Hepatoprotective activity of aqueous methanolic extract of *Viola odorata* against paracetamol–induced liver injury in mice
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Abstract

Traditionally Viola odorata is used for liver protection. To provide scientific support to its traditional use, aqueous methanolic extract of V. odorata (250 and 500 mg/kg) was given to mice intoxicated with paracetamol. Obtained results demonstrated that the extract significantly (p<0.01-0.001) reduced paracetamol induced increase levels of serum hepatic enzymes and total bilirubin. Histopathological studies showed that the plant attenuated the hepatocellular necrosis and inflammation. HPLC results showed the presence of hepatoprotective flavonoids (isorhamnetin and luteolin) in the extract. It was concluded from the present study that V. odorata has hepatoprotective activity against paracetamol-induced liver injury in mice.

Materials and Methods

Plant material

Aerial parts of V. odorata were collected from Faisalabad (Pakistan) and identified from Department of Botany, University of Agriculture, Faisalabad, Pakistan. The plants were kept in herbarium for future reference. The plants were shade-dried and powdered by a mechanical grinder. The powdered material was extracted by the
A method of cold maceration using aqueous methanol (70:30) as solvents. After soaking the powdered extract for 48 hours, it was passed through muslin cloth and then through filter paper for filtration. Rotary evaporator was used for concentrating the extracts.

**Experimental animal**

Swiss Albino mice weigh about 20-35 g were used. Mice obtained from the NIH Islamabad. The animals were kept in animal house of College of Pharmacy, housed in standard condition and feed with standard pellet diet and water ad libitum (Vadivu et al., 2008).

**Experimental protocol**

Animals were divided into five groups containing six animals each. Group I was control group received distilled water only. Group II received paracetamol at the dose of 250 mg/kg orally dissolved in water daily for 7 days. Group III was treated with silymarin as standard drug at the dose of 50 mg/kg p.o daily for 7 days and received paracetamol 3 hours after silymarin. Group IV was treated with 250 mg/kg p.o of aqueous methanolic extract of *V. odorata* for 7 days and received paracetamol 250 mg/kg p.o for 7 days 3 hours after the extract dose. Group V was treated with 500 mg/kg p.o aqueous methanolic extract of *V. odorata* for 7 days and received paracetamol 250 mg/kg p.o for 7 days 3 hours after the extract dose (Sabir and Rocha, 2008).

After completing the last dose mice sacrificed by cervical dislocation. By decapitate sedated mice we can draw the maximum Blood from mice and the blood was collected in eppendorff tubes and the clot was dispersed with glass rod and then centrifuged at 3000 rpm for 15 min. Then separated out the serum that was used to estimate SGPT, SGOT and serum alkaline phosphatase to present the parameter of LFT and serum total bilirubin used as excretory function test. The liver from animals was separated for histopathological studies (Ali et al., 2013).

**Identification of active constituent by HPLC**

Hepatoprotective activity may be due to flavonoids (Ali et al., 2013). HPLC may be used to separate and identify the flavonoids. HPLC protocol was used as described elsewhere (Saddiqe et al., 2011). By using the said protocol, retention time for quercetin is 2.05, myricetin 1.75, isorhamnetin 2.72, kaepferol 2.6, luteolin 1.98.

**Statistical analysis**

One-way ANOVA (analysis of variance) was used for statistical analysis. Results were represented by mean ± SE.

**Results**

Serum biochemical parameters are shown in Table I. Paracetamol administration caused damage to hepatocytes demonstrated by raised level of liver enzymes (ALT, AST and ALP) and total bilirubin as compared to control. Elevated levels of these enzymes are indicative of cellular damage and loss of functional integrity of hepatocytes. *V. odorata* pretreatment significantly restored elevated levels of ALT with 250 and 500 mg/kg (p<0.01), AST with 250 and 500 mg/kg (p<0.01), ALP with 250 mg/kg (p<0.01) and with 500 mg/kg (p<0.001) and total bilirubin with 250 mg/kg (p<0.01) and with 500 mg/kg (p<0.001). Both doses (250 mg/kg and 500 mg/kg) provided significant approximately similar reduction in biochemical parameters except ALP and total bilirubin. Results for histopathological examination are given in Figure 1. The normal (control) group receiving distilled water only has normal nuclei. While the administration of paracetamol causes necrosis, inflammation and ballooning of cells. The group treated with silymarin has mild inflammation and necrosis and ballooning was also absent. The groups treated with both doses (D, E) of *V. odorata* have few inflammatory cells but most of the cells gained normal structures as were present in control group (A).

Phytochemical analysis for qualitative determination of

| Treatment groups | ALT (U/L) | AST (U/L) | ALP (U/L) | Total bilirubin (g/dL) |
|------------------|-----------|-----------|-----------|-----------------------|
| Normal (control) | 44.0 ± 2.7 | 62.8 ± 3.0 | 218.0 ± 8.3 | 0.7 ± 0.0 |
| Paracetamol (250 mg/kg) | 134.5 ± 12.1 | 90.5 ± 5.2 | 325.0 ± 12.6 | 1.6 ± 0.1 |
| Silymarin 50 mg/kg + paracetamol (250 mg/kg) | 47.3 ± 4.0b | 44.8 ± 3.6b | 172.3 ± 18.1b | 0.7 ± 0.0b |
| Extract (250 mg/kg) + paracetamol (250 mg/kg) | 81.0 ± 6.3a | 65.8 ± 3.6a | 287.8 ± 11.1a | 0.8 ± 0.0a |
| Extract (500 mg/kg) + paracetamol (250 mg/kg) | 57.8 ± 3.6a | 58.8 ± 4.2a | 205.8 ± 8.9b | 0.8 ± 0.0b |

Data are mean ± SE; *p<0.01; **p<0.001
active constituents through HPLC showed presence of two flavonoids: isorhamnetin and luteolin (Figure 2).

**Discussion**

The liver is the major organ of the body and metabolizes many nutrients carbohydrates, proteins, lipids; and also metabolize the drugs, and excrete them from the body (Saleem et al., 2010). Many drugs and chemicals cause the damage to the liver. The existing hepatoprotective drugs are not full of the efficacy. Efficacy of a drug may be increased by increasing the drug delivery to the target site by the use of polymers (Khalid et al., 2009; Hussain et al., 2011) or through nanotechnology (Naz et al., 2012; Ehsan et al., 2012), synthesis of new drugs, either by the use of proteomics (Qadir, 2011), or synthesis from lactic acid bacteria (Masood et al., 2011), or marine microorganisms (Javed et al., 2011). However, now-a-days, the trend is also being changed to the use of herbal products for their use as their anti-inflammatory (Qadir, 2009), hypotensive (Qadir, 2010), hypoglycaemic (Qadir and Malik, 2009; Qadir and Malik, 2010), amoebicidal (Asif and Qadir, 2011), anti-fertility, cytotoxic, antibiotic (Amin et al., 2012), spasmolytic, bronchodilator (Janbaz et al., 2013), anti-oxidant (Janbaz et al., 2012) and hepatoprotective (Ahmad et al., 2012).

*V. odorata* was considered to be a hepatoprotective traditionally (Amiri et al., 2014), which was proven by...
this study. Results showed that paracetamol damages the liver but after taking standard hepatoprotective drug (silymarin), liver cells cured. Mice which treated with V. odorata were also looking healthy but the results were less significant than the silymarin. Significant reduction (p<0.01) of biochemical markers was observed with aqueous methanolic extract of V. odorata in paracetamol-treated mice. Two doses (250 and 500 mg/kg) were screened to evaluated hepatoprotective potential of this plant. There were no significant difference (p>0.05) between two doses in terms of liver enzymes e.g. ALT, AST and ALP. This is indicative that V. odorata doesn’t cause dose-dependent reduction of liver enzymes. On the other hand, the two doses have significant difference (p<0.01) when compared each other for reduction of total bilirubin. Direct proportion relationship of dose-response can be confirmed by more focused studies.

The biochemical results were further supported by histopathological findings. Paracetamol administration lead to damages normal hepatocyte’s structure that were restored with silymarin. Similar effects were observed with plant extract showing improvement in inflammation-necrosis score and reversal of normal hepatocytes.

Results were further supported by HPLC, in which two flavonoids were found isorhamnetin and luteolin. These flavonoids belong to flavonols class which is well reported to have hepatoprotective activity (Singab et al., 2005; Chattopadhyay et al., 2003).

It was concluded from the present study that V. odorata has hepatoprotective activity against paracetamol-induced liver injury in mice.

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