Exploration of in vitro thrombolytic, anthelminthic, cytotoxic and in vivo anxiolytic potentials with phytochemical screening of flowers of *Brassica nigra*

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**Abstract**

**Background:** *Brassica nigra* is a plant of Brassicaceae family, which possesses numerous medicinal values. Our present study is intended to assess the potential in vitro thrombolytic, anthelminthic, cytotoxic and in vivo anxiolytic properties of MCE of *B. nigra* flowers. MCE was fractioned for separating the compound on the basis of polarity by using chloroform, n-hexane and ethyl acetate solvent. Thrombolytic and anthelminthic activities were explained by collecting human erythrocytes and earthworms as test models, respectively. Anxiolytic activity was evaluated by elevated plus maze and hole board models while cytotoxic test was conducted through brine shrimp lethality bioassay.

**Results:** MCE revealed the presence of alkaloids, flavonoids, tannin, diterpenes, glycosides, carbohydrates, phenols, fixed oils and fat. In case of thrombolytic test, the MCE, CSF, ASF and n-HSF had produced maximum clot lysis activity at 5 and 10 mg/ml dose conditions. Two different concentrations (10 and 20 mg/ml) of MCE and its fractions showed significant (*p* < 0.05) anthelminthic activities in a dose-dependent manner. Significant anxiolytic activity was observed for all fractions which was comparable to the standard drug diazepam (*p* < 0.05). Again, the cytotoxic screening also presented good potentials for all fractions.

**Conclusion:** From the findings of present study, we can conclude that MCE of *B. nigra* flowers and its fraction possess significant anxiolytic, anthelminthic, anticancer and thrombolytic properties which may be a good candidate for treating these diseases through the determination of bio-active lead compounds.

**Keywords:** *Brassica nigra*, Anxiolytic, Anthelmintic, Thrombolytic, Cytotoxic activity

**Background**

Various types of plants have been used in conventional medicine throughout the world for thousands of years, and they are providing new medications for today [1]. It has been stated that about 64% of world population is using conventional medicine to meet their healthcare needs [2]. A number of current medications used in the treatment of various chronic and severe conditions have been originated from plants which were discovered by the human being in the course of time [3]. Such type of conventional medicines is used in both developed and developing countries for primary healthcare due to their wide biological and medicinal activities, higher safety and low prices [3]. By this way, allopathic medicine has replaced by medicines from natural sources for the treatment of many diseases in recent times that show considerable therapeutic activity [4]. Plants are also providing the raw materials for the treatment of various life-
threatening disorders [5]. This information encouraged us to search for new plants in order to disclosing different pharmacological activities.

Human beings are suffering from various diseases. Among these diseases, thromboembolic disorders such as pulmonary emboli, deep vein thrombosis, strokes and heart attacks are the main causes of morbidity and mortality in both developing and developed countries [6]. Various thrombolytic agents such as alteplase, streptokinase, urokinase and tissue plasminogen activator (TPA) are widely used to dissolve clots [6]. These drugs have some limitations which cause severe and sometimes fatal disorders including haemorrhage, severe anaphylactic reaction and lacked specificity [2]. That is why efforts have been focused towards the findings of natural products from various plant sources which belong to antiplaetelet, anticoagulant, antithrombotic and thrombolytic activities. Again, anxiety disorders are the most prominent problems among many psychiatric disorders and approximately 10–30% of the general population are suffering from this globally [7]. It is a psychiatric disorder that causes significant impairment of personal functioning [7]. Benzodiazepines are the mostly prescribed synthetic drugs for anxiolytic, muscle relaxant, sedative-hypnotic and anticonvulsant actions [8]. These drugs cause impairment in cognitive functions, physical dependence and harmful effects on respiratory, digestive and immune systems of the body [9]. For this reason, the search of new anxiolytic and antidepressant agents with less adverse effects is still an area of great interest to the researchers [7].

Helminth infection, a parasitic disease, is thought to be caused by various parasitic worms such as flukes (Trematodes), roundworms (Nematodes) and tapeworms (Cestodes) [10]. It is highly seen in developing countries including Bangladesh owing to the lack of management and insufficient control measures [11, 12]. Again, cancer is another leading disease in which abnormal cellular growth, lack of control of the cell proliferation, differentiation and death usually occur [13]. There are various ways to address cancer such as surgery, radiation therapy and chemotherapy which aim to eradicate all cancerous cells from the body, but each of the treatment strategy has several limitations like drug resistance, toxicity and low specificity [14]. The synthetic chemotherapeutics which have currently been used possess several limitations, and thus, the initiation of novel chemotherapeutic agents from plant origin is increasing in recent years [15].

Human beings are greatly depending on nature from the very ancient times for their any primary health complaints, and nature is a good source of medicine which can be obtained from various types of plants [2]. Brassica nigra is such type of plant with a great medicinal value. Brassicaceae family plants are thought to be native to the eastern Mediterranean or the surrounding area, but nowadays, it has been grown all over Europe [16]. It is also grown in Bangladesh. It has been cultivated for millennia for fulfilling the demand of spice and is also considered as one of the most important ingredients in mustard industries [16]. B. nigra is non-mycorrhizal, and the presence of it decreases the soil biodiversity [17]. Brassica vegetables are greatly enriched with several nutritional values such as vitamin C and nutrients with anticancer properties including 3,3-diindolylmethane, sulforaphane and selenium [18]. According to Kiasalari et al, the seeds of Brassica nigra possesses essential oils (omega-3 fatty acids), the minerals (selenium, phosphorus, magnesium, manganese, iron, zinc and calcium), vitamins (A, C and B-complex), protein (dietry fibre) and phytoneutrients [19]. The phytochemical screening of seeds of Brassica nigra showed the presence of alkaloid, flavonoid, carbohydrate, fixed oil and glycosides [20]. It also showed the significant growth inhibitory activities against HeLa, HepG2, HCT, HEp2 and MCF-7 tumour cells [21]. Brassica vegetables contain indole-3-carbinol which inhibits the growth of cancer cell [22]. The seeds of Brassica are traditionally used in the treatment of cardiovascular diseases, neurotic pain, rheumatoid arthritis, brain and lung edema, paralysis, migraine, diabetes and its complications [19, 23]. It also belongs to antiepileptic and anxiolytic activity [20]. Furthermore, the leaves of B. nigra contain 3,3-diindolylmethane that is a potent modulator of the innate immune response system with potent antibacterial, antiviral and anticancer activity [24].

Considering the biomedical importance of B. nigra, we became interested to learn about the thrombolytic, anxiolytic, anticancer properties including 3,3-diindolylmethane, sulforaphane and selenium [18]. According to Kiasalari et al., the seeds of Brassica nigra possess essential oils (omega-3 fatty acids), the minerals (selenium, phosphorus, magnesium, manganese, iron, zinc and calcium), vitamins (A, C and B-complex), protein (dietry fibre) and phytoneutrients [19]. The phytochemical screening of seeds of Brassica nigra showed the presence of alkaloid, flavonoid, carbohydrate, fixed oil and glycosides [20]. It also showed the significant growth inhibitory activities against HeLa, HepG2, HCT, HEp2 and MCF-7 tumour cells [21]. Brassica vegetables contain indole-3-carbinol which inhibits the growth of cancer cell [22]. The seeds of Brassica are traditionally used in the treatment of cardiovascular diseases, neurotic pain, rheumatoid arthritis, brain and lung edema, paralysis, migraine, diabetes and its complications [19, 23]. It also belongs to antiepileptic and anxiolytic activity [20]. Furthermore, the leaves of B. nigra contain 3,3-diindolylmethane that is a potent modulator of the innate immune response system with potent antibacterial, antiviral and anticancer activity [24].

Considering the biomedical importance of B. nigra, we became interested to learn about the thrombolytic, anxiolytic, cytotoxic and anxiolytic activities of the flowers of B. nigra which is not still investigated. For this purpose, the study has been conducted.

Methods

Collection of plant materials

The flower of B. nigra (Brassicaceae) has been selected for the present study. For this investigation, B. nigra was collected from the local area of Noakhali, Bangladesh, in the month of March 2019 and was identified in Bangladesh National Herbarium (BNH), Dhaka, Bangladesh. A voucher specimen (DACB Accession No 45837) of the plant has been deposited in the herbarium.

Preparation of plant extract

The dried and powdered flowers (500 g) were soaked in 1500 ml of 99% methanol for about 21 days at room temperature with occasional stirring. After 21 days, the solution was filtered using filter cloth and Whatman filter paper. A rotary evaporator was used under reduced pressure at 40°C to obtain the concentrated methanol extract.
Solvent-solvent partitioning
The residue (40 g) derived from methanol extract was subjected to vacuum liquid chromatography using n-hexane, chloroform, ethyl acetate and water in order to increase polarity. The solvent-solvent partition of crude extract was performed according to Hossain et al. [14].

Experimental animal
For continuing the present experimentation, Swiss albino mice (both male and female) weighed between 20 and 25 g were used. These mice were collected from Jahangirnagar University, Dhaka, Bangladesh. Before the current research work, these animals were adjusted to standard animal house condition for 4 days [25]. Feeding of the animal was performed using standard laboratory pellet and water ad libitum, subjecting them to an alternate cycle of 12 h dark and light at relative humidity 55 ± 10% and temperature 25 ± 2 [25]. ‘Principles of Laboratory Animal Care’ (NIH publication no. 85-23, revised 1985) as well as specific national laws was followed by all authors.

Drugs and chemicals
Standard diazepam, dimethyl sulfoxide, vincristine sulphate (VS), albendazole and streptokinase were purchased from Square Pharmaceutical Ltd., Bangladesh. Other reagents of analytical grade for conducting this research work were supplied by the ethno pharmacology laboratory of the Pharmacy Department of NSTU.

Phytochemical evaluation
In order to determine the chemical groups present in the MCE, the preliminary phytochemical studies were conducted [26–28]. In each test, 10% (w/v) solution of extract in methanol was taken for the detection of presence of chemical groups.

Cytotoxicity by brine shrimp lethality bioassay
In brine shrimp lethality bioassay, dimethyl sulfoxide (DMSO) was used as both solvent and negative control whereas vincristine sulphate (VS) was applied as the positive control [14]. DMSO solutions of the test samples were subjected to Artemia salina for a day in order to determine in vivo cytotoxic screening. Four milligrams of the each sample was dissolved in DMSO, and solutions of different concentrations (400, 200, 100, 50, 25, 12.50, 6.25, 3.125, 1.563, 0.781 μg/ml) were attained by serial dilution technique. Then, mature shrimps (10 pieces) were kept in each of the experimental vials. The vials were observed after 24 h using a magnifying glass, and the number of subsisting nauplii in each vial was determined. Using these data, the percentage (%) of lethality of brine shrimp nauplii was calculated for each concentration by applying the following formula:

\[
\% \text{ of mortality} = \frac{N_t}{N_0} \times 100
\]

Here, \( N_t \) = the number of dead nauplii after 24 h of incubation and \( N_0 \) = the number of total nauplii transferred (10)

Then, the median lethal concentration (LC_{50}) was calculated from the log concentration versus percentage of mortality curve.

Anthelmintic assay
To study the anthelmintic activity, adult earthworms were used [10, 29]. The earthworms (Pheretima posthuma) were collected from moist soil of NSTU campus and washed with normal saline to eradicate all the soil. The earthworms 4–6 cm in length and 0.1–0.3 cm in width weighing 0.8–4.04 g were used for all experiment protocols. Both anatomically and physiologically, the earthworms were assimilated with the intestinal worm parasites of human beings, and for this reason, it was used to study the anthelmintic activity. Albendazole was treated as the reference standard.

Thrombolytic activity
The thrombolytic activity of B. nigra was conducted according to the method described by Prasad et al. and Islam et al. [10, 30]. Five milliliters of venous blood was taken from healthy volunteers (n = 5) and transferred to different pre-weighed sterilised micro-centrifuge tube (1 ml/tube). The micro-centrifuge tubes were subjected to incubation at 37 °C. After 45 min, the formation of clot occurred and serum was completely separated from the tubes (carried out without disturbing the clot formed) and the remaining clot of each tube was again weighed to calculate the weight of the clot [clot weight(W3) = weight of clot containing tube (W2) – weight of tube alone (W1)]. Each micro-centrifuge tube was labelled appropriately, and MCE and fractions with concentrations of 5 and 10 mg/ml were added to the tubes accordingly. One hundred microliters of streptokinase was used as a positive control whereas 100 µl of sterilised distilled water was distinctly added to the control tubes numbered as a negative non-thrombolytic control. After that, the tubes were incubated again at 37 °C for 90 min for observing clot lysis. Then, following the incubation, the obtained fluid was removed from the tubes and these fluids were again weighed to observe the difference in weight after clot disruption. Finally, difference obtained in weight was determined and the result was expressed as the percentage of clot lysis using the following equation:
% of clot lysis = \frac{W_3 - W_4}{\text{weight of clot}} \times 100

Here, \( W_3 \) = to weight (g) of clot after 45 min incubation and \( W_4 \) = the weight (g) of lysis clot after 90 min incubation.

Evaluation of anxiolytic activity

Elevated plus maze test

To evaluate the anxiolytic activity, the elevated plus maze was constructed according to the description given by the researchers [10, 31]. The EPM apparatus consists of two open arms (35 × 5 cm²) elapsed by two closed arms (35 × 5 × 15 cm³). The two arms were attached at one place with a central square of 5 × 5 cm². The apparatus was placed in a dimly enlightened room. The Swiss albino mice (20–25 g) were kept fast for overnight and split into seven groups; each group consists of 5 animals. Distilled water 10 ml/kg, i.p (negative control group); diazepam 1 mg/kg, i.p (positive control group); and plant extracts 200 and 400 mg/kg, i.p were administered to the individual groups of mice before the 60 min of test. The time spent and the number of entry in both open and close arms were recorded for 5 min [31].

Hole board test

The hole board apparatus used for anxiolytic test consists of 16 holes with a wooden chamber (40 × 40 × 25 cm³), and the diameter of each hole is 3 cm. The apparatus was raised to the height of 25 cm from the floor so that the mice could peep through the holes. The Swiss albino mice (20–25 g) were divided into the control, standard and test groups. The control group received distilled water 10 ml/kg, i.p and the standard group received diazepam 1 mg/kg, i.p while the test group received extract 200 and 400 mg/kg, i.p before the 60 min of test. Each mouse was kept at one corner of the board where the mouse could move freely and dip its head into the holes. The number of head dipping was recorded for 5 min [10, 32].

Acute toxicity test

Acute toxicity test was conducted according to Gandhare et al. [33]. The MCE of flowers of Brassica nigra was applied separately in various doses (50, 500, 1000, 2000 mg/kg) in four groups of animals by oral route. The animals were observed continuously for the first 2 h and 24 h to identify the behavioural changes. However, there are no noticeable behavioural changes and mortality occurred. Therefore, we considered the extract be safe at a dose level 2000 mg/kg and LD₅₀ be > 2000 mg/kg.

Statistical analysis

The data obtained in the studies were subjected to one-way analysis of variance (ANOVA) for determining the significant differences. The intergroup significance was analysed by using Dunnett’s t test. *\( p < 0.05 \), **\( p < 0.01 \) and ***\( p < 0.001 \) indicate weak, moderate and high significance, respectively. All the values were expressed as mean ± SEM (n = 5).

Results

Phytochemical screening

The phytochemical screening of MCE of B. nigra showed the presence of alkaloids, flavonoids, saponin, tannin, diterpenes, glycosides, carbohydrates, phenols fixed oils and fat which is exhibited in Table 1. Based on these results, the MCE of B. nigra was subjected to pharmacological investigations such as cytotoxic activity, anthelmintic activity, anxiolytic activity and thrombolytic activity. The study showed that the plant can be used for various purposes.

Cytotoxic investigation

The results of the toxicity of the MCE, CSF, ASF, EASF and n-HSF against brine shrimp (LC₅₀ values) are shown in Table 2. By using the process developed by Meyer, we determined LC₅₀ (lethal concentration, 50%) of MCE and its four fractions. From the results of the brine shrimp lethality bioassay, it is observed that the CSF demonstrated greater toxicity (1.563 μg/ml) compared with others and ASF exerted the lowest percentage of mortality. So it can nicely be predicted that the MCE and its fractions possess cytotoxic properties compared with positive control VS. However, no mortality was observed for the negative control group.

Table 1 Preliminary phytochemical analysis of B. nigra flowers

| Phytochemicals | Comment |
|----------------|---------|
| Alkaloid       | ++      |
| Saponin        | +       |
| Tannin         | ++      |
| Flavonoids     | ++      |
| Carbohydrate   | ++      |
| Diterpenes     | +       |
| Protein        | −       |
| Phytosterols   | +       |
| Glycosides     | +       |
| Phenol         | ++      |
| Fixed oils and fat | ++ |

Here, ‘+’ means present, ‘++’ means strongly present and ‘−’ means absent.
Anthelmintic activity

The result of anthelmintic activity is shown in Table 3. From the result, it was seen that all fractions cause the death and paralysis of *Phertima posthuma* in a dose-dependent manner which is significant (*p* < 0.05) when compared with standard drug albendazole. Here, n-HSF at the concentration 10 mg/ml showed paralysis and death at 13.20 ± 0.58 min and 22.40 ± 1.36 min, respectively. Again, time taken for paralysis and death was 8.80 ± 0.97 min and 15.60 ± 1.08 min, respectively, at a concentration of 20 mg/ml for n-HSF. Moreover, in case of albendazole (20 mg/ml), the paralysis time was 20.80 ± 1.28 min whereas death time was 54.00 ± 1.45 min. Thus, other fractions also exhibited significant anthelmintic activity in a dose-dependent fashion. Here, n-HSF demonstrated higher anthelmintic activity against the earthworm.

Anxiolytic activity

In the EPM test, the MCE and its fractions at dose 200 and 400 mg/kg showed (Table 4) an increase of time spending in open arm in a dose-dependent manner which indicates the anxiolytic activity of the plant [10]. All the fractions exhibited good anxiolytic activity but n-HSF showed relatively good activity as compared to standard. Times spent by mice in open and closed arm for n-HSF at dose 200 mg/kg were 72.20 ± 2.71 s and 192.80 ± 2.35 s while it was 80.80 ± 1.46 s and 183.00 ± 2.43 s at dose 400 mg/kg, respectively. On the other hand, time spent by mice in open and close arm for standard was 102.20 ± 3.71 s and 170.00 ± 3.49 s, respectively. Again the number of entries in open arm was also increased with the increase of dose whereas it was decreased in case of close arm. In the hole board model, the number of head dipping was measured (Table 5). The result showed significant increase in head dipping counts with the increase of dose which is significant when compared with standard and indicates the presence of anxiolytic activity.

Thrombolytic activity

The MCE of *B. nigra* flowers was assessed for thrombolytic activity, and the results are presented in Table 6. The addition of 100 μl streptokinase as a positive control showed 46.39% clot lysis. On the other hand, distilled water was treated as a negative control which exhibited a negligible percentage of lysis of clot (6%). Among the different fractions, the highest percentage of clot lysis activity was 30.77 ± 4.09% and 36.89 ± 1.94% at the concentration of 5 and 10 mg/ml, respectively, for CSF. Other fractions also exhibited potentiality of clot lysis at a dose-dependent manner which was significant when compared with the negative control (*p* < 0.05).

### Table 2

| Sample          | LC50 (μg/ml) | Regression equation | R²  |
|-----------------|--------------|---------------------|-----|
| VS              | 0.709        | y = 39.765x + 55.949 | 0.96|
| MCE             | 5.586        | y = 30.709x + 27.058 | 0.8848|
| n-HSF           | 3.019        | y = 25.912x + 37.516 | 0.8028|
| CSF             | 1.563        | y = 22.029x + 45.724 | 0.9454|
| EASF            | 3.730        | y = 17.37x + 40.069  | 0.9291|
| ASF             | 10.168       | y = 21.394x + 28.451 | 0.9739|

*VS* vincristine sulphate, *MCE* methanol crude extract, *EASF* ethyl acetate soluble fraction, *n-HSF* n-hexane soluble fraction, *CSF* chloroform soluble fraction, *ASF* aqueous soluble fraction

### Table 3

| Sample          | Concentration (mg/ml) | Time (min) | Death time |
|-----------------|-----------------------|------------|------------|
|                 |                       | Paralysis  |            |
|                 |                       |            |            |
| Standard (albendazole) | 10                     | 35.40 ± 0.51 | 73.40 ± 1.21 |
|                 |                       | 20.80 ± 1.28 | 54.00 ± 1.45 |
| CSF             | 10                     | 17.80 ± 0.58*** | 30.80 ± 1.11*** |
|                 | 20                     | 11.80 ± 1.02**  | 22.00 ± 1.00*** |
| ASF             | 10                     | 87.60 ± 0.51*** | 99.40 ± 0.81*** |
|                 | 20                     | 54.40 ± 1.29*** | 67.60 ± 2.42** |
| MCE             | 10                     | 53.60 ± 0.81*** | 62.20 ± 0.97** |
|                 | 20                     | 33.80 ± 1.24**  | 47.40 ± 1.57*** |
| n-HSF           | 10                     | 13.20 ± 0.50*** | 22.40 ± 1.36*** |
|                 | 20                     | 8.80 ± 0.97**   | 15.60 ± 1.08*** |
| Control (1% CMC in normal saline) | – | – | – |

Each value represents the mean ± SEM (*n* = 5)

* MCE methanol crude extract, *EASF* ethyl acetate soluble fraction, *n-HSF* n-hexane soluble fraction, *CSF* chloroform soluble fraction, *SEM* standard error mean

The inter group significance was analysed by one-way ANOVA using Dunnet’s t-test. *P* < 0.05*, *P* < 0.01** and *P* < 0.001*** indicates weak, moderate and high significance respectively.
Discussion

Human beings are relying on plants for the treatment of many diseases from the very beginning of civilization [34]. At present, phyto-pharmacological study has opened a new area to discover plant derivative drugs which are effective for the treatment of certain diseases and draw the attention in herbal medicines [34]. About 30% of pharmaceuticals are thought to be prepared from plant derivatives [35].

A number of studies have been conducted to explore the safe and effective drugs in order to address the treatment of cancer or tumour. Here, we tried to find out any ingredient that may lie in MCE and may be effective in cancer or tumour through cytotoxic investigation. In this investigation, varying degree of lethality was found with exposure to different dose levels of the test samples. The degree of lethality was found, which is directly proportional to the concentration gradient which indicates that mortality increases gradually with the increase of concentration of the test samples. The MCE and its fractions exhibited (Table 1) significant toxicity to brine shrimp as the LC50 values were less than

Table 4 Effects of MCE and its fractions of *B. nigra* flowers on mice in the open and closed arm of the EPM

| Sample  | Doses (mg/kg) | Time spent in seconds (mean ± SEM) | Number of entries (mean ± SEM) |
|---------|---------------|------------------------------------|-------------------------------|
|         |               | Open arm | Closed arm | Open arm | Closed arm |
| Control | 10            | 19.40 ± 2.64 | 254.00 ± 2.07 | 7.00 ± 0.71 | 28.33 ± 2.03 |
| Standard| 1             | 102.20 ± 3.71 | 170.00 ± 3.49 | 18.60 ± 0.51 | 7.33 ± 1.45 |
| n-HSF   | 200           | 72.20 ± 2.71** | 192.80 ± 2.35** | 13.60 ± 0.81** | 6.40 ± 0.51*** |
|         | 400           | 80.80 ± 1.46* | 183.00 ± 2.43 | 14.80 ± 0.80 | 8.68 ± 2.11*** |
| CSF     | 200           | 59.80 ± 4.53** | 205.00 ± 1.87** | 11.40 ± 0.68** | 7.60 ± 0.51*** |
|         | 400           | 72.60 ± 2.01** | 195.40 ± 1.75** | 13.00 ± 0.71** | 9.62 ± 0.61*** |
| EASF    | 200           | 43.20 ± 1.43** | 230.60 ± 1.75*** | 7.60 ± 0.51*** | 10.60 ± 0.51*** |
|         | 400           | 48.80 ± 1.46*** | 224.20 ± 1.88*** | 9.60 ± 0.60*** | 12.43 ± 0.60*** |
| MCE     | 200           | 59.20 ± 0.97** | 210.20 ± 1.85** | 10.60 ± 0.51** | 11.40 ± 0.68*** |
|         | 400           | 66.80 ± 1.50** | 199.60 ± 2.50*** | 12.40 ± 0.60** | 13.12 ± 0.71*** |
| ASF     | 200           | 38.60 ± 0.93*** | 234.80 ± 2.42*** | 6.40 ± 0.51*** | 13.60 ± 0.81** |
|         | 400           | 44.00 ± 2.10*** | 226.40 ± 0.67**** | 8.60 ± 2.10*** | 14.80 ± 0.80* |

Each value represents the mean ± SEM (n = 5)
MCE methanol crude extract, EASF ethyl acetate soluble fraction, n-HSF n-hexane soluble fraction, CSF chloroform soluble fraction, ASF aqueous soluble fraction, SEM standard error mean
The inter group significance was analysed by one-way ANOVA using Dunnet’s t-test. P < 0.05*, P < 0.01** and P < 0.001*** indicates weak, moderate and high significance respectively

Table 5 Effects of MCE and its fractions of *B. nigra* flowers on mice in hole board test

| Sample  | Doses (mg/kg) | Number of head dipping (300 s test) in hole board (mean ± SEM) |
|---------|---------------|-------------------------------------------------------------|
| Control | 10            | 36.40 ± 3.26                                                |
| Standard| 1             | 39.80 ± 2.22                                                |
| n-HSF   | 200           | 36.40 ± 3.26                                                |
|         | 400           | 64.20 ± 1.77***                                             |
| CSF     | 200           | 72.20 ± 2.71***                                             |
|         | 400           | 71.60 ± 2.20***                                             |
| EASF    | 200           | 81.20 ± 2.60***                                             |
|         | 400           | 46.00 ± 1.41                                                |
| MCE     | 200           | 57.00 ± 1.22**                                              |
|         | 400           | 63.60 ± 1.72***                                             |
| ASF     | 200           | 75.20 ± 1.71**                                              |
|         | 400           | 45.00 ± 3.21                                                |

Each value represents the mean ± SEM (n = 5)
MCE methanol crude extract, EASF ethyl acetate soluble fraction, n-HSF n-hexane soluble fraction, CSF chloroform soluble fraction, ASF aqueous soluble fraction, SEM standard error mean
The inter group significance was analysed by one-way ANOVA using Dunnet’s t-test. P < 0.05*, P < 0.01** and P < 0.001*** indicates weak, moderate and high significance respectively
Table 6 Effects of different fractions of MCE of B. nigra flowers on in vitro clot lysis

| Sample       | Dose (mg/ml) | Percentage of clot lysis (mean ± SEM) |
|--------------|--------------|--------------------------------------|
| Blank        | 6 ± 1.09     |                                      |
| Streptokinase| 46.39 ± 2.92 |                                      |
| n-HSF        | 5            | 18.92 ± 2.94**                       |
|             | 10           | 22.78 ± 2.05***                      |
| CSF          | 5            | 30.77 ± 4.09                         |
|             | 10           | 36.89 ± 1.94***                      |
| ASF          | 5            | 10.92 ± 2.32**                       |
|             | 10           | 17.32 ± 2.48*                        |
| MCE          | 5            | 9.59 ± 1.82***                       |
|             | 10           | 21.59 ± 1.52***                      |

Each value represents the mean ± SEM (n = 9)  
MCE methanol crude extract, EASF ethyl acetate soluble fraction, n-HSF n-hexane soluble fraction, CSF chloroform soluble fraction, SEM standard error mean  
The inter group significance was analysed by one-way ANOVA using Dunnet’s t-test.  
* P < 0.05, ** P < 0.01 and *** P < 0.001 indicates weak, moderate and high significance respectively

100 μg/ml [14]. From the LC50 values, we conclude that the CSF (1.563 μg/mL) contains more potent cytotoxic compounds compared to the other fractions. On the other hand, the ASF (10 μg/ml) showed relatively less toxicity. There was no mortality in the negative control groups indicating the test was a valid one. Various studies revealed that plant extract shows cytotoxicity due to the presence of phytochemical compounds like flavonoids, glycosides, tannins, saponins and alkaloids which have been observed in this study [36, 37]. Saponins exert their cytotoxic activity through the apoptosis pathway which is a process of programme cell death [38]. Another study reported that saponins work by stopping cellular mutation that leads to cancer [30]. It is believed that tannins exert cytotoxicity by blocking induction of enzyme needed for cancer cell line growth [36]. Again, flavonoids exhibit cytotoxicity through increasing the production of intracellular ROS level [39] whereas alkaloids act by inhibiting the proliferation of a number of tumour cells [40]. These phytochemicals have been noticed in our study. Therefore, we can claim that the positive response obtained in our investigation suggests that the plants may contain cytotoxic agents which could be developed for medicinal use.

In anthelmintic activity test, all the fraction of extract provoke not only paralysis but also death of the earthworms in dose-dependent manner. From the study, we observed that the paralysis and death time of earthworm were inversely proportional to the extract concentrations. Among the fractions, n-HSF showed less paralysis (13.20 ± 0.58) and death (22.40 ± 1.36) time which was potential when compared with standard (35.40 ± 0.51 and 73.40 ± 1.21, respectively). Several studies claimed that tannins, flavonoids, alkaloids and phenolic compounds are accountable for anthelmintic activity [41, 42]. Tannins can bind to free proteins in the gastrointestinal tract of host animal or glycoprotein on the cuticle of the parasite and thereby cause deaths [41, 42]. Tannins may exhibit their activity by intervening with energy production of worms through uncoupling oxidative phosphorylation that leads to death [42]. On the other hand, alkaloids induce paralysis of worms by acting on its central nervous system [42]. Our phytochemical screening exhibited the presence of alkaloids, flavonoids, tannins and phenolic compounds. Hence, we can consider the flower of B. nigra as an alternative source of anthelmintic drugs.

In case of thrombolytic test, it can be demonstrated that our findings may have significant implications in cardiovascular health. A widely used thrombolytic agent streptokinase acts by converting additional plasminogen to plasmin. But this agent has several adverse effects which encouraged the researcher to discover alternative agent [43]. Therefore, we tried to find out whether the plant possesses clot lysis property or not. The comparative study between positive and negative control clearly showed that clot lysis did not occur when water was added to the clot. On the other hand, addition of different fractions of extract revealed a significant clot lysis. Among the different fractions, CSF showed highest clot lysis (30.77 ± 4.09). Several studies reveal that tannins, alkaloids and saponins are responsible for clot lysis activity [44, 45]. It is assumed that these phytochemicals exert their activity by disrupting the fibrinogen and fibrin in a clot that ultimately leads to fibrinolysis [44, 45]. As phytochemical analysis revealed that the crude extract contains tannins, alkaloids and saponins, it may predict that these phytochemicals may be responsible for its clot lysis activity.

Elevated plus maze and hole board were used to estimate the psychomotor performance and emotional aspects of mice. Moreover, we know that anxiolytic agents raise the time spent and a number of entries in open arms of the EPM [10, 46]. The result of the study showed that different fractions of plant extract at both dose 200 mg/kg and 400 mg/kg treated into mice showed significant increases in time spent and the number of entries in open arms in comparison to the control group. On the other hand, both the time spent and number of entry decreased in close arms in comparison to the control group. Again, in case of hole board test, the head dipping of animals is inversely proportional to their anxiety state in the moderately aversive environment [10, 47]. Therefore, an increasing number of head dipping of mice into the holes of board indicate declined anxiety state. Previous investigations suggested that alkaloids, flavonoids and terpenoids are accountable for anxiolytic activity which was observed in our study [10]. Flavonoids induce anxiolytic effect by
opening activated chloride channel to increase permeability of neuronal membrane chloride ions that results in GABA’s inhibitory effect [10]. Again, alkaloids such as montanine initiate anxiolytic activity by reducing the locomotor activity [48]. Thus, we may conclude that the methanol extract of \textit{B. nigra} flowers and its fractions have satisfactory anxiolytic potential.

From the light of above study, it can easily be mentioned that the methanol extract and its different fractions of \textit{B. nigra} flowers are a vital source of therapeutic agents having the thrombolytic, anthelmintic, anxiolytic and cytotoxic potentiality.

**Conclusion**

On the basis of the findings of the present study, it can be determined that methanol extract of \textit{Brassica nigra} flowers possesses different types of pharmacological activities in varying concentration and fractions. At the end, it can be concluded that the experimental evidence obtained in the laboratory test model could provide a rationale for the traditional use of this plant as an anthelmintic, thrombolytic, anti-tumour and anxiolytic agent. Future scope involves the isolation of phyto-constituents that are responsible for anthelmintic, thrombolytic, cytotoxic and anxiolytic activity and the study of its pharmacological actions.

**Abbreviations**

TPA: Tissue plasminogen activator; BNH: Bangladesh National Herbarium; VS: Vincristine sulphate; MCE: Methanol crude extract; EASF: Ethyl acetate soluble fraction; n-HSF: n-Hexane soluble fraction; CSF: Chloroform soluble fraction; LCO\textsubscript{50}: Lethal concentration 50; ROS: Reactive oxygen species; SEM: Standard error mean; DNA: Deoxyribonucleic acid; GABA: Gamma-amino-butyric acid; DMSO: Dimethyl sulfoxide; EPM: Elevated plus maze

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**Source of the plant**

The plant was collected from the local area of Noakhali.

**Authors’ contributions**

We declare that this study was conducted by the authors named in this article: MSU, MF and MSM designed the study. MSH, MGU and SAS carried out the laboratory work, analysed the data and wrote the manuscript. MSU and MGU helped to supervise the work and collaborated in the data analysis while MSM revised and corrected the manuscript. All authors read and approved the final manuscript.

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**Availability of data and materials**

All data and material are available upon request.

**Ethics approval and consent to participate**

The ethical approval for conducting the study was taken from the ethical approval committee of Noakhali Science & Technology University. The reference number is 03/2019.

**Consent for publication**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

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