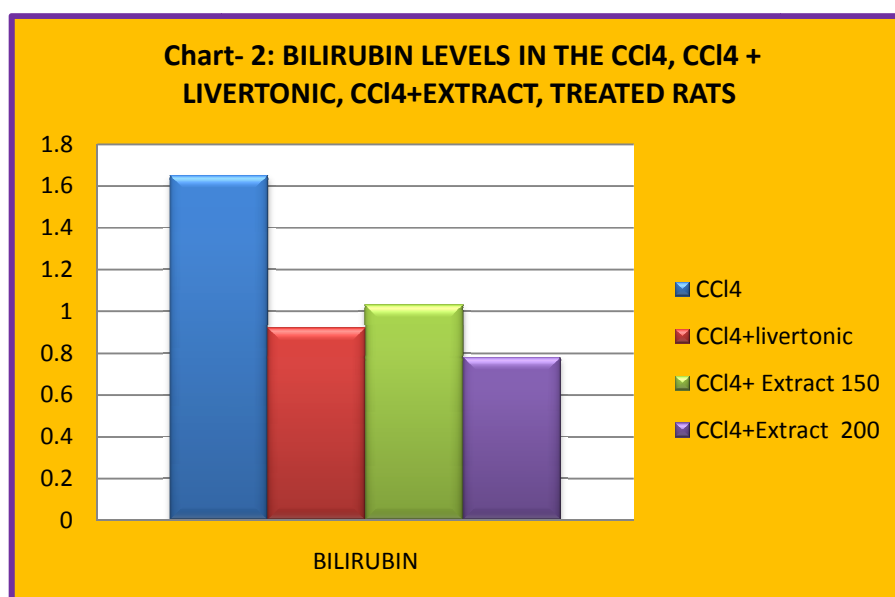
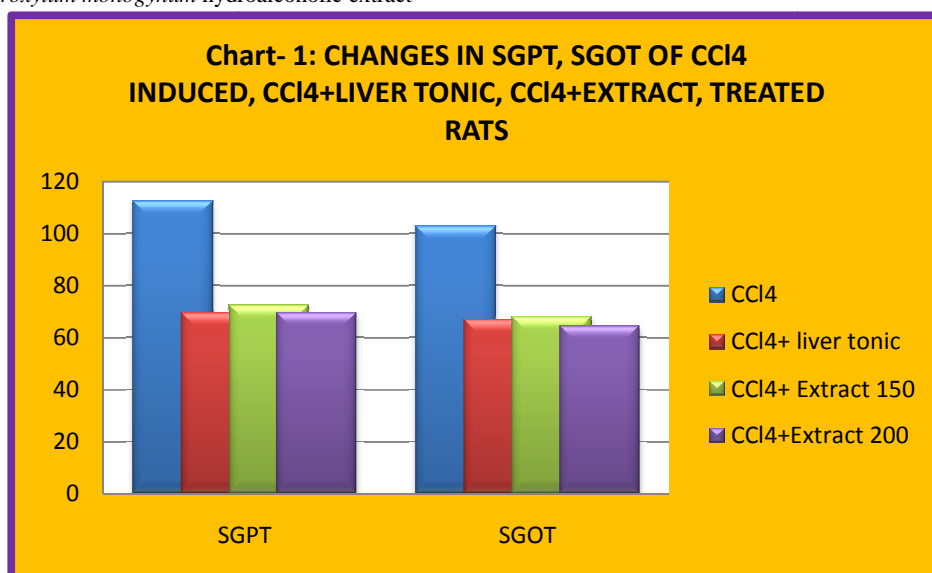
Table – 1: Hepatoprotective Activity of *Erythroxylum monogynum* in CCl₄ induced albino rats

GROUPS	SGPT (IU/L)	SGOT (IU/L)	BILIRUBIN (mg/dL)
1.CONTROL	52.73±0.73	42.66 ± 0.55	0.64±0.08
2.CCl ₄	112.60±4.51	102.88±1.79	1.65±0.15
3.CCl ₄ + Liver Tonic	69.54±1.63	66.99±0.86	0.92±0.14
4.CCl ₄ + Extract 150 mg	72.48±1.72 ^a	68.03±1.23 ^a	1.03±0.16 ^a
5.CCl ₄ + Extract 200 mg	69.56±3.90 ^a	64.49±2.30 ^a	0.78±0.06 ^a

All values are expressed in mean ± SD; n=6

a= p <0.01 compare to CCl₄ induced group,

Extract- *Erythroxylum monogynum* hydroalcoholic extract



Histological sections of liver

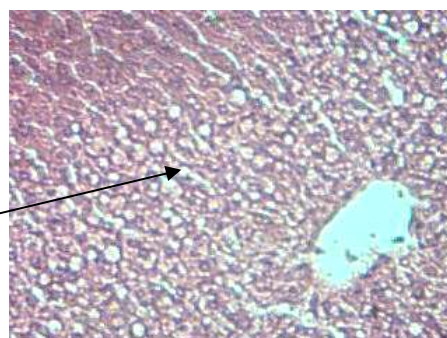
Fig. 1



Normal cells

Liver of control rat showing normal histology

Fig. 2



Damaged cells

Histology of liver with CCl₄ damage

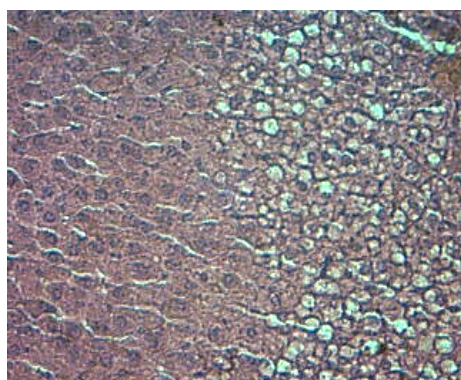
Fig. 3



Cells recovering to normal stage

Histology of liver treated with CCl₄ + liver tonic

Fig. 4



Histology of liver treated with CCl₄+Extract 150mg/kg

Fig. 5



Histology of liver treated with CCl₄ + Extract 200mg/kg

REFERENCES

1. Mascolo, N. Sharma, R. Jain, S. C., and Capasso, F., Ethnopharmacology of Calotropis Procera flowers. *J Ethnopharmacol.*, **22(2)**: 211-221(1998)
2. Chowdhary, G.D. Kamboj, P. Singh, I. and Kaila, A.N., Herbs as liver savers – A review, *Indian J. of Natural Products and Resources.*, **14**: 397-408(2010)
3. Venkatesalu, V. Gopalan, N. Pillai, C.R. Singh, V. Chandrasekaran, M. Senthilkumar, A. and Chandramouli, N., *In vitro* anti-plasmodial activity of some traditionally used medicinal plants against Plasmodium falciparum, *Parasitol Res.*, **111(1)**: 497-501(2012)
4. Kavitha, B.T. Shruthi, S.D. Padmalatha, R.S. and Ramachandra, Y.L., Phytochemical analysis and hepatoprotective properties of Tinospora cordifolia against carbon tetrachloride-induced hepatic damage in rats, *J of Basic and Clinical Pharmacy.*, **2(3)**: 139-142 (2011)
5. Chaudhari B. P., Chaware V. J., Joshi Y. R. and Biyani K. R., Hepatoprotective activity of hydroalcoholic extract of Momordica charantia Linn. leaves against Carbon tetra chloride induced hepatopathy in rats., *Intern J of Chem Tech Research.*, **2**: 355-358 (2009)
6. Brattin W.J., Glend F.A.J. and Recknagel R.O., Pathological mechanism in carbon tetrachloride hepatotoxicity, *J Free Radical Boil Med.*, **1**: 27-28(1985)
7. Durga, P.N. Dinda, S.C. Swain, P.K., Kar, B. and Patro, V.J., Hepatoprotective activity against CCl₄-induced hepatotoxicity in rats of Chenopodium album aerial parts, *J of Phytotherapy and Pharmacology.*, **2**: 33-41 (2012)
8. Siddhatha, S. Archana, M. and Pradeep, M., Hepatoprotective activity of Cajanus cajan against CCl₄ induced liver damage, *Inter J of Pharm. and Pharmaceu. Sciences.*, **3(2)**: 146-147 (2011)
9. Vilas, A.A. Ganjiwale, R.O. and Yeole, P.G., Phytochemical and pharmacological standardization of polyherbal tablets for hepatoprotective activity against carbon tetrachloride induced hepatotoxicity, *Inter J of Pharmaceu. Sciences and Drug Research.*, **2(4)**: 265-268 (2010)
10. Rajesh, K. Sushil, K. Arjun, P. and Jayalakshmi, S., Hepatoprotective activity of aerial parts of Plumbago zeylanica linn, against CCl₄ induced hepatotoxicity in rats, *Inter J of pharmacy, Pharmaceu Sciences.*, **1(1)**: 171-175(2009)
11. Jeong, H.G. You, H.J. Park, S.J. Moon, A. Chung, Y.C. Kang, S.K. and Chun, H.K., Hepatoprotective effects of 18 β -glycerrheinic acid on carbon tetra chloride induced Liver injury: Inhibition of cytochrome P450 2E1 expression, *Pharmacological Res.*, **46**: 221-227 (2002)
12. Prabhat, K.D. Prasanna, P. Somya, R.P. and Ranjan, S., Hepatoprotective activity of Plant Argemone mexicana (Linn) against CCl₄ induced hepatotoxicity in rats, *Inter J of Pharmaceu Research and Development.*, **8**: 1-20 (2009)
13. Satheesh, B. Sathyanarayana, J. Krishna Reddy, M. and Prasad, M.S.K., Evaluation of hepatoprotective activity of aqueous extract from leaves of Solanum americanum. *Inter J of Plant Animal and Environm Sciences.*, **1(2)**: 150-154 (2011)
14. Zahara, K. Malik, M.A. Mughal, M.S. Arshad, M. and Sohail, M.I., Hepatoprotective role of extracts of Momordica charantia.L. in Acetaminophan – induced toxicity in rabbits. *The J Animal and Plant Sciences.*, **22(2)**: 273-277(2012)