



## Article

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## Original article

Quantification of biochemical compounds in *Bauhinia Variegata* Linn flower extract and its hepatoprotective effect

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## ABSTRACT

Liver disorders may occur as a result of exposure to chemical compounds capable of inducing the oxidative stress and hepatic injuries. The aim of present study was to investigate the effects of flower extracts of *B. Variegata* for the treatment of liver injury induced by the CCl<sub>4</sub>. About 1 ml/kg body weight (b.w) of CCl<sub>4</sub> was induced to experimental mice by intraperitoneal way for 14 days. The methanol and chloroform extracts (100, 200 and 300 mg/kg b.w) were administered to experimental animals for 14 days along with standard drug Silymarine (100 mg/kg b.w). The extracts alone showed no evidence of hepatic toxicity but animals exposed to CCl<sub>4</sub> without the treatment with *B. Variegata* presented variations in levels of liver enzymes, antioxidant enzymes, proteins and blood cells as well as injuries in liver cells were also observed during histopathological study. However, after the treatments especially with 300 mg/kg b.w of methanol flower extracts levels of liver markers (ALT, AST and ALP), antioxidant enzymes and blood cells decreases and turned towards normal levels. Whereas level of total proteins and bilirubin was improved and damaged liver cells were repaired. The curative activity of flower extracts can be correlated to the higher potential of antioxidants and occurrence of Quercetin and some other organic compounds those were investigated from flower extracts of *B. Variegata* during HPLC and GC-MS analysis. The finding of this study supports the use of *B. Variegata* flower formulation in folk medicines.

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## 1. Introduction

Medicinal plants play significant role as raw material for important drugs; those are providing remedies against many human disorders. Report of world health organization (WHO) interpret that

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almost 80% of the world's population rely on traditional system of folk remedies for primary health care and that is providing considerable economic benefit for the treatment of various diseases (Igoli et al., 2005; Munazir et al., 2012). Medicinal plants produce diversity of secondary metabolites like polyphenols (flavonoids, tannins), saponins, alkaloids and carotenoids which are acts as potent antioxidant compounds (Kaur et al., 2006). Even though flavonoids possess different biochemical activities however the pre-eminent defined property of flavonoids is the capability to act as antioxidant compounds (Cody et al., 1986).

Oxidative stress is triggered by free radicals that deplete the sources of antioxidant enzymes and decline the functioning of the liver cells. Liver is prone to oxidative stress induced damage caused by free radicals. To scavenge free radicals, antioxidant

defense system can reduce their harmful effects (Choudhary and Tandon, 2009). Natural antioxidants either in the form of raw extracts or their chemical constituents are very effective to prevent the destructive processes caused by oxidative stress which is accountable for a lot of disorders in humans such as infections, cancer, diabetes, Alzheimer's diseases, arthritis, cardiovascular diseases and AIDS (Oluyemi et al., 2007; Zengin et al., 2011). Drugs derived from natural sources play a considerable role in the prevention and treatment of human diseases (Farnsworth, 1993).

About 6,000 diverse varieties of plants are found in Pakistan's flora and assert enormous exotic and intuitive plant species as the possible resource for improving health standards of local occupants in long term future across numerous cultures. Rural communities rely on medicinal plants for curing their ailments due to their quick accessibility (Mustafa et al., 2016; Irum et al., 2019).

*Bauhinia Variegata* Linn. is an edible plant belongs to family (Fabaceae/Leguminosae) consists of approximately 300 species. The plant is broadly dispersed in most tropical countries, including Africa, Asia and South America. The leaves, flowers and stem-bark of *Bauhinia* have been used over and over again in folk medicine as a remedy for different kinds of pathologies (da Silva and Filho, 2002; Cavalcanti and Favoretto, 2005; Filho, 2009). Young flowers have been used by inhabitants of hilly areas from northern east of Punjab as vegetable to get relief from different diseases including liver disorders. It is a common practice to isolate compounds from plant extracts, a number of different separation techniques including HPLC and GC-MS are widely used to get possible information about pure compounds. In present work flower extracts of *B. Variegata* were assessed for their possible phytochemicals contents as well as for *in vivo* and *in vitro* bioactivities.

## 2. Material and methods

### 2.1. Plant samples collection

The fresh young flowers of *Bauhinia Variegata* Linn were collected from Tehsil Kotli Sattian (District Rawalpindi) in the flowering period of March - April 2016. The flower samples were identified by experienced taxonomist Dr. Mushtaq Ahmad, Associate Professor in the Department of Plant sciences, Quaid-i-Azam University Islamabad Pakistan. The voucher specimen was maintained in the Herbarium of Pakistan (ISL) Department of Plant sciences, Quaid-i-Azam University for future reference (BV115).

### 2.2. Plant extraction

The fresh flowers of *Bauhinia Variegata* Linn (BV) were washed with double distilled water and dried in the shade for 4–5 days. The dried flowers were crushed into fine powder by using Willy machine. There has been no mesh sizing done and the coarse powder was used for extract preparation using organic solvents. A 10% solution of flower powder in methanol (BVM) and chloroform (BVC) was prepared followed by solvent extraction. The obtained extract was dried and evaporated in the rotary evaporator. Then the dried extracts were stored at 4 °C for more chemical analysis.

### 2.3. Phytochemicals estimation

The dried flower extracts were used for the quantitative assessment of phytochemicals to quantify the total phenols (Harborne, 1998), total flavonoids (AOAC 2003), saponins (Marinova et al., 2005), alkaloids and tannins (Abbasi et al., 2015).

## 2.4. Antioxidants assays

### 2.4.1. DPPH assay

The free radical scavenging ability of the plant extracts were assessed by using 1,1 diphenyl 1–2 -picryl-hydrazyl (DPPH) assay. Plant concentrations (20 to 100 µg/ml) with DPPH solution (2 ml) were incubated in darkness (30 min). Measurement of test sample was recorded at 517 nm against blank mixture (DPPH). Standard or positive control gallic and ascorbic acid were used for comparison. Inhibition concentration (IC<sub>50</sub>) was obtained from linear regression equation. Antioxidant potential is achieved by following formula:

$$\text{DPPH}\% = [Z^x - Z^y / Z^x] \times 100$$

x = Absorbance without plant extract

y = Absorbance with plant extract

### 2.4.2. ABTS scavenging activity

2,2'-azino-bis-3-ethylbenzthiazoline-6-sulphonic acid (ABTS) method is defined by (Ashafa et al., 2010). Oxidation of ABTS solution (3 mM) was done by potassium persulfate (2.5 mM) and the above mixture solution was placed in dark (12 h). After incubation, mixture was diluted with the help of distilled water to get an absorbance of 2.5. Concentrations of plant extract (20–100 µg/ml) and positive controls (standard) were analyzed at 734 nm. The Scavenging activity was determined by using formula:

$$\text{ABTS}\% = [Z^x - Z^y / Z^x] \times 100$$

x = Absorbance without plant extract

y = Absorbance with plant extract

### 2.4.3. H<sub>2</sub>O<sub>2</sub> assay

The oxidant scavenging of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) was used according to the method of (Ruch et al., 1989). H<sub>2</sub>O<sub>2</sub> solution (4 mM) was formulated in phosphate buffer (50 mM) with 7.4 pH. Put the mixture in plant concentrations to get their volume up to 4 ml followed by shaking and incubation (10 min). Absorbance of blank (phosphate buffer), test samples and standards were taken at 230 nm. Scavenging capacity was assessed by the formula:

$$\text{Scavenging}\% = 1 - \frac{\text{Sample absorbance}}{\text{Control absorbance}} \times 100$$

### 2.4.4. Estimation of flavonoids with high performance liquid Chromatography (HPLC)

HPLC analysis was performed via Shimadzu HPLC system (Tokyo, Japan) specified with C18 column (250 × 4.5 mm, 5 µm), gradient pump with detector (UV/Visible). Biocompounds were extracted by different mobile phases (Acetonitrile & 0.1% phosphoric acid with the ratio of 36:64), and the taken volume of injection was 20 µl. Quercetin from the samples was quantified at 285 nm per flow rate (1 ml/min). The standard used in above analysis was Quercetin and all the measurements were taken in triplicates.

### 2.4.5. Gas chromatography-Mass Spectrometry analysis (GC-MS)

GC-MS model QP2010 (Shimadzu®) equipped with following specifications; RTX- 5MS capillary column (cross bond 5% diphenyl - 95% dimethylpolysiloxane), film thickness (30 m × 0.25 mm × 0.25 µm) and carrier gas (helium) in split less inject mode @ 250 °C. Provisions of column include 1.2 ml/min flow rate @ linear velocity mode, oven temperature programmed @ 150 °C for 1 min and then at 4 °C /min to 150 °C for 10 min. Volume of injection (0.2 µl), 50:1 split ratio and injector temperature (275 °C) with N<sub>2</sub> carrier gas (1.0 ml/min). A mass spectrum was operated at 70 eV electron ionization in SIM (Selected Ion Monitoring) manner and ions were

taken (*m/z*). Peaks obtained were tinted and compared with NIST library's spectral data base for compound identification.

#### 2.4.6. Identification of compounds

GC–MS mass spectrum interpretation was performed by using NIST (National Institute of Standard and Technology). Compound name, structure, molecular weight and percentage were estimated by the comparison of an average area with total area. Spectra of unidentified components were matched with Software Turbo mass 5.2 (2005 version).

#### 2.4.7. Hepatoprotective effect

Total 50 albino male mice weighing ( $50 \pm 2.0$  g) were obtained from NIH (National institute for health) Islamabad. Experimental work followed all precautions and guidelines proposed by ethical committee of an Institute (National Veterinary Lab, Islamabad). Mice were kept in the animal house with following conditions: Standard animal house conditions were observed with special monitoring and maintenance. commercial chow (Feed Mills, Islamabad), water *ad libitum* and proper.

#### 2.4.8. Acute oral toxic study

The study was performed to standardize the dose of botanicals for animals as previously manifested by (Gulfraz et al., 2008).

#### 2.4.9. In-vivo study

Animals were categorized into ten groups and each group consists of five mice. Group **I** and **II** were named “Normal group” and “Olive oil group”. They were provided normal feed till 14 days with 1 ml olive oil to olive group. Group **III** were given  $\text{CCl}_4$  (1 ml/kg b. w) by intraperitoneal way for 14 days to induce toxicity. Group **IV**, **V** and **VI** were given methanol extracted plant drug @ 100 mg/kg b. w, @ 200 mg/kg b.w and @ 300 mg/kg b.w when toxicity is created by  $\text{CCl}_4$ . Group **VII**, **VIII** and **IX** were delivered 100 mg/kg b.w, 200 mg/kg b.w and 300 mg/kg b.w chloroform extracted plant drug after  $\text{CCl}_4$  induction. Group **X** was provided standard control drug “Silymarine” 100 mg/kg b.w to cure toxicity of  $\text{CCl}_4$ . All the plant and standard drug doses given by gavage. On fifteenth day animals were initially weighed and then sacrifice for the removal of blood and liver for different examinations. Blood was centrifuged @3000 rpm (10 min) for serum separation. Liver was removed, washed and weighed for the comparison. Subsequently, blood, serum and liver were stored at  $-20^\circ\text{C}$  prior to analysis.

#### 2.4.10. Biochemical analysis

The biochemical markers (AST, ALP, ALT and Bilirubin) were verified with kits (AMS diagnostic, Italy). Red Blood Cells (RBCs), White Blood Cells (WBCs) and Platelets were calculated with the procedures by (Dacie, 2006). Liver derived antioxidant enzymes i.e. SOD, GPx, CAT & total protein were assessed by the procedures of (Winterbourn and Metodiewa, 1999; Misra and Fridovich, 1972; Flohe and Günzler, 1984; Lowry et al., 1951).

#### 2.4.11. Histopathology assessment

Mice liver was wisely handled after slaughtering and saved in formalin (10%) to protect it from oxidation or damage. Tested specimens were dehydrated in alcohol then passed into different xylene changes and fix into paraffin section. For histopathological investigations, specimen was cut by microtome with 5  $\mu\text{m}$  thickness, mounted at glass slides and after that stained through Ehrlich's hematoxylin as well as eosin counter stain.

#### 2.4.12. Statistical analysis

Statistical software Graph pad prism 7 was used to calculate mean ( $\pm$ ), SE (standard error), probability and Pearson coefficient correlation.

### 3. Results and discussion

#### 3.1. Quantitative phytochemical screening

The beneficial properties of medicinal plants are probably due to occurrence of various secondary metabolites, those serves as defense mechanism against many microorganisms (Britto and Sebastian, 2012). Therefore due to strong evidence obtained during screening of phytochemicals various secondary metabolites like flavonoids, phenols, tannins, saponins, alkaloids and carotenoids were quantified from various flower extracts of *B. Variegata* (Table 1). *B. Variegata* possesses significantly higher quantity of total phenols  $21.34 \pm 2.63$  and  $17.66 \pm 1.3$  mg/g in both DVM and DVC. Subsequently  $18.16 \pm 3.1$  and  $12.18 \pm 0.21$  mg/g flavonoid contents are present in DVM and DVC. Methanol extracted more quantity of phytochemicals as compare to chloroform. Methanol has a potential for the diverse richness of phytochemical extraction and found to be more effective for isolating numerous phytoconstituents (Maqsood et al., 2017). The initial approximation may be useful in uncovering the bioactive compounds and subsequently may lead to the drug discovery and development (Bhuiyan et al., 2009).

All the values were taken in triplicates, Mean  $\pm$  standard deviation (SD) while  $p < 0.01$ ,  $p = p < 0.05^\circ$ , QE = Quercetin equivalent and GAE = Gallic acid equivalent.

#### 3.2. Antioxidant assays

In the present study, three different antioxidant methods have been used for evaluation of free radical scavenging activity of methanol and chloroform extracts of *B. Variegata* (Table 2). It was observed that flower extracts of *B. Variegata* has produced strong antioxidant activities (BVM:  $\text{IC}_{50} = 21.93 \pm 0.43$ ,  $47.25 \pm 0.39$  and  $47.88 \pm 0.16$   $\mu\text{g/ml}$ ) (DVC:  $\text{IC}_{50} = 34.85 \pm 0.33$ ,  $48.2 \pm 0.61$  and  $59.02 \pm 0.80$   $\mu\text{g/ml}$ ) against DPPH, ABTS and hydrogen peroxide antioxidant assay. BV extracts were more active against DPPH scavenging assay. Antioxidant potential of medicinal plant is thought to be accredited by the presence of their active constituents like flavonoids and phenols. Excessive production of reactive oxygen species or insufficient antioxidant resistance induces oxidative stress in human body and cause degenerative diseases. Antioxidant activity control series of mechanisms e.g. scavenges free radicals, decomposes peroxides, reducing capacities, prevent binding of transition metal ions and inhibit chain initiation. The achieved results in the present study concord the previous findings described for the root and stem analysis of *B. variegata* (Rajani and Ashok, 2009) (see Table 3).

#### 3.3. Quercetin analysis by HPLC

Analysis of methanol flower extracts of *B. Variegata* with HPLC revealed that significant quantity of quercetin (3.5 mg/g) at peak area of 248,178 and 12.5 retention time (Figs. 1 and 2). Flavonoids being an important group of secondary metabolites play a dynamic role for the protection against various diseases. Plant phytochemicals such as flavonoids control the activity of enzymes and human cells, and found to have antitumor, anti-hepatotoxic, anti-inflammatory and anti-allergic activities (Simpson et al., 2011). Moreover flavonoids provide health benefits and have oxidant quenching activities in both *in vivo* and *in vitro* systems by their role as dietary component (Cook and Samman, 1996). Quercetin is a known bioflavonoid has an antioxidant & anti-inflammatory activity and exhibited unique anticancer properties.

Quercetin ( $\text{C}_{15}\text{H}_{10}\text{O}_7$ ) is a poly-phenolic heterocyclic flavonoid compound contains 302.235 g/mol molecular weight and

**Table 1**  
Quantification of various phyto-constituents from flower extracts of *Bauhinia Variegata*.

Sample extracts	Total Flavonoids (mg QE/g)	Total phenols (mg GAE/g)	Alkaloids (mg/g)	Tannins (mg/g)	Saponins (mg/g)
Methanol extracts	18.16 ± 3.1 <sup>b</sup>	21.34 ± 2.63 <sup>o</sup>	11.89 ± 0.25 <sup>o</sup>	9.63 ± 0.35 <sup>b</sup>	5.29 ± 0.85 <sup>p</sup>
Chloroform extracts	12.18 ± 0.21 <sup>b</sup>	17.66 ± 1.3 <sup>b</sup>	8.18 ± 0.24 <sup>o</sup>	7.38 ± 0.56 <sup>o</sup>	3.38 ± 0.24 <sup>o</sup>

**Table 2**  
IC<sub>50</sub> values of radical scavenging potential of various extracts of *B. Variegata*.

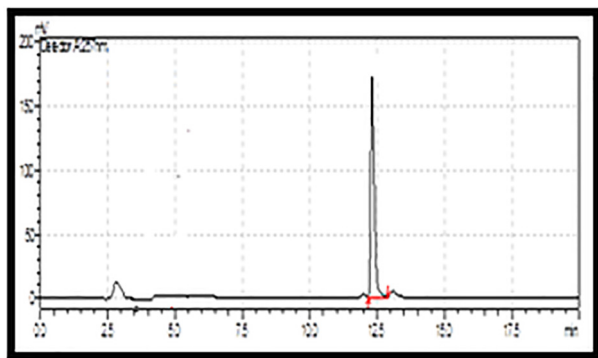
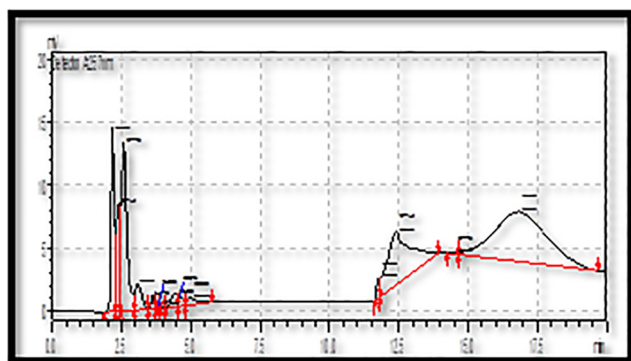
Extracts	DPPH Assay	H <sub>2</sub> O <sub>2</sub> Assay	ABTS Assay	Ascorbic acid	Gallic acid
Methanol	21.93 ± 0.43 <sup>b</sup>	47.88 ± 0.16 <sup>o</sup>	47.25 ± 0.39 <sup>o</sup>	7.22 ± 0.38 <sup>o</sup>	2.434 ± 0.74 <sup>p</sup>
Chloroform	34.85 ± 0.33 <sup>b</sup>	59.02 ± 0.80 <sup>o</sup>	48.2 ± 0.61 <sup>o</sup>	8.69 ± 0.39 <sup>b</sup>	5.87 ± 0.22 <sup>b</sup>

Results are Means ± SD, (n = 3), expressed in µg/ml whereas <sup>o</sup> = p < 0.01<sup>b</sup> = p < 0.05

**Table 3**  
GC–MS analysis of Organic compounds identified in *Bauhinia Variegata*.

Peak	R.T	Area (%)	Molecular Formula	Molecular weight	Name of compound
1	1.136	4.213	C <sub>4</sub> NiO <sub>4</sub>	170	Nickel tetracarbonyl
2	1.178	10.321	C <sub>3</sub> H <sub>6</sub> O	58	Propanone
3	15.776	35.092	C <sub>38</sub> H <sub>68</sub> O <sub>8</sub>	652	1-(+)-Ascorbic acid 2, 6-dihexadecanoate
4	19.215	35.740	C <sub>21</sub> H <sub>40</sub> O <sub>4</sub>	356	2,3-Dihydroxypropyl elaidate SS 2,3-Dihydroxy
5	19.626	10.314	C <sub>18</sub> H <sub>36</sub> O <sub>2</sub>	284	Stearic acid
6	21.907	1.580	C <sub>44</sub> H <sub>86</sub> O <sub>3</sub>	662	Docosanoic anhydride
7	27.710	0.616	C <sub>23</sub> H <sub>40</sub> O <sub>4</sub>	380	Sebacic acid
8	27.940	0.710	C <sub>60</sub> H <sub>122</sub>	842	Hexacontane
9	29.510	0.271	C <sub>36</sub> H <sub>70</sub> O <sub>2</sub>	534	Hexadecenoic acid
10	30.750	1.140	C <sub>54</sub> H <sub>108</sub> Br <sub>2</sub>	914	Tetrapentacontane

Organic compounds along with retention time (R.T) and Peak area (%) obtained by GC–MS analysis.

**Fig. 1.** Chromatogram of Quercetin standard.**Fig. 2.** HPLC analysis of Quercetin from *B. variegata*.

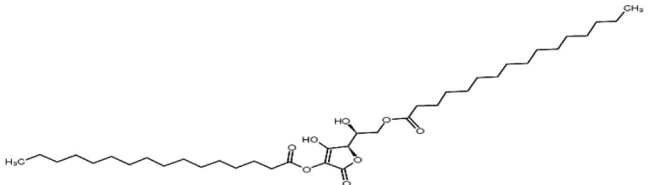
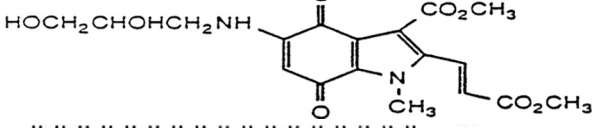
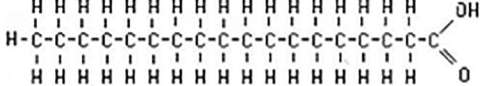
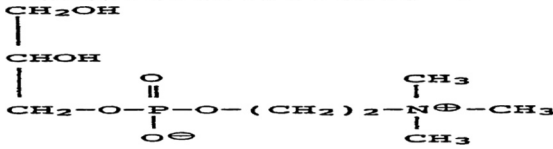
commonly found in fruits and vegetables as most abundant dietary flavonols with average intake of 25–50 mg (Formica and Regelson, 1995) (see Table 4).

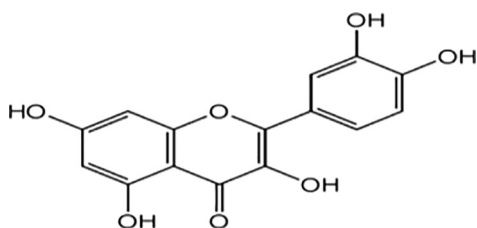
### 3.4. Identification of compounds by GC–MS

Methanol flower extracts of *B. Variegata* was subjected to Gas Chromatography Mass Spectrometry analysis and results are demonstrated in Fig. 3 and Table 2. Comparison of chromatogram and compounds gave peak area (%) and further information was attained from the data base of NIST library. GC–MS spectrum showed 10 compounds and out of which 4 compounds illustrated higher substantial peaks. Results showed that 1-(+)-Ascorbic acid 2, 6-dihexadecanoate (35.092%), Dihydroxypropylelaidate (35.740%), Stearic acid (10.314%) and Docosanoic anhydride (1.580%) was observed on Peak 3, 4, 5 and 6 in BV flower extract.

After comprehensive survey of literature it was known that these organic compounds were not earlier reported from leaves, flower and root extracts of *B. Variegata*, therefore our claim for first time investigation of these compounds from flower of extracts of *B. Variegata* is strongly justified. Hence according to our results presence of biologically important organic compounds in *B. Variegata* is a good proof for the use of this extract in folk medicines against human disorders (Pinheiro et al., 2017). Furthermore 1-(+)-Ascorbic acid 2, 6-dihexadecanoate (C<sub>38</sub>H<sub>68</sub>O<sub>8</sub>) is an essential fatty acid and ascorbic acid is a strong antioxidant compound and being use as food additive, whereas 2,3-Dihydroxypropylelaidate SS 2,3-Dihydroxy (C<sub>21</sub>H<sub>40</sub>O<sub>4</sub>) is a very important organic compound that is not reported largely from plant extracts but scientists believed this compound have very important role to prevent many human disease. Fatty acids reduce oxidative stress by playing an important role in the functionality of liver cells during injury (Beeharry et al., 2003; Hernandez et al., 2016).

**Table 4**  
Molecular structure of the compounds identified by GC–MS.

Compounds	Molecular formula	Molecular Structures
1-(+)-Ascorbic acid 2, 6-dihexadecanoate	C <sub>38</sub> H <sub>68</sub> O <sub>8</sub>	
2,3-Dihydroxypropyl elaidate SS 2,3-Dihydroxy	C <sub>21</sub> H <sub>40</sub> O <sub>4</sub>	
Stearic acid	C <sub>18</sub> H <sub>36</sub> O <sub>2</sub>	
Docosanoic anhydride	C <sub>44</sub> H <sub>86</sub> O <sub>3</sub>	



**Fig. 3.** Chromatogram showing analysis of methanol flower extracts of *B. Variiegata* by using GC–MS.

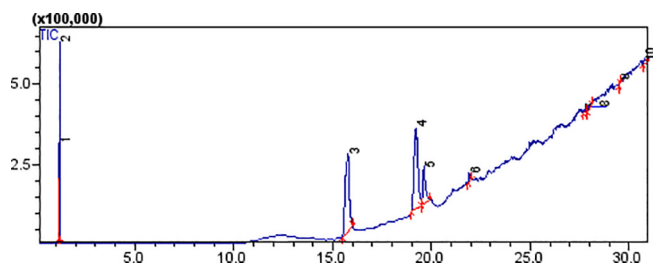
### 3.5. Hepatoprotective effect

#### 3.5.1. Acute toxicity

Study showed no mortality or noticeable behavioral changes in the animals (mice) of all groups. Methanol flower extracts of *B. Variiegata* were observed innocuous up to 500 mg/kg body weight as our earlier study reported same dose for other plant (Gul et al., 2017).

#### 3.5.2. Response of plant extracts on body weight

The weight of mice was suddenly reduced ( $50 \pm 2.0$  to  $48.3 \pm 0.5$ ) when CCl<sub>4</sub> was induced in mice from all groups. However with the treatment of plant extracts particularly 300 mg/kg body weight (b.w) methanol flower extract has shown increased in the body weight which was comparable with animals from



**Fig. 4.** Comparison of initial and final body weight of mice.

normal control group (Fig. 4). Carbon tetrachloride is readily absorbed from the gastrointestinal tract causes vomiting gastrointestinal pain, cell injury and liver necrosis followed by cell degeneration and decreased in body weight (Carlberg and Mannervik, 1975). The recovery of body weight of animals proved the protective effects of *B. Variiegata* extracts and due to its treatment weight of animals became normal after passage of time.

#### 3.5.3. Liver and antioxidant enzymes

The levels of different biochemical markers i.e. (ALT, AST and ALP) and total direct Bilirubin in the blood serum of trial animals were assessed. It was observed that after administration of CCl<sub>4</sub>, intensities of various parameters were increased (Table 5). While by the consumption of plant drug, ranges of enzymes and Bilirubin became normal and significant results were noted for 300 mg/kg b. w of methanol flower extracts. The effects of flower extracts on hepatic enzymes such as CAT, SOD, GPx and total proteins were also evaluated (Table 6).

When CCl<sub>4</sub> was incorporated in the body of mice it decreased the level of all enzymes when compared to normal level (normal group). However, after treatment of 300 mg/kg b. w methanol flower extracts, levels of these parameters remained increased on dose depended way. It was reported earlier that flavonoids improved viability of cells and repress CCl<sub>4</sub> caused hepatocyte's cellular leakage (Wu et al., 2006). Similarly, several doses of flower extracts have significantly moderated the ranges of aspartate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphatase (ALP) together with direct Bilirubin.

#### 3.5.4. Hematological study

Effects of B.V extracts on RBC, WBC and Platelets levels were assessed. After administration of CCl<sub>4</sub> to experimental animals, levels of RBC, WBC and Platelets were decreased as relate to normal (Fig. 5). Current findings specify that the flower extracts and silymarine standard drug increased above parameters to normal ranges (normal group) and further improvement can depends on amount of dose given. Rapid rise or decline in hematological parameters is the source to weaken animal immune system and these persistent conditions end in severe consequences (Carvalho et al., 2011).

**Table 5**  
Role of flower extracts on Hepatic enzymes.

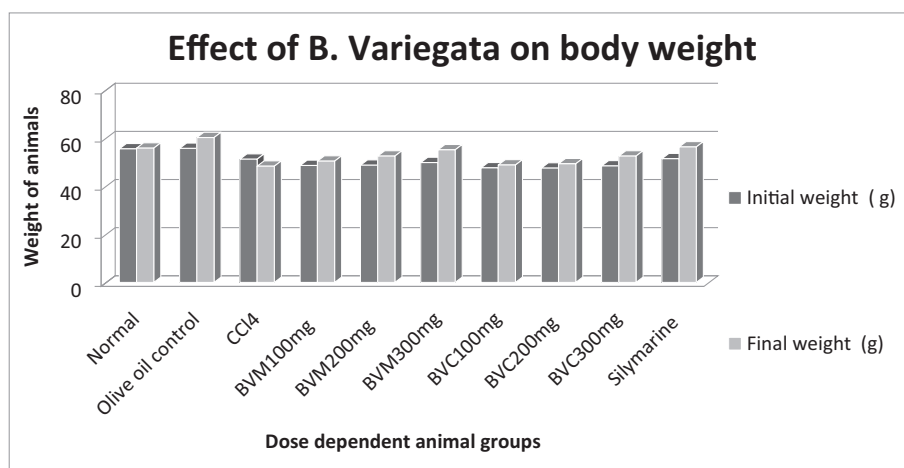
S #	Trial Animals	ALT U/L	AST U/L	ALP U/L	Direct bilirubin g/dl
1	Normal control	38.5 ± 0.5	80.6 ± 0.5	110.6 ± 1.2	0.2 ± 0.01
2	Olive oil	40.6 ± 1.2	63.5 ± 3.6	175.2 ± 1.5	0.3 ± 0.01
3	CCl4	120.4 ± 0.7	127.6 ± 2.5	294.5 ± 2.6	1.9 ± 0.2
4	Methanol flower extract(100 mg/kg b.w) + CCl4	70.6 ± 3.5	84.5 ± 4.7	319.7 ± 4.5	0.9 ± 0.1
5	Methanol flower extract(200 mg/kg b.w) + CCl4	57.4 ± 2.6	79.5 ± 3.5	301.5 ± 2.5	0.7 ± 0.1
6	Methanol flower extract(300 mg/kg b.w) + CCl4	56.5 ± 3.1	63.5 ± 2.8	238.3 ± 1.5	0.7 ± 0.02
7	Chloroform flower extract (100 mg/kg b.w) + CCl4	74.1 ± 3.7	110.5 ± 2.4	337.7 ± 4.6	1.1 ± 0.3
8	Chloroform flower extract (200 mg/kg b.w) + CCl4	72.2 ± 3.2	89.4 ± 4.2	295.6 ± 5.3	1.2 ± 0.4
9	Chloroform flower extract (300 mg/kg b.w) + CCl4	64.5 ± 1.9	81.3 ± 3.5	275.6 ± 2.8	0.8 ± 0.4
10	Silymarine (100 mg/kg b.w) + CCl4	66.4 ± 2.1	76.1 ± 2.5	185.6 ± 2.6	0.5 ± 0.01

Effect of dosage of flower extract on liver markers were recorded (n = 5).

**Table 6**  
Assessment of hepatic anti-oxidant enzymes and total proteins.

Sr.No.	Test animals	CAT (m mol/min mg protein)	SOD (USOD/mg Protein)	GPx (µmol/min/mg Protein)	Total protein g/dl
1	Normal group	8.2 ± 0.3	10.15 ± 0.6	32.10 ± 3.3	3.15 ± 0.4
2	Olive oil	7.9 ± 0.4	10.22 ± 0.7	34.5 ± 3.2	3.1 ± 0.5
3	CCl4	4.75 ± 0.6	5.75 ± 0.4	17.83 ± 1.5	0.75 ± 0.1
4	Methanolic flower extract(100 mg/kg b.w) + CCl4	7.42 ± 0.3	8.29 ± 0.5	24.1 ± 2.6	1.84 ± 0.5
5	Methanolic flower extract(200 mg/kg b.w) + CCl4	7.91 ± 0.7	9.95 ± 0.6	28.72 ± 2.8	2.29 ± 0.7
6	Methanolic flower extract(300 mg/kg b.w) + CCl4	8.15 ± 0.8	11.03 ± 0.8	32.51 ± 3.6	3.10 ± 0.8
7	Chloroform flower extract (100 mg/kg b.w) + CCl4	7.05 ± 0.6	8.04 ± 0.7	22.4 ± 2.1	1.76 ± 0.6
8	Chloroform flower extract (200 mg/kg b.w) + CCl4	7.61 ± 0.4	9.52 ± 0.8	26.25 ± 2.5	2.35 ± 0.3
9	Chloroform flower extract (300 mg/kg b.w) + CCl4	7.99 ± 0.6	10.78 ± 1.5	30.63 ± 3.1	2.92 ± 0.2
10	Silymarin (100 mg/kg b.w) + CCl4	8.11 ± 1.2	11.62 ± 0.6	33.7 ± 1.5	3.22 ± 0.8

Values were taken (n = 5)



**Fig. 5.** Response of the animal group against different doses were recorded n = 5. Whereas RBC, WBC and Platelets count is measured in 10<sup>6</sup> /µl, 10<sup>3</sup> /µl and 10<sup>3</sup> /µl.

**3.5.5. Histopathology**

Alteration observed in the symmetry of liver cells was created by CCL<sub>4</sub> shown in (Fig. 6a) and healing property of methanol flower extract of *B. variegata* is presented in (Fig. 6b). In CCL<sub>4</sub> group impaired liver tissues with altered cellular organization, inflamma-

tion and hepatocyte pycnosis were noticed. Animals treated with 100 and 200 mg/kg b.w recovered the cellular organization to some extent, while the flower extract 300 mg/kg b.w has reinstated the structural arrangement of tissues which verified its curative potential and was analogous to silymarine effect. Histological studies

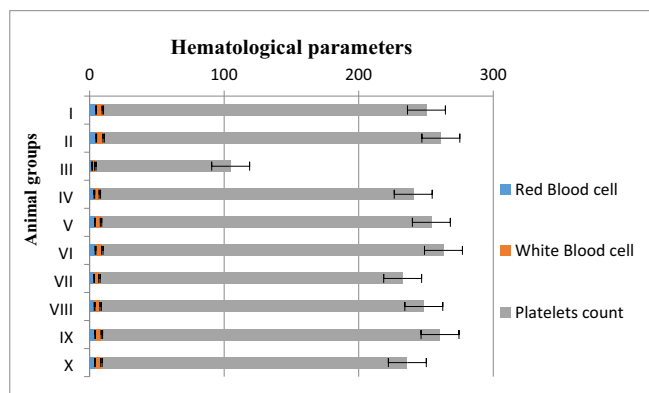


Fig. 6. a: CCl<sub>4</sub> induced toxicity in the liver by changing its symmetry. b: Damaged liver treated with B.V flowers tried to cure the injury.

discovered the dynamic role of flavonoids for the treatment of damaged cells of liver (Hollman et al., 1997; Wu et al., 2006).

Based on the existing results, it is concluded that *B. Variegata* young flower can be used as a natural antioxidants source and its consistent ingestion could provide health benefits to human population by protecting against oxidative stress, hepatic injury and similar other liver disorders. Further detailed in vitro and in vivo correlation based studies along with separation of active constituents are required to unravel the novel approaches for curing free radical induced diseases.

#### Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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