STUDIES ON TOXICOLOGICAL AND NEUROBEHAVIORAL PROFILE OF METHANOL EXTRACT OF MUSSAENDA ROXBURGHII HOOK. F. LEAVES IN MICE

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ABSTRACT

Objective: The aim of this study was to evaluate the toxicity of Mussaenda roxburghii with special reference to the nervous system.

Methods: For the study, 15 mice were obtained from Pasteur Institute, Shillong. The mice were then divided into three groups containing five mice in each group. The groups were, namely Group I, Group II, and Group III; Group I mice received distilled water and acted as a control group, Group II mice received plant extract at a dose of 600 mg/kg body weight (BWT), and Group III mice received plant extract at a dose of 800 mg/kg BWT. The doses were administered orally through oral gavage tube for 28 days and the BWT of the mice was measured at 7th, 14th, 21st, and 28th days. The behavior of mice was evaluated for anxiety, locomotion, immobility, learning, and memory with the elevated plus maze test (EPMT), open field test (OFT), forced swim test (FST), and Morris water maze test (MWMT), respectively.

Results: The result of the experiment showed a decrease in the BWT of mice exposed to plant extracts significantly as compared to the control. In the OFF, there is a significant decrease in total distance travel in OFT and also total distance travel in the central zone in mice treated with the plant extracts as compared to the control mice. In the EPMT, the plant extract treated mice showed a decreased in the time spent in open arms. The FST results in increased immobility in mice exposed to plant extracts as compared to control. In the present study, MWMT results in an increased escape latency and path length and in decreased annulus crossovers in plant extract treated group as compared to control.

Conclusion: The results of the present study suggest that the plant extract alters the behavior of the treated mice and possesses neurotoxic activity.

Keywords: Northeast, Toxicity, Neurobehavioral, Mice.

INTRODUCTION

Mussaenda species member of the Rubiaceae family is used in West Africa, Indian subcontinent, Southeast Asia, and Southern China. This plant is used in Chinese, Fijian, and Indian folkloric preparations to treat various diseases and possesses medicinal activity such as the diuretic, antiphlogistic, and antipyretic [1]. Mussaenda roxburghii (MR) Hook. is one of the species found in Bhutan, Bangladesh, Myanmar and in the Northeast part of India. This plant has been reported to use to treat boils in tongue, bacterial and fungal infection as traditional folk medicine and also used among tribal people of Arunachal Pradesh, India, as food [2]. Studies reported that these plant ethyl acetate and dichloromethane root extracts possess good thrombolytic, anticancer, and anti-inflammatory activity. Methanol leaves extract at a dose of 20, 40, and 60 mg/kg showed anti-neoplastic properties in cancer cell lines, highest anticancer activity showed at 60 mg/kg [3]. Leaves of this plant have anti-inflammatory, thrombolytic, antitumor, and antiarthritic activity effect. Methanolic extract showed 53.79% and 86.93% of membrane stabilizing activity at 31.25 μg/ml and 1000 μg/ml concentration, respectively [4,5]. Another author demonstrated that methanolic extract has strong cytotoxic activity with the LC50 value of 0.52 and 0.62 μg/ml and antimicrobial activity on Staphylococcus aureus MR [6]. Hence, the present study was carried out to evaluate the toxicity, neurobehavioral alterations induced by this plant.

MATERIALS AND METHODS

Plant materials
The plant material was collected in August 2016 from Dhalai District, Tripura, India. The plant was identified and authenticated by the Botanical Survey of India, Shillong, with No. BSI/ERC/Tech/Identification/2016/315.

Animals
Adult male Swiss albino mice weighing between 20 and 30 g were used for the study which was obtained from the Pasteur Institute, Shillong. The animals were housed in cages under standard environmental conditions and had free access to food and water ad libitum. All the procedures were performed after the approval institutional ethical committee (Ref. No. AUS/IAEC/2017/PC/18).

Phytochemical analysis
The phytochemical analysis carried out using standard methods [10].

Quantitative estimation of chemical constituency

Total alkaloid estimation
A total of 200 ml of 20% acetic acid was added to 5 g of leaf powders taken in a separate 250 ml beaker and covered to stand for 4 h. This mixture containing solution was filtered and the volume was reduced to 1 quarter using a water bath. To this sample, concentrated ammonium solution was added dropwise until the precipitate was complete. The whole solution was allowed to settle and the precipitate was collected after filtration and weighed [11].
Total terpenoids estimation
About 2 g of the dried leaf powder was weighed and soaked in 50 ml of 95% ethanol for 24 h. The extracts were filtered and the filtrate extracted with petroleum ether (60–80°C) and concentrated to dryness. The dry ether extracts were treated as total terpenoids [11].

Total flavonoids
The total flavonoids content was estimated using the procedure described by Thakur and Sahani [12].

Total phenol estimation
The total phenol content of MR was estimated using Folin–Ciocalteu reagent [13]. About 20 µg leaf extract was taken separately and it was made up to 1 ml with DDW. Then, 500 µl of diluted Folin–Ciocalteu reagent and 2.5 ml of sodium carbonate added. The mixture was shaken well and then incubated in dark condition for 40 min. After incubation, the absorbance was read at 725 nm. A calibration curve of gallic acid was prepared and used for determining the total phenol content and expressed as gallic acid equivalents.

Preparation of methanolic extract
The plant leaf was shade dried, powdered, and extracted at 60°C with 75% methanol using Soxhlet apparatus. The solvent was evaporated under reduced pressure to obtain crude extracts. The extracts were suspended in dissolved in normal saline and were used for the following toxicological studies [14].

Toxicity studies
Determination of LD₅₀
Swiss albino male mice of approximately the same weight were used. About 20 mice divided into five groups each group containing four mice. Acute toxicity was tested in response to 400, 800, 1200, 1600, or 2000 mg/kg of methanolic leaf extracts. Animals were observed for 72 h for behavioral changes or mortality. The number of mice died within 72 h was recorded for each group, and subsequently, the median lethal dose (LD₅₀) was determined by probit analysis [14,15].

Experimental design
For the study, 15 mice were obtained from Pasteur Institute, Shillong. The mice were then divided into three groups containing five mice in each group. The groups were namely, Group I, Group II, and Group III; Group I mice received distilled water and acted as a control group, Group II mice received plant extract at a dose of 600 mg/kg BWT, and Group III mice received plant extract at a dose of 800 mg/kg BWT. The doses were administered orally through oral gavage tube for 72 h for behavioral changes or mortality. The number of mice died for 72 h was recorded for each group, and subsequently, the median lethal dose (LD₅₀) was determined by probit analysis [14,15].

Neurobehavioral evaluation tests
Open field test (OFT)
Open field activity was measured in a square area (25 cm × 25 cm). At the beginning of a test, mice were placed in the central part of the area. We recorded the total distance traveled in the central zone and the total distance traveled in the whole area. The mice were tracked during the test by video-tracking software animal tracker [29].

Forced swim test (FST)
FST was conducted by Porsolt et al. [16]. The animal was individually forced to swim in a transparent vessel (25 cm high, 15 cm in diameter) filled with (12.5 cm high) water and a temperature of 21–24°C was maintained. The duration of immobility (in second) was measured for 5 min. “Immobility” was defined as floating and treading water just enough to keep the nose above water. The water was changed after every other trial. The total immobility period during a 5 min test was recorded on day 28 [34].

Elevated plus maze test (EPMT)
The EPM apparatus consists of two open arms (16 cm × 5 cm) and two closed arms (16 cm × 5 cm × 12 cm) having an open roof. The whole apparatus was elevated 25 cm from the floor in a dimly lit room. The mice were placed individually at the center of the maze with their head facing a closed arm. The mice were allowed to explore the maze for 5 min. The number of entries in open arms and time spent in open arms was recorded [18].

Spatial learning and memory examinations: Morris water maze test (MWM)
After the PH administered for 28 days, the Morris water maze was made up of a circular pool of diameter 110 cm and height of 30 cm, the circular pool was filled with water and maintained a temperature at 23±10°C. The circular pool was divided into four quadrants of equal area. A hidden transparent escape platform of 9 cm in diameter was placed in the center of one of the four quadrants of the pool. Mice have administrated trials for 4 days repeatedly with an interval of 24 h. For each daily trial, the mouse was placed into the water maze and allowed to find the hidden platform. Once the mouse located the platform, it was allowed to remain on it for 10 s, if the mouse did not locate the platform, it was placed in a platform for 15 s. After finding the hidden platform, the trial was stopped and escape latency time was recorded [23,32]. The animal movement path was recorded using a video camera and AnimalTracker software [33].

BWT
The BWT of mice was recorded once a week and the food and water consumption by each mouse was measured.

Statistical analysis
The data were analyzed using SPSS. The data are expressed as mean±SEM. The statistical significance of difference was evaluated using one-way analysis of variance followed by Tukey’s procedure for multiple comparisons. p<0.05 was considered statistically significant.

RESULTS
Preliminary phytochemical screening
Results obtained from the experiments showed the presence of alkaloids, flavonoids, saponin, carbohydrates, tannin, terpenoids and fat, oil, etc. (Table 1).

Quantitative phytochemical analysis
Results of the quantitative phytochemical analysis of the plant extract are represented in Table 2.

Acute toxicity (LD₅₀)
Most of the animals showed a decrease in the water consumption, food consumption, increased in respiratory rate, and dyspnea before death. The first death was recorded 4 h after oral administration of the highest

Table 1: Phytochemical screening of methanol extract of MR

| Plant constituents | Methanol extracts |
|--------------------|------------------|
| Alkaloid           | +                |
| Saponin            | –                |
| Terpenoid          | +                |
| Phenol             | +                |
| Flavonoids         | +                |
| Steroid            | +                |
| Protein            | +                |
| Glycoside          | +                |
| Carbohydrates      | +                |
| Sterol             | +                |
| Tannin             | +                |

MR: Mussaenda roxburghii, +: Present, –: Absent
dose (2400 mg/kg), which caused 100% mortality. The LD$_{50}$ of MR was calculated to be 1778.27 mg/kg BWT (Fig. 1 and Table 3).

The result of the BWT after subchronic administration of mice is reported in Fig. 2. A significant decrease in BWT was observed in the 2nd week after the treatment of the plant extract in Group III (p<0.05) and in the 3rd and 4th weeks in both Group II and Group III as compared to the control group.

**OFT**

The effect of administration of crude extracts of MR (600 or 800 mg/kg) in OFT on mice is reported in Fig. 3, Fig. 4 and Fig. 5 for the total distance traveled and total distance traveled in the central zone, respectively. The total distance traveled was significantly lower in both plant extract treated groups, i.e., Group II (p<0.05) and Group III (p<0.001) as compared to the control group. The total distance traveled in the central zone was also significantly lower in both plant extract treated groups, i.e., Group II (p<0.01) and Group III (p<0.001) as compared to the control group.

**EPM**

In the EPM, the time spent in open arm and the percentage of time spent open arm were significantly decreased in Group II (p<0.05) and Group III (p<0.01) and compared to that of the control group (Fig. 6 and Fig. 7).

**FST**

In the FST, the immobility was recorded in second for 5 min. The immobility was increased in the Group III significantly (p<0.01) as compared to the control group (Fig. 8).

**MWMT**

The result of the MWMT for escape latency is reported in Fig. 9 and Fig. 10. On the 3rd, 4th, and 5th days, there is a significant increase in escape latency in Group III, and on the 5th day, there is a significant increase in Group II. The result of path length is reported in Fig. 11. On the 2nd, 3rd, 4th, and 5th days, there is a significant increase in path length in Group III, and on the 4th and 5th days, there is a significant increase in Group II in path length as compared to the control group. The target annulus crossover is reported in Fig. 12. A significant decrease in Group II was observed as compared to that of the control group.

**DISCUSSION**

*M. roxburghii* is an ethnomedicinal plant used in traditional medicine of Tripura and also other parts of Northeast India. However, scientific data on the toxicity of this plant are not available yet. Till now, no data have been published concerning its safety on the central nervous system. The preliminary phytochemical analysis showed that the methanolic extract of MR presents alkaloid, terpenoid, glycoside, phenol, flavonoids, steroids, and protein. The quantitative phytochemical analysis of the plant extract showed the presence of total alkaloid of 3 g/100 g and total plant terpenoids (mg/10 g) of 14.5621 mg/10 g.

### Table 2: Quantitative analysis of phytochemical of the leaves extracts of MR

| Total plant terpenoids (mg/10 g) | Total plant alkaloids (mg/100 g) | Methanol extracts total flavonoids content (g/100 g) | Methanol extracts total phenol content (g/100 g) |
|----------------------------------|---------------------------------|-----------------------------------------------|-----------------------------------------------|
| 14.5621 mg/10 g                  | 3 g/100 g                       | 13.9736842 g/100 g                             | 9.74137931 g/100 g                          |

MR: *Mussaenda roxburghii*

### Table 3: Determination of acute oral toxicity of MR

| Dose mg/kg (body weight) | Log dose | Total number of animal died (72 h) | Mortality (%) | Corrected mortality (%) | p value |
|-------------------------|----------|-----------------------------------|---------------|-------------------------|---------|
| 400                     | 2.6021   | 0                                 | 0             | 2.50                    | 3.04    |
| 800                     | 2.9031   | 0                                 | 0             | 2.50                    | 3.04    |
| 1200                    | 3.0792   | 0                                 | 0             | 2.50                    | 3.04    |
| 1600                    | 3.2041   | 1/4                               | 25            | 25.00                   | 4.33    |
| 2000                    | 3.3011   | 2/4                               | 50            | 40.00                   | 5       |
| 2400                    | 3.3802   | 4/4                               | 100           | 97.50                   | 6.96    |

Corrected formula: For 0% dead=100 (0.25)/n; for 100% dead=100 (n−0.25)/n, where n=Number of animals in each group. MR: *Mussaenda roxburghii*
total terpenoids of 14.56 mg/10 g, total phenol of 9.74 g/100g, and flavonoids of 13.9736842 g/100 g. Studies reported that this plant contains sterol glucosides which are toxic to the motor neurons [9].

In the present study, the effects of methanolic extract MR leaf on certain behavioral paradigms have been carried out on male Swiss albino mice. An important step in evaluating the action of a substance in the central nervous system (CNS) is to observe its effect on the behavior of the animal.

The LD<sub>50</sub> study of the methanolic extract of MR leaf showed no mortality at dose from 400 to 1200, whereas at doses from 2000 to 2400 mg/kg BWT, all the animals showed abnormal behavior and diarrhea developed within 4 h of administration of the extract. In the present study, the LD<sub>50</sub> of the extract by the oral route was 1778.27 mg/kg BWT. A recent study on acute toxicity by Islam et al. reported the LD<sub>50</sub> of methanolic extract of MR as 600 mg/kg BWT administered through i.p in mice [6].

Analysis of organ weight in toxicology study is important for identification of the harmful effect of chemicals [19]. In this study, changes in the BWT were observed in the mice groups fed with the plant extracts. Relative organ weights have been observed in toxicity study to be a relatively sensitive indicator for particular organs and define toxicity. Changes in mice brain weight or BWT, decrease or increase in food consumption and organ weight act as hallmark of stress [20].
It is an important step to evaluate the action of a substance on the behavior of the animal to observe its effects on CNS of the animal. The results of the present study showed a decrease in the locomotor activity in the mice exposed to MR which gives an indication of toxicity in the CNS and the decrease in locomotor activity may be closely related to sedation resulting from the depression of the CNS. This sedative effect may be due to the presence of toxic compounds of alkaloids, terpenoids, etc. The similar effect has been reported by many other plants extract [17,21,22,30].

The EPMT is used to investigate the anxiolytic and anxiogenic activity drugs. Anxiolytic drugs increase the amount of time spent in open arm, and anxiogenic drugs decrease the time spent in open arm. The animals which spent less time spent in open arms are associated with significant anxiety-related behaviors [18,26]. In this study, plant extract showed anxiogenic activity by decreasing the time spent in open arms in the plant extract treated mice as compared to that of the control mice. Similar results were found in studies of other plants where higher doses showed anxiogenic activity in EPMT [28].

The FST is used to monitor depression as behavior by estimating immobility [24]. When mice are forced to swim in an inescapable situation, they tend to become immobile after an initial vigorous activity. The immobility reflects a state of lowered mood in which the animals have given up hope of finding an exit and have resigned themselves to the experimental situation. In the present study, the significant increase in immobility in the mice treated with plant extract indicates depressant activity of the extract on the treated mice at the given doses [31].
In the present study, it was observed that in the MR treated groups, the escape latency and the path length (Fig. 9, Fig. 10, Fig. 11) were increased as compared to that of the control group, whereas the number of annulus crossovers (Fig. 12) in the MR treated groups was increased significantly which indicates that MR extract could cause learning and memory impairment in mice [27].

CONCLUSION

From the experiments of this study, it could be concluded that the methanolic extract of the plant possesses neurotoxic properties. The acute toxicity study showed that the extract was moderately safe at lower doses. Chronic exposure of mice of plant extract for 28 days changes the BWT. The neurobehavioral performance in the test mice was altered, which resulted in anxiety, depression, loss of locomotor activity, and learning and memory in mice. In conclusion, care should be taken while using this plant for medicinal purposes.

AUTHORS’ CONTRIBUTIONS

All the authors contributed equally.

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