

Hepatoprotective activity of *Punica granatum* leaf extract against Carbon Tetrachloride induced Hepatotoxicity in Rats

Manoj Kumar^{1,*}, Sukumar Dandapat¹ and Manoranjan Prasad Sinha¹

Department of Zoology, Ranchi University, Ranchi – 834008, Jharkhand (India)

*Correspondence address: Department of Zoology, St. Xavier's College, Ranchi – 834001, Jharkhand (India)



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Corresponding author: Manoj Kumar

E-mail address: dr17mk@gmail.com

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Florina Ojoga, National Institute of Rehabilitation, Physical Medicine and Balneology, "Carol Davila" University of Medicine and Pharmacy

Abstract

In the present study Hepatoprotective activity of aqueous leaf extract of *Punica granatum* on total protein, bilirubin, aspartate aminotransferase (AST), alanine transaminase (ALT) and alanine phosphatase (ALP) in CCl₄ intoxicated rat were studied. Administration of CCl₄ showed significant increase ($p < 0.01$) in liver marker enzymes in serum namely AST (52.30 ± 1.15 to 107.2 ± 13.7 IU/L), ALT (146.63 ± 5.79 to 206.2 ± 28.82 IU/L), ALP (176.24 ± 5.8 to 508.2 ± 10.22 IU/L), bilirubin (0.52 ± 0.09 to 2.61 ± 0.27 mg/dl) and significantly decreased total protein (9.57 ± 0.17 to 5.2 ± 0.085 mg/dl), when compared to normal. Aqueous extract of *Punica granatum* at 250 mg/kg and 500 mg/kg body weight showed significant increase in total protein (5.2 ± 0.085 to 6.1 ± 0.023 ; 6.5 ± 0.033) as compared to CCl₄ treated rats. The extract lowered enzyme levels which is designation of Hepatoprotective action of extract. The serum AST, ALT and ALP levels are reliable markers of liver function. Thus the present study concludes the aqueous leaf extract of *Punica granatum* to possess Hepatoprotective activity.

Key words: alanine transaminase, alkaline phosphatases, CCl₄, aspartate aminotransferase

INTRODUCTION

Diseases like jaundice, cirrhosis and fatty liver are commonest liver diseases worldwide. In India numerous medicinal plants such as *Adhatoda vasica*, *Psidium guajava*, *Coccinia indica* etc... are employed for treatment of liver related disorders. *Punica granatum* is traditionally used medicinal plant in India. *Punica granatum* has been reported to possess antimicrobial, antioxidant and reducing power ability (Kumar *et al.*, 2015). Free radicals or oxidative injury now appears to be the fundamental cause behind a number of mammalian diseases (Parvathi *et al.*, 2013). The mammals have a complex antioxidant system to combat the oxidative stress. However excessive reactive species derived from oxygen and nitrogen may still lead to oxidative damage to tissue organs. Oxidative stress has been considered as a conjoint pathological mechanism and it contributes to initiation and progression of liver injury (Sha *et al.*, 2015). Various plants are scientifically proven to possess Hepatoprotective activity, and the underlying mechanisms involve the antioxidant property of the plants (Kumar *et al.*, 2013a). Effects of antioxidants or free radical scavengers have been widely tested for the prevention and treatment of acute and chronic liver injuries. In some of the studies, antioxidants have shown beneficial effects, specially for prevention and

treatment of liver injury (Kukongviriyapan *et al.*, 2013; Kumar *et al.*, 2014a). The present study was undertaken to investigate the Hepatoprotective activity of aqueous leaf extract *Punica granatum* leaf extracts in CCl₄ induced Hepatotoxicity in rats.

Materials and Methods:

Plant materials: The fresh tender leaves of *Punica granatum* was collected. The leaves were washed with deionised water and disinfected with 0.1% HgCl₂ solution for 5 min and dried in shade away from direct light for 20 days and ground to fine powder using electrical grinder. The powder obtained was sieved and stored in air tight containers for future use (Kumar *et al.*, 2014b).

Preparation of leaf extracts: the fine powder of *Punica granatum* was made into thimble for loading in Soxhlet apparatus and extraction was done using distilled water. The extraction was continuously done for 72 hours. The extracts thus obtained were concentrated in vacuum rotary evaporator and extracts were kept in dessicator until used (Kumar *et al.*, 2014b; Dandapat *et al.*, 2014c)

Phytochemical screening: preliminary phytochemical screening were conducted on *Punica*

granatum in accordance to previously published standards (Kumar *et al.*, 2014b).

Antioxidant properties: The antioxidant properties of plant sample was determined by Spectrophotometric quantitation method (Prieto and pineda, 1999; Ferreira *et al.*, 2007; Ghosh *et al.*, 1984). Various concentrations of samples (5 µg, 50 µg, 100 µg) were taken in a series of test tubes. Then 1.9mL of reagent solution (0.6m Sulphuric acid, 28mm Sodiumphosphate and 4mm Ammonium molybdate) was added to the test tubes. The tubes were incubated at 95°C for 90 min and allowed to cool down. The absorbance of aqueous solution of each was measured at 695nm against blank. Antioxidant capacity was expressed as equivalents of ascorbic acid. Butylated hydroxyl anisole (BHA) was used as reference standard.

Animals: Albino rats weighing about 175-200 g were used in the study. They were maintained under standard laboratory conditions at ambient temperature of $25 \pm 2^\circ\text{C}$ and relative humidity at $50 \pm 15\%$, with dark-light cycle of 12h. Animals were fed with a commercial pellet diet and water ad libitum. The experiment was performed after prior approval of Ethics committee of Ranchi University, Ranchi (Proceeding no. 46, page no. 137).

Acute Toxicity Study: the acute toxicity studies was carried out as per stair case method [20]. 50 albino mice of either sex weighing 20-25g and 90 days were used to determine LD50 of various extracts. The mice were divided into 3 groups of 10 mice each as follows for leaf extract of *Punica granatum*.

Group 1: Received 1 ml of distilled water orally.

Group 2: Received 250 mg/kg body weight of extract orally.

Group 3: Received 500 mg/kg body weight of extract orally.

No mortality was observed up to 500 mg/Kg body weight of leaf extract.

Assessment of Hepatoprotective Activity: All the animals were sacrificed on day 14 under light ether anesthesia. 5 ml of blood was collected from each animal by cardiac puncture using sterile needle and syringe. Part of blood sample was put into test tubes

and allowed to clot for 30 min at 37°C . The clear serum was separated at 2500 rpm for 10 min and biochemical investigations were carried out to assess liver functions *viz.* Total bilirubin (Kingsley and Frankel, 1939), total protein (Kingsley and Frankel, 1939), serum transaminases (Reitman and Frankel, 1957) and alkaline phosphatase (Bessey *et al.*, 1964).

Results and Discussion:

The results of phytochemical screening of aqueous leaf extract of *Punica granatum* is presented as figure 1. Aqueous leaf extract of *Punica granatum* showed the presence of alkaloids, flavonoids, saponins, tannins and phenolic compounds. The results revealed that flavonoids are present in highest concentration (81.16 ± 1.099 mg/ml) and alkaloids are present in lowest concentration (3.075 ± 1.00 mg/ml). The results of antioxidant activity of aqueous leaf extract of *Punica granatum* is presented as figure 2. The leaf extract of *Punica granatum* showed strong antioxidant activity as compared to standard BHA (Butylated Hydroxyanisole). The results of different parameters of Liver function tests are presented in figure 3 and figure 4. Figure 3 shows the change in liver function parameters in rats administered with CCl_4 . The CCl_4 has been utilized as an implement to induce Hepatotoxicity in experimental animals (Recnagel, 1983; Okuno *et al.*, 1986). Liver is the main metabolic centre where detoxification and drug metabolism take place which makes it greatly vulnerable to damage by toxic substances (Reddrop *et al.*, 1983). CCl_4 caused peroxidative degradation in adipose tissue resulting in fatty infiltration of hepatocytes. The incrementation in the calibres of serum bilirubin reflected the depth of jaundice and incrementation in serum transaminases and alkaline phosphatases was the clear designation of cellular damage and loss of functional integrity of the cell membrane of hepatocytes (Sarwath *et al.*, 1993). The results showed elevation in all parameters under examination.

Aminotransferases include AST (Aspartate transaminase) and ALT (Alanine transaminase). They participate in gluconeogenesis by catalysing the transfer of amino group from aspartic acid or alanine to ketoglutaric acid to produce oxaloacetic acid and pyruvic acid respectively. AST and ALT are an excellent markers of hepatocellular injury. AST is

present in cytosolic and mitochondrial isoenzymes and is found in the liver, cardiac muscle skeletal muscle, kidneys, brain, pancreas, lungs leucocytes and red cells (Cohen and Kaplan, 2000). It is comparatively less sensitive as compared to ALT. The ALT is a cytosolic enzyme found in highest concentration in the liver, and its test is more specific in determination of liver functional health (Cohen and Kaplan; 2000).

The lowering of enzyme levels is a definite designation of Hepatoprotective action of the drug. The serum AST, ALT and ALP levels are reliable markers of liver function (Sarwath *et al.*, 1993; Cohen and Kaplan, 2000). In our study, the consequential elevation of marker enzymes following administration of CCl₄, betokened earnest toxicity engendered by the chemical. The administration of aqueous leaf extract of *Punica granatum* engendered reduction in AST, ALT and ALP levels and total bilirubin levels. Proteins are the building units of the body and are also the most abundant macromolecules in the cells constituting half of their dry weight and they regulate various physiological and metabolic processes (Jaleel *et al.*, 1996). On administration of CCl₄ the total protein content decreased. Shalaby (2009) attributes the significant hypoproteinaemia to the cellular destruction or necrosis of liver with subsequent impairment of the protein synthesis machineries of liver, which synthesized most of the plasma proteins (Guyton and Hall, 1996). An increase in total protein was observed in leaf extract administered animals (Group 1 and Group 2) which indicates the recovery of the protein synthesis machineries of liver. Various liver protective herbal drugs contain a variety of chemical constituents like carotenoids, flavonoids, alkaloids, reducing sugars and saponins etc. [25] (Gupta and Misra, 2006), and in the present study alkaloid, flavonoid, saponin, tannin were found in the aqueous leaf extract of *Punica granatum* and may be responsible for the Hepatoprotective effect of the extract. Thus it could be suggested that *Punica granatum* aqueous leaf extract possesses Hepatoprotective activity in this model and may be used in abnormalities related to liver.

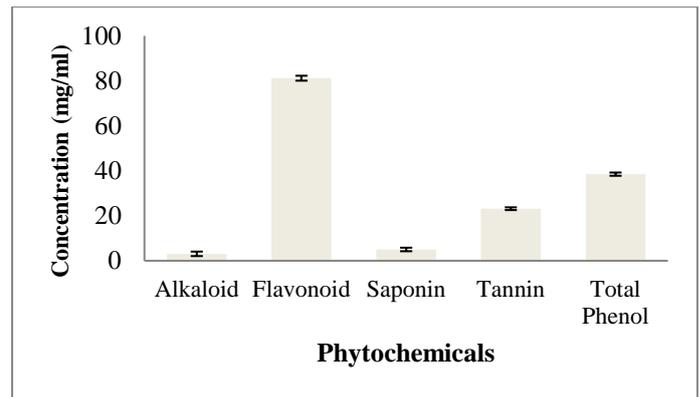


Figure 1: showing results of phytochemical screening of aqueous leaf extract of *Punica granatum*

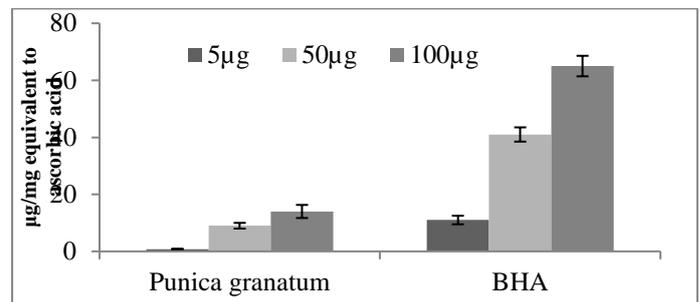


Figure 2: antioxidant activity of aqueous leaf extract of *Punica granatum* as compared to BHA (Butylated hydroxyanisole)

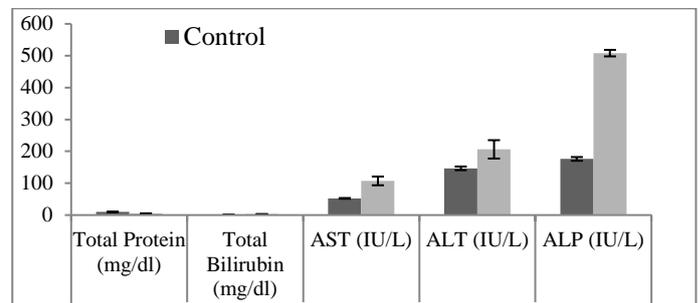


Figure 3: Effects of CCL4 treatment on Liver function parameters of albino rats

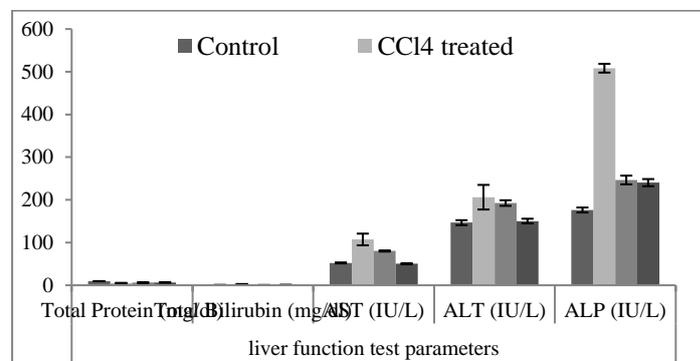


Figure 4: Effects of Punica granatum leaf extracts on liver function parameters of albino rats

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