

Evaluation of Anti-Anxiety Effect of *Brassica oleracea* L. Extracts in Experimental Animals

Divneet Kaur¹, Richa Shri², Anjoo Kamboj³

ABSTRACT

Introduction: Oxidative stress in the brain causes nervous system impairment. Hence the use of antioxidants could be an effective approach to prevent disorders such as anxiety. Plants containing phenolic constituents are reported to be potent antioxidants. *Brassica oleracea* L. commonly known as broccoli, is rich in phenolics mainly flavonoids and possesses excellent antioxidant property, however its anxiolytic potential has not been investigated. Thus in the present study antianxiety activity of extracts of broccoli was evaluated in experimental animals. **Materials and Methods:** Various pharmacognostic parameters of the plant material were studied following standard procedures. Groups of six mice each were taken as different treatment groups. Control group received vehicle (1% Carboxy Methyl Cellulose), positive control received Diazepam (2mg/kg) as standard drug while test groups received petroleum ether and hydroalcoholic extracts (50, 100 and 200 mg/kg body weight, p.o.). The effects were evaluated on Elevated Plus Maze (EPM), hole board and mirror chamber tests. Statistical analysis was performed using ANOVA, followed by Turkey's post-hoc test. **Results:** Hydroalcoholic extract shows dose dependent increase in the average time spent and frequency of entries in the open arms of the EPM; decreased latency, increased time spent and frequency of entries in the mirror chamber; increased number of head dips in hole board test as compared to control was observed. These effects were comparable to the effect produced by diazepam. Phytochemical screening of the bioactive extract revealed the presence of alkaloids, phenols, flavonoids and tannins. **Conclusion:** From our study, it can be concluded that hydroalcoholic extract of *Brassica oleracea* at the dose of 200 mg/kg possesses marked anxiolytic activity.

Key words: Antianxiety, *Brassica oleracea*, Elevated Plus Maze, Hole Board, Mirror Chamber Test.

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INTRODUCTION

Anxiety is an adaptive emotion that permits physiological and behavioural changes, to appropriately react to a stressful situation to resolve it by fighting or escaping. Anxiety disorder is defined as a feeling of apprehension, uncertainty or tension stemming from the anticipation of imagined or unreal threat.¹ Anxiety affects one-eighth of the population worldwide and has debilitating effects on the quality of life of many people.²

Anxiety disorders can be identified by early diagnosis and effective treatment includes psychotherapy and use of drugs like benzodiazepines.³⁻⁴ Conventional drug therapy have a narrow margin of safety between the anxiolytic effect and unwanted side effects, thus prompting researchers to evaluate new compounds specially plant based drugs having less undesirable effects.⁵

Oxidative stress in the brain causes nervous system impairment. Oxidative stress has also been implicated in depression and anxiety disorders.⁶ Studies have

revealed that antioxidants have a key role as an adjuvant therapy in patients with generalized anxiety disorder and depression.⁷ Flavonoids are reported to be potent antioxidants.⁸⁻¹⁰ Many plants containing flavonoids are reported to possess anxiolytic potential.¹¹⁻¹² Some of the plants containing flavonoids demonstrating antianxiety activity are *Abies pindrow*,¹³ *Calotropis gigantean*,¹⁴ *Matricaria recutita*,¹⁵ *Gelsemium sempervirens*,¹⁶ *Medicago sativa*,¹⁷ *Passiflora coerulea*,¹⁸ *Passiflora incarnate*,¹⁹ *Tilia Americana*,²⁰ *Turnera aphrodisiaca*.²¹⁻²² Flavonoids which possess anxiolytic effects include apigenin,²³ chrysin, kaempferol,²⁴ quercetin.²⁵

In the present study *Brassica oleracea* L., also known as broccoli, which belongs to family Brassicaceae has been evaluated for its anxiolytic potential. Broccoli is an edible green plant in the cabbage family whose large flower head is eaten as a vegetable. Like other species of the Brassica family, broccoli is a rich source of health promoting phytochemicals.^{26,27}

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It is rich in phenolic compounds, particularly flavonoids. It also contains ascorbic acid, vitamins C and E, amino acid, the flavonols quercetin and kaempferol, the carotenoids β -carotene, lutein, and the glucosinolates.²⁸ This is reported to have anti-oxidant, anti-microbial, gastroprotective, anti-cancer, anti-diabetic, cardioprotective and hepatoprotective effects.²⁹⁻³⁵

Broccoli (*Brassica oleracea*) contains phenolic compounds and has marked antioxidant activity has been selected and evaluated for its anxiolytic potential. This plant has never been evaluated for their anti-anxiety potential, therefore it was investigated for its anxiolytic potential. So the aim of the present investigation was to study pharmacognostic parameters and to evaluate the antianxiety activity of petroleum ether and hydroalcoholic extract of *Brassica oleracea* using Elevated Plus Maze (EPM), hole board and mirror chamber tests of anxiety.

MATERIALS AND METHODS

Collection and authentication of plant material

Flower heads of *Brassica oleracea* were procured from Punjab Agricultural University, Ludhiana, India in the month of January 2015. Identity of the plant was confirmed by Head, Raw Materials, Herbarium & Museum at National Institute of Science Communication and Information Resources (NISCAIR), New Delhi, India, vide letter number/ NISCAIR/RHMD/Consult/2014/2577-156 dated 13.02.2015

Drugs and chemicals

Diazepam was obtained from GlaxoSmithKline and petroleum ether, methanol (LR grade) were procured from S.D. Fine-Chem Ltd, Mumbai.

Physicochemical evaluation

The determination of physicochemical parameter is important in determination of adulterants and improper handling of drugs. Various physicochemical parameters were studied following standard procedures.³⁶ All parameters such as moisture content (LOD method), ash values, extractive values were determined in triplicate.

Preparation of extract

The plant was dried in shade and powdered. Powdered drug (750 g) was subjected to successive exhaustive extraction with petroleum ether and methanol:water (70:30) in the Soxhlet apparatus. Petroleum ether (PE) and hydroalcoholic (HA) extracts were dried using rotary evaporator and yield calculated on dry weight basis. Dried extracts were preserved in vacuum desiccator. These solvents selected for extraction were selected broadly to get a non polar and a polar extract.

Phytochemical screening

A phytochemical examination was carried out for both the extracts as per standard methods.³⁷

Animals

Swiss albino mice (weighing 15-30g) were taken from Chandigarh College of Pharmacy, Landran (Regd. No. 1201/a/08/CPCSEA) and kept in polypropylene cages of 5 mice at $22 \pm 1^\circ \text{C}$ on a 12-h light/dark cycle. Water and food were available *ad libitum*. Groups of six mice were randomly assigned to different treatment groups. Control group received vehicle, one group was given standard drug Diazepam (2mg/kg p.o.) while test groups received PE and HA extract (50, 100 and 200 mg/kg p.o.).

Different treatments

Three sets of test doses (50, 100 and 200 mg/kg of body weight) were prepared by suspending dried extracts in vehicle. Diazepam 2mg/kg

suspended in vehicle was used as a standard anxiolytic drug. Vehicle (1% CMC) was used as control. The effects of the various treatments were estimated 45 min after the oral administration of vehicle/ standard drug/ test extracts. The entire tests were carried out between 8:00-14:00. In each experiment, apparatus was cleaned using 5% ethanol before introducing the next animal to preclude the possible cueing effects of odours left by previous subjects.

Animal Model for Anxiety Elevated Plus Maze

The plus maze apparatus consisting of two open arms (16 x 5 cm) and two closed arms (16 x 5 x 12 cm) having an open roof with a plus maze elevated (25 cm) from the floor was used to observe anxiolytic behaviour of animals. The animals were fasted 18 h prior to experiment. The dose administration schedule was so adjusted that each mouse was having its turn on plus maze after 45 min of administration of dose. Each animal was placed in the centre of the elevated plus maze with its head facing the open arms.

During the 5 min experiment, behaviour of the mice was noted as a) preference of the animal for its first entry to open/closed arm b) number of entries into the open arm c) average time spent by the animal in open arm. The antianxiety activity was recorded as average time spent by the animals in the open arms of the EPM. During the entire experiment, each animal was allowed to socialize. Every precaution was taken to ensure that no external stimuli disturbed the animal.³⁸

Mirror chamber Test

The mirror chamber consisted of wooden chamber (40x40x30.5 cm) having a mirror (30x30x30 cm) enclosed within it. Placement of the mirrored cube into the centre of the container forms a 5 cm corridor that completely surrounds the mirror chamber. The animals were placed individually into the chambers of the mirrors at a fixed corner.

During the 5 min experiment following parameters was noted a) latency to enter mirror chamber b) number of entries into the mirror chamber c) total time spent in the mirror chamber.³⁸

Hole-Board Test

The hole board apparatus consisted of a wooden box (40 x 40 x 25 cm) with 16 holes (each of diameter 3 cm) evenly distributed on the base of box. The apparatus was elevated to a height of 25 cm. The number of head dips and the time of head dipping during a 5 min period were recorded.³⁹

Statistical Analysis

All the values were expressed as mean \pm SEM. Statistically significant difference between the groups were calculated by the application of one way analysis of variance (ANOVA) followed by Tukey's, post-hoc test. ***P<0.001 and **P<0.01 were considered statistically significant.

RESULTS

Yield of extracts: The yield of petroleum ether extract was 1.5% w/w and hydroalcoholic extract was 2.5% w/w on dry weight basis.

Phytochemical Screening: The results of phytochemical screening of *B. oleracea* are shown in Table 1.

Physicochemical evaluation: The results of various pharmacognostic parameters assessed are shown in Table 2.

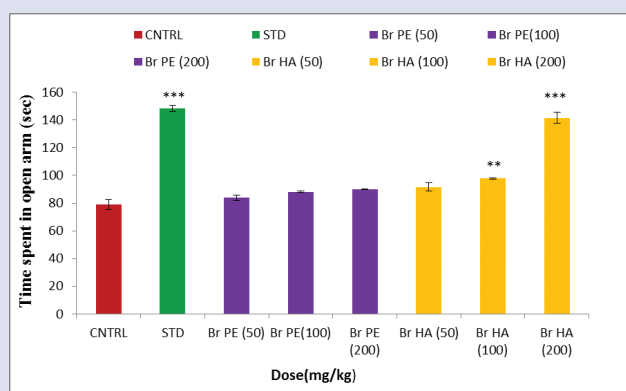
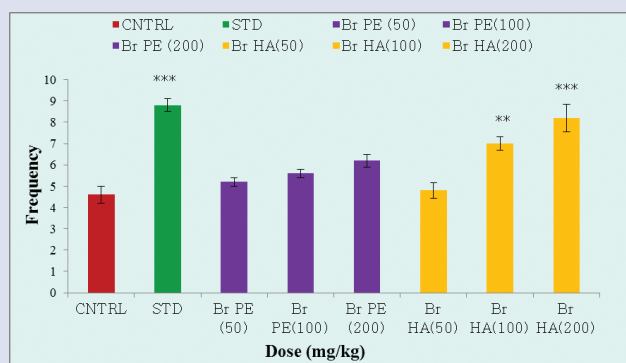
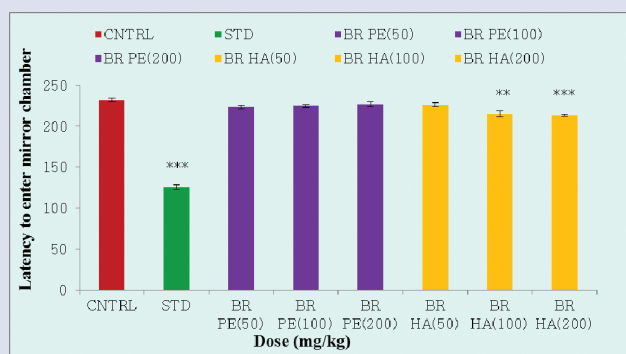
Elevated Plus maze model: Diazepam (2 mg/kg) showed anxiolytic activity, by showing significant (**P < 0.001) increase in average time spent and frequency in open arms. The groups of mice treated with HA (200 mg/kg) and HA (100mg/kg) have shown anxiolytic activity, signifi-

Table 1: Phytochemical analysis of extracts of *Brassica oleracea* L.

Tests	Petroleum ether extract	Hydroalcoholic extract
1. Alkaloids		
(a) Mayer's Test	-	+
(b) Wagner's Test	-	+
(c) Hager's Test	-	+
(d) Dragendroff Test	-	+
2. Carbohydrates		
(a) Molisch's Test	-	+
(b) Benedict's Test	-	+
(c) Fehling's Test	-	+
3. Glycosides		
(Anthraquinone glycosides)		
(a) Borntrager's Test	-	-
(b) Modified Borntrager's Test	-	-
(Cardiac glycosides)		
(c) Killer Killiani's test	-	-
(d) Legal's Test	-	-
4. Saponins		
(a) Froth test	-	-
5. Phytosterols		
(a) Salkowski's Test	-	+
(b) LibermannBurchard's test	-	+
6. Phenols		
(a) Ferric Chloride Test	-	+
7. Tannins		
(a) Gelatin Test	-	+
8. Flavonoids		
(a) Alkaline Reagent Test	-	+
(b) Lead acetate Test	-	+
(c) Shinoda Test	-	+
9. Proteins and aminoacids		
(a) Xanthoproteic Test	+	+
(b) Ninhydrin Test	+	+
10. Diterpenes		
(a) Copper acetate Test	+	+

Table 2: Results of pharmacognostic parameters

Pharmacognostic parameters	Mean±SEM (n=3)
1. Loss on Drying	0.33±0.165 %w/w
2. Ash value	
a) Total ash	1.82±1.82 %w/w
b) Acid insoluble ash	0.146±0.122 %w/w
c) Water soluble ash	2.42±1.22 %w/w
3. Extractive value	
a) Water soluble extract	2.5±1.52 %w/w
b) Alcohol soluble extract	2.0±1.9 %w/w

**Figure 1:** Effect of different extracts of *Brassica oleracea* on the time spent by mice in the open arms of the EPM CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), Br HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.**Figure 2:** Effect of different extracts of *Brassica oleracea* on number of entries (frequency) by mice in the open arms of the EPM CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), Br HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.**Figure 3:** Effect of different extracts of *Brassica oleracea* on the latency to enter mirror chamber by mice in the mirror chamber test CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), Br HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.

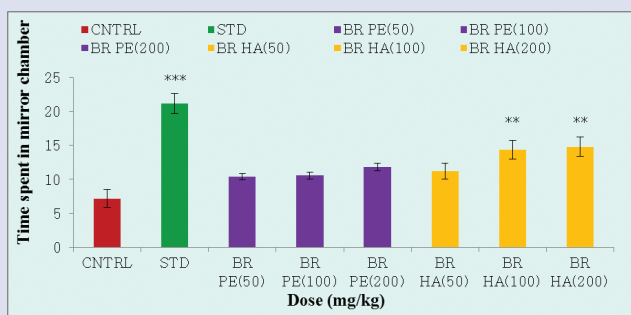


Figure 4: Effect of different extracts of *Brassica oleracea* on the time spent by mice in the mirror chamber in the mirror chamber test CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), BR HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o.). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.

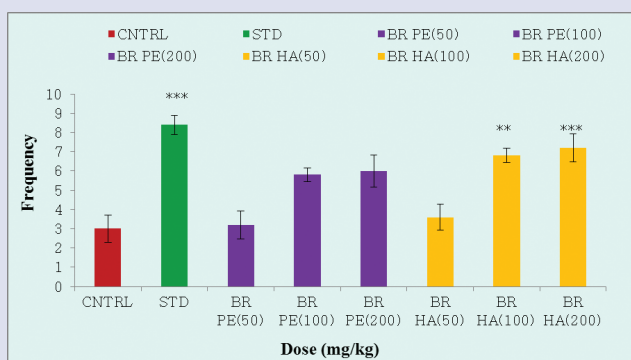


Figure 5: Effect of different extracts of *Brassica oleracea* on the number of entries (frequency) by mice in the mirror chamber in the mirror chamber test CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), BR HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o.). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.

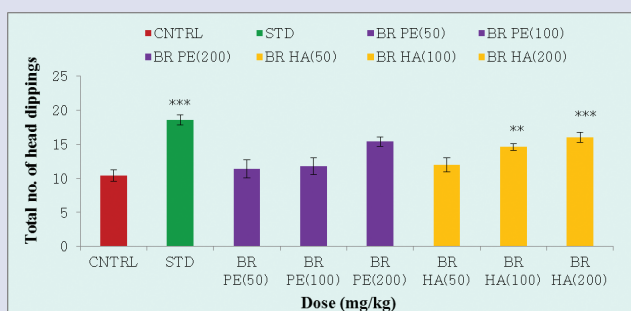


Figure 6: Effect of different extracts of *Brassica oleracea* on the number of head dippings by mice in the the hole board test CNTRL=control (2% CMC p.o.), STD=Diazepam (2 mg/kg p.o.), Br PE=petroleum ether extract (50, 100 and 200 mg/kg p.o.), BR HA=hydroalcoholic extract (50, 100 and 200 mg/kg p.o.). Results are expressed as mean±SEM (n=6); *** P<0.001 and ** P<0.01 as compared to control.

cant (**P < 0.01, ***P < 0.001) in the open arms in comparison to control group as shown in Figure 1 and 2.

Mirror chamber Test: The groups of mice treated with HA (200 mg/kg) and HA (100mg/kg) have shown decrease in latency, increase in time spent and frequency in the mirror chamber significant (**P < 0.001, ***P < 0.01) in comparison to control as shown in Figure 3, 4 and 5.

Hole board Test: With the dose of 50 mg/kg, HA extract of broccoli, there was no significant increase in number of head dips as compared to control. HA extract at 200 and 100 mg/kg increased the number of head dips significantly (**P < 0.001, ***P < 0.01). The standard group showed significant increase in number of head dips (**P < 0.001) as shown in 6.

DISCUSSION

The brain is highly vulnerable to oxidative stress. Oxidative stress can alter neurotransmission, neuronal function and overall brain activity. Oxidative stress has been implicated in neurodegenerative diseases and neuropsychiatric diseases including stress and anxiety.⁶ Antioxidant therapy has been reported to have a protective effect in anxiety disorders. Polyphenols, flavonoids, specific foods and diets rich in antioxidants have been shown to improve antioxidant status and have anxiolytic effects.^{7,9} Plant extracts, which contain phenolic constituents mainly flavonoids, have shown beneficial effects on mental health. Extracts of plants like *Pulsatilla nigricans*,⁴⁰ *Tephrosia purpuria*,⁴¹ *Coriandrum sativum*,⁴² *Gastrodia elata*,⁴³ *Citrus paradisi*⁴⁴ contain flavonoids and possess anti-anxiety activity. Flavonoids have been reported to possess antianxiety potential.¹² Flavonoids and Diazepam are structurally similar.⁴⁵⁻¹² Searching for safer BDZ-receptor (BDZ-R) ligands has demonstrated the existence of a new family of ligands which have a flavonoid structure. *Brassica oleracea* contains large amounts of polyphenols specially flavonoids.⁴⁶ Thus, for the present investigation, Broccoli (*Brassica oleracea*) that contains phenolic compounds and has marked antioxidant activity has been selected and evaluated for its anxiolytic potential. Thus, the antianxiety effect of *B. oleracea* extracts was evaluated on experimental animals using various models. Various physicochemical parameters of the plant were also evaluated which have not been determined so far.

The Elevated Plus Maze is considered to be a valid animal model of anxiety because it uses natural stimulus that is the fear of a new, brightly light open space and the fear of balancing on a relatively narrow raised platform.⁴⁷ Moreover, it is known that anxiolytic agents increase the frequency of entries and the time spent in open arms of the Elevated Plus Maze.⁴⁸ In the present study, HA extract markedly increased the frequency of entries and the time spent by the animals in the open arms.

The Hole-Board test provides a simple method for measuring the response of an animal to an unfamiliar environment. It has been showed that head-dipping behaviour was sensitive to changes in the emotional state of the animal, and suggested that the expression of an anxiolytic state in animals may be reflected by an increase in head-dipping behaviour. In the present study, HA extract at 200 and 100 mg/kg increased the number of head dips significantly.

Animal species exhibit approach avoidance conflict upon placement of a mirror. The extended latency to enter mirror chamber is a parameter of anxiety analogy. In the present study, HA extract has shown decrease in latency, increase in time spent and frequency of entries in the mirror chamber.

The results of the behavioural tests clearly demonstrate that the HA extract shows distinct potential for being developed as an anti-anxiety drug. The phytochemical study showed the presence of phenols and flavonoids in the bioactive extract.

CONCLUSION

From our research study, it is concluded that hydroalcoholic extract of *Brassica oleracea* after acute dosing possess significant anxiolytic activity at dose of 200 mg/kg. The petroleum ether extract was devoid of anti-anxiety effect. Further studies are being conducted to ascertain the bioactive constituent responsible for the activity.

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CONFLICT OF INTEREST

Authors have no conflict of interest.

ABBREVIATION USED

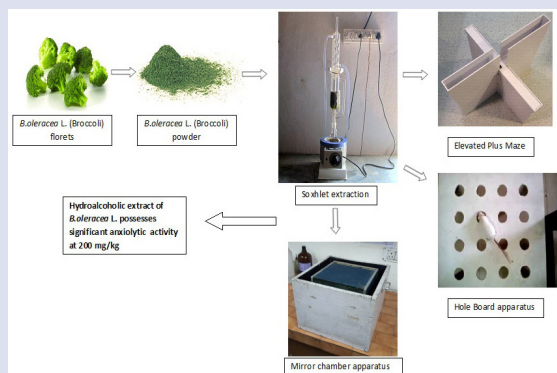
EPM: Elevated Plus Maze; **PE:** Petroleum ether; **HA:** Hydroalcoholic.

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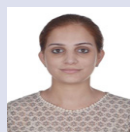
GRAPHICAL ABSTRACT



HIGHLIGHTS OF PAPER

- Various pharmacognostic parameters of the plant material were studied following standard procedures.
- Powdered drug was subjected to successive exhaustive extraction with petroleum ether and methanol:water (70:30) in the soxhlet apparatus and PE and HA extracts were prepared.
- The anxiolytic effects were evaluated by Elevated Plus Maze (EPM), hole board and mirror chamber tests and statistical analysis was done by using one way analysis of variance (ANOVA) followed by Tukey's, post-hoc test.
- Results showed that hydroalcoholic extract of *Brassica oleracea* after possess significant anxiolytic activity at dose of 200 mg/kg. The petroleum ether extract was devoid of anti-anxiety effect.
- The phytochemical study showed the presence of phenols and flavonoids in the bioactive extract.

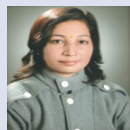
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