Original Research

**Citrus limon** L. (lemon) seed extract shows neuro-modulatory activity in an *in vivo* thiopental-sodium sleep model by reducing the sleep onset and enhancing the sleep duration

Md. Momnir Rahman¹, Fahadul Islam¹, Anwar Parvez¹, Md. A.K. Azad¹, Chulam Md Ashraf²,³, Mohammad Fahad Ullah⁴,⁵, Muniruddin Ahmed¹,⁶

¹ Department of Pharmacy, Faculty of Allied Health Sciences, Daffodil International University, 1207 Dhaka, Bangladesh
² Pre-Clinical Research Unit, King Fahd Medical Research Center, King Abdulaziz University, 21589 Jeddah, Saudi Arabia
³ Department of Medical Laboratory Technology, Faculty of Applied Medical Sciences, King Abdulaziz University, 21589 Jeddah, Saudi Arabia
⁴ Prince Fahd Research Chair, Department of Medical Laboratory Technology, Faculty of Applied Medical Sciences, University of Tabuk, 47311 Tabuk, Saudi Arabia

*Correspondence: drmuniruddin@gmail.com (Muniruddin Ahmed); m.ullah@ut.edu.sa (Mohammad Fahad Ullah)

DOI: 10.31083/j.jin2101042

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Submitted: 24 May 2021 Revised: 28 July 2021 Accepted: 14 September 2021 Published: 28 January 2022

**Citrus limon** L. is an ingenious alternative medication and has a broad scope in managing several health conditions as part of natural remedies. Recently, medicinal plants have witnessed incredible consideration worldwide in the field of neuroscience for remedial intervention. The present work has investigated the phytochemical compounds and neuropharmacological potential of the seed extract of *Citrus limon* as a step to partially validate its formulations as nutraceuticals using an *in vivo* model. Diverse phytochemical groups such as alkaloids, flavonoids, tannins, gums, saponins, steroids were qualitatively identified through colorimetric methods utilizing standard compounds. The neuropharmacological properties were studied in Swiss albino mice with the sleep time induced by thiopental sodium taken as an end-point, in standard hole cross, hole board, and open-field experiments at varying doses of 50 and 100 mg/kg body weight. Phytochemical screening showed that alkaloids, flavonoids, saponins, tannins, steroids, and glycosides are present in the aqueous extract of the seed. The extracts demonstrated a significant reduction in sleep onset and enhanced the sleep duration in a dose-dependent manner in thiopental sodium-induced sleeping time, along with a marked decrease in unconstrained locomotors and explorative properties in both hole cross and open field tests. Moreover, in the hole board study, the extracts minimized the count of head dips observed in the treated mice. The results shown in this study demonstrate that *Citrus limon* extracts have neuropharmacological properties that can be further examined for their potential role as an adjuvant with conventional medications or nutraceuticals.

**Keywords**

Neurological disorders; Phytochemical compounds; Latent period; Neuropharmacological potential; Nutraceuticals

1. Introduction

Recently, a stressed lifestyle has been linked to an assortment of psychiatric disorders, and evidence suggests that depression is an unavoidable medical concern in these disorders [1–4]. Different antipsychotic drugs, namely, tricyclic antidepressants, monoamine oxidase inhibitors, and serotonin re-uptake inhibitors, are available to manage various psychotic problems, but several side effects are gradually hindering their use in a long-term therapeutic regimen [5, 6]. Thus, the quest for new anxiolytic substances with minimal unfavorable impact is still an area of interest being explored for an appropriate pharmacological effect. Certain organic substances, particularly restorative herbs, are commonly known to be the primary armamentarium of concoction compounds with enormous remedial potential. Neuropsychiatric disorders such as depression are common, corresponding to a state of mental illness that may result in severe symptoms such as sadness, feelings of guilt and loss of interest which affect sleeping, dietary intake, cognitive and psychomotor guided daily activities. Depression and anxiety, if they remain untreated, may aggravate the symptoms, worsen the complications, and ultimately lead to suicidal ideation risks [7, 8]. The actual etiology of depression and anxiety is still unknown. However, these may arise due to continued disruption of the antioxidant defense system and an enhanced redox imbalance or oxidative stress [9]. The redox stress in the brain supports cellular conditions that cause neurological damage and related cognitive deteriorations [10–13].

*Citrus limon* belongs to the Rutaceae family, with a predominant presence in South Asia, including Northeastern India. *Citrus limon* is the most notable citrus species, with an affluent mercantile value. Most types of citrus plants are substantial evergreen bushes or small trees, 5–15 meters tall. The tree’s ellipsoidal yellow fruit is used worldwide for culinary and non-culinary purposes and is an active ingredient of folk medicine and functional foods. The lemon juice is around 5 to 6 percent citrus extract, having an acrid flavor with a
pH of about 2.2. The distinctively harsh flavor of lemon juice makes it an essential constituent of drinks and nutritional supplements, including lemonade and lemon meringue pie. Its berry is essentially rich in alkaloids which show anticancer properties and demonstrates antibacterial activities in unrefined concentrations of lemon on clinically challenging bacterial strains [14–21]. The biological activity of C. limon is attributed to the rich quantity of phenolic compounds, for instance, flavonoids; limonoids, carotenoids and bioactive monoterpenoids such as D-limonene, β-pinene and γ-terpinene [22]. These chemical compounds are the factor for the significant utilization of C. limon in the food and cosmetics industries [23–28]. The scientifically proven therapeutic activities of C. limon include anti-inflammatory, antimicrobial, antioxidant, anticancer, and antiparasitic actions. Citrus flavonoids may also modulate blood platelet functions and have an anti-aggregatory effect [29, 30]. To further explore the pharmacological properties of C. limon, the current investigation focused on the neuro-pharmacological benefits of this plant in the in vivo mouse model.

2. Materials and methods

2.1 Chemicals and drugs

Thiopental Sodium and diazepam were purchased from Incepta Pharmaceuticals Ltd (Dhaka, Bangladesh) and Opsonin Pharma Ltd (Dhaka, Bangladesh). Distilled water and different reagents were obtained from BDH Chemicals Ltd (Dhaka, Bangladesh).

2.2 Plant materials

The seeds were obtained from 45–50 completely ripened fruits of the 4–5 Citrus limon plants, which were mature and aged 2–2.5 years, grown in fields close to Jahangirnagar University in Dhaka, Bangladesh, on 23rd October 2020. These were authenticated from Bangladesh National Herbarium, Mirpur, Dhaka.

2.3 Preparation of plant extract

The seeds collected from Citrus limon were carefully washed with distilled water to remove undesired natural parts. The seeds of the plants were dried under sunlight (temperature approximately 30–35 °C) for two days and stored for seven days in an air-tight container. The dried seeds were ground to a coarse powder. The prepared fine powder was washed with distilled water to remove undesired natural parts. These seedsof the plants weredried undersunlight (temperature approximately 30–35 °C) for two days and stored for seven days in an air-tight container. The dried seeds were packed in a container and stored in cool, dark, and dry conditions to prepare fresh extracts before the investigation. A quantity of 500 gm of seed powder of Citrus limon was dissolved in 1000 mL of water in a glass compartment for 2–3 days, with standard shaking and mixing. Afterward, the entire mixture was passed through a fine, white cotton cloth. Further filtration was done using Whatman filter paper. Extracts were obtained by the evaporation of solvent that was kept in open space. The yield value of the extracts from the lemon seeds was 2.65% w/w [31].

2.4 Experimental animals

Ninety Swiss albino mice (22–25 g) aged 6–7 weeks were obtained from Jahangirnagar University, Dhaka, Bangladesh. The animals were caged under standard natural conditions (22–25 °C, humidity 60–70%, 12-hour light: 12-hour dull cycle). These were given a standard pellet diet, which was obtained from Jahangirnagar University, Dhaka. All protocols considered in our study, including animals, were approved by the Faculty of Allied Health Sciences Research Ethics Committee, Daffodil International University, Dhaka-1207, Bangladesh. (Ref: FAHSREC/DIU/2020/1006).

2.5 Qualitative phytochemical screening

Qualitative assessment was done using colorimetric procedures with standard substances, indicating specific phytochemical groups, for example, glycosides, flavonoids, tannins, alkaloids, gums, steroids, saponins [32]. For carbohydrates, Molisch and Fehling’s tests were used. Identification of flavonoids was based on the standard flavonoid test. The Dragendorf’s, Mayer’s, and Hager’s tests were used for alkaloids. Potassium dichromate test, ferric chloride, and lead acetate derivation tests were pursued for tannins. For glycosides, Keller-Kiliani tests were performed. A frothing test was done for the presence of saponins. To detect steroids, the sulphuric acid analysis was performed. For detecting the presence of gum in the extracts, the Molisch test was further used.

2.6 Quantitative determination of some phytochemical constituents

Harborne’s (1973) method was used to analyze alkaloids: 5 g of the sample was weighed into a 250 mL beaker, and 200 mL of 10% acetic acid in ethanol was added, capped, and set aside for 4 hours. This was filtered, and the extract was concentrated to one-quarter of its original volume using a water bath. Dropwise additions of concentrated ammonium hydroxide to the extract were made until the precipitation was accomplished. The entire solution was allowed to settle and rinsed with weak ammonium hydroxide before being filtered. The residual alkaloid content was desiccated and weighed. Van-Burden and Robinson’s (1981) method was used to determine tannin: A 50 mL plastic bottle filled with 500 mg of the sample. In a mechanical shaker, 50 mL of distilled water was added and agitated for 1 hour. This was then filtered into a 50 mL volumetric flask and brought up to the required concentration. The filtrate was then pipetted into a test tube with 2 mL of 0.1 M FeCl₃ in 0.1 N HCl and 0.008 M potassium ferricyanide. Within 10 minutes, the absorbance was taken at 120 nm. Bohm and Kocipai Abyazan’s (1994) procedure was used for flavonoid determination: At room temperature, 10 g of the plant sample was extracted many times with 100 mL of 80 % aqueous methanol. Whatman filter paper No 42 was used to filter the entire solution (125 mm). The filtrate was then placed in a beaker and dried over a water bath before being weighed at a consistent weight [33–35].
2.7 Sleeping time induced by thiopental sodium

The technique defined by Raihan MO et al. [36] was used to study the impact of the lemon seed extracts on sleeping time experiments induced by thiopental sodium. For such an objective, mice were grouped into six groups, with five \((n = 5)\) mice in each group. Group I was used as a control and given distilled water (10 mL/kg b.w.), group II, as usual, diazepam (0.50 mg/kg b.w.) was used as standard and groups III, IV, V and VI were used for the seed extracts. Group II, III, IV, V and VI were administered orally using cannula, the standard drug diazepam (0.50 mg/kg b.w.) and seed extracts in the doses: 25, 50, 100 and 200 mg/kg b.w. After thirty minutes, thiopental sodium (20 mg/kg b.w.) was injected intraperitoneally to all assortments to induce sleep. For monitoring adverse effects (if any), single mice from each group were placed on the table and examined. The length of sleep or monitoring adverse effects (if any), single mice from each group were placed on the table and examined. The length of sleep induced by thiopental sodium in animals was determined as the percentage increase in thiopental sodium-induced sleep time dependent on the dose (Table 3). The percentages of effects were calculated using the following equation:

\[
\text{Effect (\%)} = \frac{\text{The average duration of loss of righting reflex in the test group}}{\text{The average duration of loss of righting reflex in the control group}} \times 100
\]

2.8 Hole cross test

This investigation was carried out as described in the past by Uddin et al. [37]. A cage was utilized that has a size of 35 × 25 × 20 cm. In the enclosure, a segment was marked. At the height of 7.5 cm at the enclosure’s central focus, a hole with a diameter of 3 cm was created. Mice were administered with control, standard, or extract treatment and were put on one side of the platform. The number of passages taken by the mouse in each group through the hole from one chamber to another was then counted for 3 min at intervals 0, 30, 60, 90, and 120 min after administering the control, standard, and test sample (p.o.). This experiment was conducted within two days. Group I was set as control and given distilled water (10 mL/kg b.w.), and group II had taken diazepam (1 mg/kg, b.w.), which was used as standard. Groups III and IV were given seed extracts of 50 and 100 mg/kg body weight, respectively.

\[
\text{Inhibition (\%)} = \frac{\text{Mean No. of movements (control)} - \text{Mean No. of movements (test)}}{\text{Mean No. of movements (control)}} \times 100
\]

2.9 Hole board test

The test was performed as described by Kamei J et al. [38] with slight deviations. For the present test, a level platform of 90 cm × 90 cm in radius with 16 equivalently separated holes was utilized. This stage likewise had a frame of 5 cm in height. Mice were grouped into four groups; control, standard, and test. Five \((n = 5)\) mice were in each group. Group 1 (10 mL/kg b.w.) was listed for control and took distilled water. As standard, diazepam was provided to group II (1 mg/kg, b.w., p.o.). Groups III and IV independently took seed extracts in 50 and 100 mg/kg b.w. doses. The number of head dips in single mice into the holes was monitored for 10 min after the application.

\[
\text{Inhibition (\%)} = \frac{\text{Mean No. of head dips (control)} - \text{Mean No. of head dips (test)}}{\text{Mean No. of head dips (control)}} \times 100
\]

2.10 Open field test

The method described by Gould TD et al. [39] was used for this analysis. The test device is made up of a plane 0.5 m² field with a square progression. The squares on the other side are painted black and white. The experimental board is identical to a chessboard. Likewise, the mechanical system had a compartment height of 0.1 m. Mice were aligned into four groups. There were five \((n = 5)\) mice in every group. Group I was assessed as control and given distilled water (10 mL/kg b.w.). Diazepam (1 mg/kg, b.w., p.o.), which was deemed standard, was given to Group II. Seed extract doses of 50 and 100 mg/kg b.w. were distinctly provided to Groups III and IV. The number of squares moved at any pace by the animals in each treated group was measured for 3 min at the time intervals 0, 30, 60, 90, and 120 min after the oral administration of test medication.

\[
\text{Inhibition (\%)} = \frac{\text{Mean No. of movements (control)} - \text{Mean No. of movements (test)}}{\text{Mean No. of movements (control)}} \times 100
\]

2.11 Statistical analysis

Results are shown as mean ± SEM. Analyzes were carried out using one-way ANOVAs followed by Dunnett’s post hoc tests for sleeping time and hole board data. Two-way ANOVAs followed by Bonferroni’s post hoc tests were used for hole cross and open-field data. As a statistical result, \(p < 0.05\) was reported as significant.

3. Results

3.1 Aqueous seed extract shows a variety of bioactive phytochemicals

A comprehensive range of chemical tests for identifying major classes of therapeutically significant compounds exhibited alkaloids, flavonoids, tannins, steroids, and cardiac glycosides in the aqueous extract of the Citrus limon seed. In contrast, saponins were found to be absent. Phytochemical compounds of Citrus limon aqueous extract of seeds are shown in Table 1. Quantitative estimation for certain phytoconstituents such as alkaloids, flavonoids and tannins in Citrus limon seed extract is also summarized in Table 2. The estimations found tannins to be present in good proportion compared to other molecules.

3.2 Lemon seed extract induces neuromuscular sedation in the thiopental Na —sleep model

In the thiopental-induced hypnosis test, the plant extracts have shown a significant reduction in sleep onset and enhanced sleep duration in a dose-dependent manner. In this study, seed extract at doses of 25, 50, 100, and 200 mg/kg demonstrated a substantial decrease in sleep onset and increased sleeping time dependent on the dose (Table 3/\(p < 0.05\)). The length of thiopental sodium-induced sleep time
Table 1. Phytochemical compounds of *Citrus limon* (lemon) seed extracts.

<table>
<thead>
<tr>
<th>Compounds</th>
<th>Seed (aqueous extracts)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alkaloids</td>
<td>++</td>
</tr>
<tr>
<td>Flavonoids</td>
<td>+</td>
</tr>
<tr>
<td>Saponins</td>
<td>-</td>
</tr>
<tr>
<td>Tannins</td>
<td>+</td>
</tr>
<tr>
<td>Steroids</td>
<td>++</td>
</tr>
<tr>
<td>Gums</td>
<td>+</td>
</tr>
<tr>
<td>Cardiac glycosides</td>
<td>+</td>
</tr>
</tbody>
</table>

All tests were carried out in triplicates.

**KEY**: ++, Highly detected; +, Less detected; -, Not detected.

Table 2. Percentage of alkaloids, tannin and flavonoids in the *Citrus limon* seed extract.

<table>
<thead>
<tr>
<th>Phytochemical class</th>
<th>Quantity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alkaloids</td>
<td>0.36 ± 0.2</td>
</tr>
<tr>
<td>Flavonoids</td>
<td>0.34 ± 0.10</td>
</tr>
<tr>
<td>Tannins</td>
<td>12.14 ± 0.12</td>
</tr>
</tbody>
</table>

The values are represented as the Mean ± SD.

and its latency in experimental animals was effectively modulated by varying doses of the extract relative to controls (Figs. 1 and 2). As shown, the seed extract at 200 mg/kg had a maximal effect of 704% for lacking resolving reflex, while diazepam 0.50 mg/kg had a 529% effect (Table 3).

Fig. 1. The effect of *Citrus limon* (lemon) seed extracts on thiopental-Na stimulated sleeping time in mice (n = 5).

Fig. 2. The effect of *Citrus limon* (lemon) seed extracts on thiopental-Na induced latent period on mice (n = 5).

3.3 Central nervous system (CNS) depressant activity observed in animals treated with lemon seed extract as indicated by hole cross method

In the hole cross arrangement, the number of holes crossed from one compartment to another was recorded in intervals ranging from 30 to 120 min. It was observed that the seed extracts at 50 and 100 mg/kg doses caused a reduction of activity in the animals (Fig. 3). Robust ($p < 0.05$) data have been obtained in dose-dependent terms (Table 4). In this study, evaluation at the 5th period demonstrated a median of 69% suppression of locomotors activity by seed extract at 100 mg/kg. In the same experimental set-up, diazepam demonstrated 53% suppression of the activity.

Fig. 3. CNS depressant activity of *Citrus limon* (lemon) seed extracts by hole cross method (n = 5).

3.4 CNS depressant activity induced by lemon seed extracts in treated animals as indicated by hole board method

The evaluation of CNS depressant activity by seed extracts provided some interesting results based on the dosage ($p < 0.05$). The extract resulted in significantly minimizing of head-dips in the treated animals in the hole-board test (Fig. 4). The 100 mg/kg dose of seed extract resulted in the suppression of movement (72%), which was stronger than the standard diazepam (67%) (Table 5).

Fig. 4. CNS depressant activity of *Citrus limon* (lemon) seed extracts by hole board method (n = 5).
Table 3. The effects of *Citrus limon* (lemon) seed extracts on the sleeping time of mice induced by thiopental-Na.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dose (mg/kg)</th>
<th>Latent period (min)</th>
<th>Duration of sleep (min)</th>
<th>% Effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>10 mL/kg</td>
<td>9.6 ± 0.92</td>
<td>37.4 ± 1.4</td>
<td>0</td>
</tr>
<tr>
<td>Standard (Diazepam)</td>
<td>0.50</td>
<td>2.6 ± 0.24</td>
<td>198.2 ± 4.71</td>
<td>529.94</td>
</tr>
<tr>
<td>Seed extract 25</td>
<td>8.5 ± 0.44</td>
<td>39.8 ± 5.46</td>
<td>106.41</td>
<td></td>
</tr>
<tr>
<td>Seed extract 50</td>
<td>6.00 ± 0.44</td>
<td>74.8 ± 4.12</td>
<td>200.00</td>
<td></td>
</tr>
<tr>
<td>Seed extract 100</td>
<td>3.5 ± 0.22</td>
<td>140.8 ± 5.44</td>
<td>376.47</td>
<td></td>
</tr>
<tr>
<td>Seed extract 200</td>
<td>1.95 ± 0.16</td>
<td>263.6 ± 4.6</td>
<td>704.81</td>
<td></td>
</tr>
</tbody>
</table>

Values are represented as Mean ± SEM (*n* = 5), *p* < 0.05, that is the comparing the control group significantly (two-way ANOVA followed by Bonferroni’s test).

Table 4. Neuropharmacological potential test of *Citrus limon* (lemon) seed extracts by Hole cross method.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dose (mg/kg)</th>
<th>0 min</th>
<th>30 min</th>
<th>60 min</th>
<th>90 min</th>
<th>120 min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>10 mL/kg</td>
<td>5.0 ± 0.70</td>
<td>6.2 ± 0.37</td>
<td>6.2 ± 0.48</td>
<td>6.6 ± 0.40</td>
<td>8.6 ± 0.24</td>
</tr>
<tr>
<td>Standard (Diazepam)</td>
<td>1</td>
<td>3.4 ± 0.92</td>
<td>5.0 ± 1.48</td>
<td>4.0 ± 1.30</td>
<td>3.6 ± 0.60</td>
<td>4.0 ± 0.63</td>
</tr>
<tr>
<td>Seed extract 50</td>
<td>2.8 ± 1.15</td>
<td>4.2 ± 1.15</td>
<td>4.4 ± 0.67</td>
<td>3.4 ± 0.67</td>
<td>2.8 ± 0.58</td>
<td></td>
</tr>
<tr>
<td>Seed extract 100</td>
<td>1.4 ± 0.50</td>
<td>2.6 ± 0.50</td>
<td>2.2 ± 0.48</td>
<td>3.0 ± 0.94</td>
<td>2.6 ± 0.81</td>
<td></td>
</tr>
</tbody>
</table>

Values are represented as Mean ± SEM (*n* = 5), *p* < 0.05, that is the comparing the control group significantly (two-way ANOVA followed by Bonferroni’s test).

Table 5. Neuropharmacological potential test of *Citrus limon* (lemon) seed extracts by hole board method.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dose (mg/kg)</th>
<th>Quantity of head dips</th>
<th>% Inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>10 mL/kg</td>
<td>21.4 ± 2.97</td>
<td>0</td>
</tr>
<tr>
<td>Standard (Diazepam)</td>
<td>1</td>
<td>7 ± 0.70</td>
<td>67.28%</td>
</tr>
<tr>
<td>Seed extract 50</td>
<td>8.4 ± 0.81</td>
<td>60.74%</td>
<td></td>
</tr>
<tr>
<td>Seed extract 100</td>
<td>5.8 ± 0.37</td>
<td>72.89%</td>
<td></td>
</tr>
</tbody>
</table>

Values are represented as Mean ± SEM (*n* = 5), *p* < 0.05, that is the comparing the control group significantly (two-way ANOVA followed by Bonferroni’s test).

3.5 CNS depressant activity observed in animals treated with lemon seed extracts as indicated by the open-field method

The extracts substantially reduced the locomotor activity in tested mice at 50 and 100 mg/kg doses (*p* < 0.05). This result was noticeable from the underlying perception duration (0 min) and was consistent with the fifth perception duration (120 min) (Fig. 5). From the 2nd inspection to the 5th inspection, diazepam (1 mg/kg) demonstrated a substantial decline in locomotor activity in mice. This test showed a maximum of 36% suppression of locomotor function with seed extract at 100 mg/kg, while 37% was reported with the standard drug diazepam (Table 6).

4. Discussion

Traditional medicines have been considered an original form of therapeutic approach that has been inexpensive and effective since ancient times. Recent efforts have focused on including neurobehavioral function and efficiency to complement modern medication with different ethnomedical products. Sedative properties for such bioactive compounds and natural products have been explored with standard protocols like hole-cross and free field experiments that can evaluate and record the naturalistic locomotor behavior of mice or its alteration under induced conditions. Our findings indicate that the oral administration of the experimental plant seed extract at the doses (50 and 100 mg/kg) resulted in a significant decrease in the count of holes passed (Table 4). The repressive behavior was shown at 30 min when the extracts were given and persisted for 120 min. The extracts at the studied doses induced a substantial locomotion reduction that was observed in the open field test from 30 min to 120 min during the inspected duration (Table 6). It is known that by observing the total distance, the open field test can be used to evaluate the anxiety behavior. In the current study, the resting time’s range was between 6–12 seconds, the duration of time spent in center squares was 3 min, and the distance traveled into the center squares was between the ranges of 8.5 m²–19.3 m². The findings demonstrated that the extract reduced the locomotor activity, supporting the extract’s CNS...
and prolonged the sleep periods (Fig. 2) and prolonged the sleep periods (Fig. 1) depending on the doses and demonstrated substantial sedative effects. Both tests suggestively diminished locomotion in all Swiss albino mice treated with the standard and seed extracts. Animal head-dipping behavior remains closely correlated with their psychological response. Gamma amino-butyric acid (GABA) can modulate the central nervous system through the voltage-gated Ca\(^{2+}\) channel or chlorine conductance [40–50]. It was revealed that the animal’s head-dipping behavior is clearly relevant to their mental state, and this test is used to investigate anxiety-related activities. Based on the research findings, it was hypothesized that an increase in head-dipping behavior could indicate an anxiolytic state in animals. At the same time, a decrease in the number of head dips was related to a depressive impact [31]. Thus, the reduced head dipping behavior and improved sleeping time of mice in our study may relate to the inhibitory effect of the extract on neuronal activity, similar to the inhibition induced by GABA [51–56]. The therapeutic benefits of traditional medications might amalgamate a combination of constituents that could serve as an adjuvant to the conventional regimen. Multiple studies have reported that phytochemical compounds possess sedative and hypnotic effects, including tannins which can cause nonspecific CNS depression [56–58]. Several such action mechanisms have been proposed for alkaloids, flavonoids, steroids, as well as terpenoids, including activation of protein kinase C, neuroprotection against oxidative and metabolic insults; enhancing nicotinic receptors that also elevate the sensation and memory; energizing and improving nervous function; activating the transient receptor calcium channels in the membrane of the nerve cell which have neuropharmacological influence [59–67].

5. Conclusions

Recent studies in literature have reported the pharmacological benefits of the lemon constituents on the neurological well-being [68–70]. The current study provides evidence of the pharmacological properties of Citrus limon (lemon) seed extract in complementing the effects of thiopental sodium by restricting the locomotor activity in the mice model. Thiopental sodium is a standard drug for the induction of anaesthesia and hypnosis. The reduction in the sleep onset and an enhanced sleep duration induced in this drug model by the lemon seed extract provide a potential adjuvant for the qualitative augmentation of the existing protocols. The decrease in unconstrained locomotors and explorative properties further demonstrates the anxiolytic potential of the extract. However, further studies are warranted to investigate the plausible modes of action responsible for such neuropharmacological effects of Citrus limon (lemon) seed extract. The study broadens the scope of the lemon fruit as a nutraceutical or functional food with neuro-modulatory pharmacological properties.

**Table 6. Neuropharmacological potential test of Citrus limon (lemon) seed extracts by the open-field method.**

<table>
<thead>
<tr>
<th>Group</th>
<th>Dose (mg/kg)</th>
<th>0 min</th>
<th>30 min</th>
<th>60 min</th>
<th>90 min</th>
<th>120 min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>10 mL/kg</td>
<td>26.8 ± 1.56</td>
<td>28.0 ± 2.38</td>
<td>28.6 ± 2.15</td>
<td>31.8 ± 2.13</td>
<td>38.6 ± 2.50</td>
</tr>
<tr>
<td>Standard</td>
<td>1</td>
<td>16.2 ± 4.11</td>
<td>19.2 ± 5.95</td>
<td>21.8 ± 5.84</td>
<td>23.0 ± 5.50</td>
<td>24.2 ± 5.66</td>
</tr>
<tr>
<td>Seed extract</td>
<td>50</td>
<td>19.4 ± 3.18</td>
<td>25.2 ± 3.15</td>
<td>24.2 ± 3.30</td>
<td>23.0 ± 2.60</td>
<td>25.4 ± 2.63</td>
</tr>
<tr>
<td>Seed extract</td>
<td>100</td>
<td>16.6 ± 2.20</td>
<td>20.4 ± 3.23</td>
<td>19.2 ± 1.74</td>
<td>21.6 ± 2.13</td>
<td>24.4 ± 2.48</td>
</tr>
</tbody>
</table>

Values are represented as Mean ± SEM (n = 5), p < 0.05, that is the comparing the control group significantly (two-way ANOVA followed by Bonferroni’s test).

### Abbreviations

C. limon, Citrus limon; Ltd, limited; BDH, British Drug House; HCL, Hydrochloric acid, Normality; M, Molarity; b.w, body weight; CNS, Central nervous system; GABA, Gamma amino-butyric acid.

### Author contributions

MMR—Experimental work; FI—Experimental work; AP: Experimental work; MAK—Statistical analysis; GMA—Conceptualization and Analysis; MFU—Study Design, Analysis and manuscript writing; MA—Conceptualization, Study design, Analysis and Manuscript writing.

### Ethics approval and consent to participate

Animals were obtained from Jahangirnagar University, Dhaka, Bangladesh. All protocols considered in our study, including animals, were approved by the Faculty of Allied Health Sciences Research Ethics Committee, Daffodil International University, Dhaka-1207, Bangladesh. (Ref: FAH-SREC/DIU/2020/1006).

### Acknowledgment

The authors are grateful to the Department of Pharmacy, Jahangirnagar University, Dhaka, Bangladesh for providing experimental animals. The authors also condole the sudden demise of Dr. Sharif Mohammad Shaheen and gratefully acknowledge his contribution to this work.
Funding
This work was funded by the Deanship of Scientific Research (DSR), King Abdulaziz University, Jeddah, Saudi Arabia under grant no. (KEP-1-141-41). The authors, therefore, acknowledge with thanks DSR technical and financial support.

Conflict of interest
The authors declare no conflict of interest.

References
[29] Olas B. A review of in vitro studies of the anti-platelet potential of citrus fruit flavonoids. Food and Chemical Toxicology. 2021; 150: 112090.
the determination of tannin and flavonoid levels and some applications in ethnomedicine and ethnopharmacology. Functional Ecosystems and Communities. 2008; 2: 88–94.


