BIOACTIVITY-GUIDED ISOLATION OF MEMORY-ENHANCING COMPOUND FROM CHLOROFORM EXTRACT OF ROOTS OF PLUMBAGO ZEYLANICA LINN.

VAIBHAV UPLANCHIWAR1, GUPTA MK1, RUPESH K GAUTAM2*

1Department of Pharmacy, Oriental University, Indore, Madhya Pradesh – 453 555, India. 2Department of Pharmacology, MM School of Pharmacy, Maharishi Markandeshwar University, Sadopur, Ambala – 134 007, Haryana, India. Email: drrupeshgautam@gmail.com

Received: 30 April 2018, Revised and Accepted: 25 May 2018

ABSTRACT

Objective: The main aim of our study is to isolate the active compound from roots of Plumbago zeylanica Linn. by bioactivity-guided isolation and evaluate its memory-enhancing effect by Morris water maze.

Methods: Roots were extracted by successive solvent methods by petroleum ether, chloroform, methanol, butanol, and finally, water. Chloroform extract was selected for isolation, and plumbagin was isolated by hexane and ethyl acetate as solvent system. Plumbagin was evaluated by Morris water test, and brain acetylcholine esterase level was measured.

Result: Plumbagin showed a significant decrease of escape latency and increase of time spent in target quadrant by mice in Morris water maze indicating improvement of learning and memory. It also significantly decreases the cholinesterase level in the brain.

Conclusion: Learning and memory of mice doubtless may be through embarrassment of brain acetyl cholinesterase activity and through involvement of GABA-benzodiazepine pathway. Further detailed study is required to explore the other possible mechanisms for the management of cognitive disorders.

Keywords: Bioactivity-guided isolation, Plumbagin, Cholinesterase level, Morris water maze, Locomotor activity.

INTRODUCTION

Alzheimer's disease is a progressive neurodegenerative disorder characterized by the gradual commencement of dementia. It is characterized by gradually progressive refuse in cognitive function, with deficits, especially in memory recovery [1]. A number of cholinesterase inhibitors are in clinical practice for the management of various cognitive disorders [2]. However, they are associated with a number of adverse effects, i.e., anorexia, nausea, vomiting, diarrhea, and insomnia [3]. Hence, there has been intense attention focused on the part of potential phytochemical to modulate neuronal function and protective mechanism against degeneration. As complementary and alternative therapy, herbal medicine or phytotherapy refers to the medical exploitation of plant components for their curative properties. A number of medicinal plants are reported and widely used in Ayurveda and other system of medicine for neurodegenerative disorders. Among all reported medicinal plants, Plumbago zeylanica (chitraka) is well-known medicinal plant for neurodegenerative disorders.

Plumbago zeylanica is an old-age Rasayana herb in traditional Ayurveda. P. zeylanica Linn. is distributed as a weed throughout the tropical and subtropical countries of the world [4] native to South Asia and cultivated throughout India and Sri Lanka. It belongs to family Plumbaginaceae [5-7]. The roots of P. zeylanica are also used as expectorant, antiarrhythmic, anti-sabies, appetite stimulant, anti-diarrheal, antiperiodic, and antimalarial and for the treatment of leprosy [4]. The second hand of whole plant, roots, and powder of the roots [8,9] is in fever and malaria, in opposition to diarrhea, dyspepsia, piles, and skin diseases including leprotic lesions, learning, and memory [10-12].

In our previous study, various extracts of roots of P. zeylanica were evaluated by Morris water test and their brain acetylcholine esterase level was measured. Among all the extracts, chloroform extract showed potent activity for learning and memory [13]. Phytochemical study of chloroform extract showed the presence of active compound, i.e., plumbagin. Hence, an attempt was made for the bioactivity-guided isolation of active compound from chloroform extract and its evaluation in various models of learning and memory.

METHODS

Procurement of plant material
The roots of P. zeylanica were purchased from the market. The roots were taxonomically identified and authenticated by senior botanist. A voucher specimen is preserved in the department for the further reference.

Successive solvent extraction method
The roots (250 g) were dried in shade and sliced into small pieces and pulverized using a mechanical grinder for the coarse powder. The coarse powder of root was subjected to Soxhlet extraction using petroleum ether to remove all fats. The marc was dried and then extracted by using chloroform, methanol, butanol, and finally with water for 72 h. After exhaustive extraction, the extracts were filtered, concentrated, and dried. All the extracts were screened for the presence of alkaloids, fatty acids, terpenoids, steroids, flavonoids, and glycosides. Since chloroform extract showed potent activity in Morris water test, it is selected for further isolation and characterization of active compounds.

Chromatographic separation and isolation of active compound
Completely dried chloroform extract (25 g) was subjected to chromatographic separation by loading of extract on glass column (90×3c). For the stationary phase, silica gel was used. Isolation of compound was followed by gradient elution method. In this method, n-hexane containing increasing amounts of ethyl acetate (90:10, 80:20, 70:30, 60:40, 50:50, 40:60, and 30:70) was used as the solvent system. The column was eluted by n-hexane containing increasing amounts of ethyl acetate till 90% of the process completed. One thirty-five fractions were together and fractions showing similar chromatogram on thin-layer chromatography plate were pooled together and total four (F1-F4) fractions were prepared. The percentage yields of fractions were...
determined. Fractions (F2) eluted showing only one spot on TLC (n-hexane:ethyl acetate, 9:1; Rf-0.61) were pooled and evaporated to dryness, yielding plumbagin (F2). After isolation of active compound, melting point of plumbagin was recorded by capillary tube method. For the spectral determination of compound, various spectral methods, i.e., ultraviolet and infrared spectra were followed [14].

Experimental animals
Disease-free Swiss male albino mice (25-35 g) were used for this experiment, and they were purchased from the Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) approved vendor. Animals were housed separately in polycarbonate cage in groups of 6–8 per cage under proper laboratory conditions with alternating light and dark cycle of 12 h each. Before the experiment, the protocol was approved by the Institutional Animal Ethics Committee and animal care was in use as per the strategy of CPCSEA, Ministry of Environment and Forests, Government of India.

Morris water maze
The method and parameters for learning and memory of mice using Morris water maze were followed as reported earlier [15-17].

Animals were divided in eight groups and six animals were placed in each group. Group 1 served as control and Group 2 as standard drug (physostigmine, 0.1 mg/kg, i.p.) treated. Animals of Groups 3 and 4 were treated by plumbagin in a dose of 1.5 and 3 mg/kg, rats of Groups 5 and 6 were treated by scopolamine and diazepam in a dose of 0.4 and 1 mg/kg, and rats of Groups 7 and 8 were treated by plumbagin and combination of scopolamine and diazepam, respectively, were administered for 15 consecutive days. Escape latency (EL) was recorded 120 min after drug administration from 11th day to 14th day. On the 15th day, time spent in target quadrant (TSTQ) was noted 120 min following the drug administration. In case of animals administered with physostigmine, EL and TSTQ were noted after 30 min of drug administration.

Measurement of locomotor activity
All the eight groups were treated by drugs, respectively, which were administered for 15 successive days. Locomotor activity was measured on the 15th day using actophotometer ([NICO, Ambala]).

Biochemical estimation
Collection of brain sample
After the 15th day using Morris water maze, the animals were sacrificed on the 16th day by cervical dislocation. Whole brain was carefully removed from the animals. The fresh whole brain was weighed first and then homogenized in 10 volumes of 0.1 M phosphate buffer (pH 8) using a glass homogenizer. The homogenate was centrifuged at 3000 rpm for 10 min at 4°C using refrigerated centrifuge (Remi, Mumbai). The resultant cloudy supernatant liquid was used for the estimation of brain acetylcholinesterase activity [18].

Estimation of acetylcholinesterase activity
0.4 ml of brain homogenate was placed into a test tube containing 2.6 ml of phosphate buffer, 5, 5-dithiobis-2-nitrobenzoic acid reagent (0.1 ml) was added to the above mixture, and absorbance was noted at 412 nm. Then, 0.02 ml of acetylcholine iodide solution was added and again absorbance was noted 15 min thereafter. Change in absorbance per minute was calculated [18].

Statistical analysis
Data were analyzed by analysis of variance followed by Tukey’s post hoc test in GraphPad prism, p<0.05 was considered as statistically significant. All the results are expressed as mean±standard error of the mean.

RESULTS
Effect of plumbagin on EL and time spent in target quadrant TSTQ
EL and time spent in target quadrant are associated with learning and memory, and decrease of EL and increase of TSTQ by mice in Morris water maze indicate an improvement of learning and memory. In our study, plumbagin (1.5 and 3 mg/kg) and physostigmine (0.1 mg/kg, i. p.) administered for 15 successive days. After administration, they significantly decreased EL of mice from 11th to 14th day and augmented TSTQ by mice on the 15th day as compared to the control, thus showing learning and memory enhancement effect. Rats treated by plumbagin in a dose of 3 mg/kg significantly upturned scopolamine- and diazepam-induced amnesia as compared to scopolamine- and diazepam-treated groups (Figs. 1 and 2).

Effect of plumbagin on locomotor activity of mice
There was a significant change in locomotor activity in mice treated by plumbagin (1.5 and 3 mg/kg) and physostigmine as compared to vehicle-treated control (Fig. 3).

Effect of plumbagin on brain acetylcholine esterase activity of mice
Animals treated by plumbagin and physostigmine for 15 successive days produced a significant reduction in brain acetylcholinesterase activity as compared to control group. Mice treated by plumbagin in a dose of 3 mg/kg showed a highly significant decreasing effect on brain acetylcholinesterase activity compared to rats of control group. Results were expressed in Fig. 4.

DISCUSSION
Herbs have been employed from 1000 of years, in several forms, under the indigenous system of medicine such as Ayurveda, Unani, and Siddha. The unbelievable development in the ground of synthetic drugs during present era is accompanied by numerous unwanted side effects, whereas plants still grasp their own unique position, with lesser
Acetylcholinesterase is decreased by plumbagin, it might be suggested that the memory-enhancing consequence of plumbagin due to inhibition of acetylcholinesterase leads to increase in brain acetycholine levels. Thus, the drugs which improve cholinergic function can be used for the treatment of dementia closely connected to Alzheimer’s disease.

CONCLUSION

By this study, it can be concluded that roots of *P. zeylanica* proves its claim for memory-enhancing effect according to Ayurveda and other system of medicine. Plumbagin was isolated as chief active phytoconstituents by bioactivity-guided isolation and it significantly improved learning and memory of mice doubtless through embellishment of brain acetylcholinesterase activity and through involvement of GABA-benzodiazepine pathway. Further detailed study is required to explore the other possible mechanisms for the management of cognitive disorders.

CONFLICTS OF INTEREST

The authors declared that they have no conflicts of interest.

AUTHORS’ CONTRIBUTION

All authors contributed uniformly.

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