Assessment of Acetylcholinesterase Inhibition and Anti-inflammatory Activity of Ethanolic Extract of Evolvulus alsinoides Linn.

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INTRODUCTION

Acetylcholine is a fast-acting neurotransmitter, alters neuronal excitability and coordinates firing of neurons in central area. ACh work through pre- and postsynaptic receptors in the brain and contribute to cortical and hippocampal function across phylogeny. ACh is released from nerve terminals in small quantum and extruded by exocytosis. ACh has a very short half-life because acetylcholinesterase (AChE) rapidly hydrolyses ACh in the terminal of nerve impulse transmission cholinergic synapses. AChE is the most essential enzymes in the family of serine hydrolases. Functions of acetylcholine in learning and memory are already known but now a day it is used in the treatment of CNS disorders like Alzheimer's and Parkinson's disease. Alzheimer's disease (AD) is well-known neurodegenerative diseases, firstly described and later named by German psychiatrist and pathologist Alois Alzheimer in 1906. In initial phase of AD, patients decline in cognitive functions and find difficulties even to remember recent events called short-term memory loss while on progression of the disease, patients experience long-term memory loss which includes difficulties in speech, speaking and cognitive thinking etc. and during the late phase language deficits, psychosis, aggressive behavior and depression etc are happened.

In 1961 Ellman, Courtney, Andres and Featherston developed a method known as Ellman's method to determination of activity of AChE enzyme. It is quantitative and photometric determination method using extract, Thiocholine + Dithiobisnitrobenzoate ion using spectrophotometer at 412 nm.

Objectives:

The objective of present study is to determine the efficacy of ethanolic extract of Evolvulus alsinoides Linn. for therapeutics at various doses. One more purpose is also to determine anti-inflammatory effects of E. alsinoides which are highly associated with CNS disease.

Methods:

Ethanolic extract of E. alsinoides has been used for estimation of acetylcholinesterase inhibitory activity at dose of 100, 200, 300 mg/kg on rat. For estimation of acetylcholinesterase, Ellman's method is used which is based on absorbance. Anti-inflammatory activity was evaluated by two in vitro experimental models i.e. heat induced hemolysis and hypotonic solution-induced hemolysis.

Results:

Ethanolic extract of E. alsinoides at the dose of 200mg/kg is more effective than 100 and 300. In heat hemolysis, inflammation was significantly reduced at 50 and 100 μg/ml in EA while hypotonic solution induced inhibition was shown significant at 50, 100 and 200 μg/ml in EA. Extract was effective in inhibition of hemolysis in a dose-dependent manner due to its membrane stabilizing action. Thus, E. alsinoides may be employed as alternative and safe drug therapy against inflammation. Conclusion: Several natural products have been used for medicinal purposes in which E. alsinoides is also one of them. It is traditionally claimed for memory enhancement and will help as therapeutics for brain related problems.

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performed using ethanolic extract and its fraction on learning behavior and memory enhancing activity.16,17

MATERIALS AND METHODS

Plants material

_E. alsinoides_ was collected from area adjoining to University campus and authenticated [Bot/H/05/111/01] in the Department of Botany, Dr. Harisingh Gour Vishwavidyalaya, Sagar, (M.P.).

Extraction of _E. alsinoides_ using soxhlet assembly

The dry herb of _E. alsinoides_ was inserted in a soxhlet apparatus with petroleum ether to defatting purpose then subjected with ethanol (90%) for 72 hr at 55°C. The collected extract was concentrated and subjected for study.

Estimation of Total phenolic contents

Preparation of drug sample

1ml of extract, 10 ml of water and 1.5ml of Folin and ciocalteu’s reagent (Phenol reagent) were taken in to 25ml volumetric flask and kept for 5 min. 4 ml of 20% w/v sodium carbonate solution was added and the volume was made up to 25 ml with double distilled water. The mixture was kept for 30 minutes and then measured the absorbance at 765nm using UV-visible spectrometer (Shimadzu, Japan). The total phenolic content was calculated by using gallic acid as standard.18

Preparation of Standard solution and standard curve

100 mg pure gallic acid was dissolved in 100 ml distilled water. This solution was again diluted with distilled water and prepared aliquots (100, 200, 300, 400, 500, 600 µg/ml). 1 ml of aliquots, 10 ml of water and 1.5ml of Folin and ciocalteu’s reagent (Phenol reagent) were transferred to the 25 ml volumetric flask and kept for 5 min. 4 ml of 20% w/v sodium carbonate solution was added and the volume was made up to 25 ml with double distilled water. The mixture was kept for 30 minutes and then measured the absorbance at 765 nm using UV-visible spectrometer (Shimadzu, Japan).

Determination of AChE inhibitory activity

Acute toxicity study

The acute toxicity study was performed as per OECD guidelines 423. Three animals were selected for each dose group i.e. 5, 50, 300 and 2000 mg/kg body weight, given p.o. Different activities of animals (change in fur color, behavior, any lethargic sign, etc.) were observed in the first 4 hours after 10 h and once a day daily for 14 days.

Experimental animals

Wistar rats of either sex (150-200g) were selected. They were procured from the animal house of the institute. The experimental protocols were approved by the Institutional animal ethics committee (IAEC No. 379/GO/ReBi/S/01/CPCSEA, Reference no. 379/CPCSEA/IAEC-2018/035) after scrutinization. The animals were fed with standard pelleted diet and water ad libitum. The animals were acclimatized to the laboratory condition before experiment. The animals were fasted for at least 24 h before treatment. The animals were divided into 6 group’s i.e.

- Group 1: Negative control group (0.2% v/v Tween-80)
- Group 2: Positive control group (scopolamine, 0.3 mg/kg; i.p)
- Group 3: Standard group (piracetam, 100 mg/kg; oral)
- Group 4: Ethanolic extract of EA (100mg/kg)
- Group 5: Ethanolic extract of EA (200mg/kg)
- Group 6: Ethanolic extract of EA (300mg/kg)

The animals were treated for 15 days, however, scopolamine was administered to the animals on the 9th day, intraperitoneally.17

Estimation of AChE inhibitory activity

The activity was determined through 96-well microplate assay according to the method of Ellman et al. 1961. In this method, thiocicholine produced by AChE reacts with 5,5′-dithiobis(2-nitrobenzoic acid) (DTNB) to form a colorimetric (412 nm) product. Acetylchicholine iodide (15 µM) in 25 µl was added into a microplate well along with the addition of DTNB (125 µl) in 50 mM of Tris/HCl (pH 8) with NaCl (0.1 M) and MgCl₂·6H₂O (0.02 M). Further, 50 µl of bovine albumin fraction V (0.1%), 25 µl of test drugs dissolved in methanol and diluted in Tris/HCl (50 mM, pH 8) at concentrations of 1.25, 12, 20, 40, 80, 160, 320 µg/ml were added into well. The absorbance was measured after 2 min of incubation at room temperature; the initial absorbance was taken at 412 nm and after 10 min, the final measurement was taken. All the analysis was performed in triplicate.19 The AChE activity was calculated using the following formula (Table 1):

\[
\text{AChE Activity (Units/L) = } \frac{(A_{412})\text{Final} - (A_{412})\text{Initial}}{(A_{412})\text{Calibrator} - (A_{412})\text{Blank}} \times n \times 200
\]

200 = Equivalent activity (Units/L) of the calibrator when assayed is read at 2 minutes and 10 minutes

n = dilution factor

\[
(A_{412})\text{Calibrator} = \text{Absorbance of the calibrator at 10 min}
\]

\[
(A_{412})\text{Blank} = \text{Absorbance of the blank at 10 min}
\]

In vitro anti-inflammatory activity of ethanolic extract of _E. alsinoides_

The anti-inflammatory activity of extract was performed using two models i.e Heat-induced haemolysis and hypotonic solution-induced haemolysis.20

Preparation of Erythrocyte suspension

It was prepared by collected whole blood followed by washing with normal saline (0.9%, NaCl). This volume was adjusted and reconstituted as a 40%, v/v suspension using isotonic buffer solution (pH 7.4). Hemolysis was induced by heat and hypotonic solution to measure the anti-inflammatory activity of test compounds by preventing hemolysis.

Heat-induced hemolysis

Test compounds were prepared by taking isotonic buffer solution (5 ml) of plants extracts in the concentrations of 50, 100, 200, 300 and 500 µg/ml into two duplicate sets. The vehicle was added to another tube separately for study. The animals were treated to the animals on the 9th day, intraperitoneally.

Table 1: AChE activity of extracts at different dose.

<table>
<thead>
<tr>
<th>Sn</th>
<th>Groups</th>
<th>AChE Activity (Units/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>01</td>
<td>Group-1</td>
<td>8.550 ± 0.3041</td>
</tr>
<tr>
<td>02</td>
<td>Group-2</td>
<td>22.91 ± 0.07339 ***</td>
</tr>
<tr>
<td>03</td>
<td>Group-3</td>
<td>15.17 ± 0.2962 ***</td>
</tr>
<tr>
<td>04</td>
<td>Group-4</td>
<td>42.70 ± 1.820 ***</td>
</tr>
<tr>
<td>05</td>
<td>Group-5</td>
<td>66.10 ± 1.518 ***</td>
</tr>
<tr>
<td>06</td>
<td>Group-6</td>
<td>22.40 ± 0.982**</td>
</tr>
</tbody>
</table>

Data are represent as mean ± SEM (n=6) and one-way analysis of variance (ANOVA) followed by Dunnett test for multiple column comparison. **P<0.01 and ***P<0.001 were considered to significant compared with control group.
of the test tube was then incubated at 54°C for 20 min. However, another pair was maintained at 0-5°C in an ice bath. The reaction mixture was placed for 15 min and the absorbance was measured at 540 nm using UV spectrophotometer.

**Hypotonic solution-induced hemolysis**

This experiment was performed in duplicate pairs with hypotonic solution. The stock erythrocyte suspension (30 µl) was mixed with hypotonic solution (5 ml) of plants extracts in the concentrations of 50, 100, 200, 300 and 500µg/ml. The vehicle was added to another tube separately as control. This mixture was incubated at 25°C for 10 min and absorbance was measured at 540 nm. The effect of test compounds was calculated as percentage inhibition or acceleration of hemolysis in both tests using following equation:

\[
\text{Percentage acceleration or inhibition of hemolysis} = 100 \left(\frac{\text{OD2-OD1}}{\text{OD3-OD1}}\right)
\]

Where, OD1 and OD2 was Test samples in isotonic and hypotonic solutions, respectively and OD3 was control sample in hypotonic solution.

**Statistical analysis**

Data are represent as mean ± SEM (n=6) and one-way analysis of variance (ANOVA) followed by Dunnett test for multiple column comparison. *P<0.05, **P<0.01 and ***P<0.001 were considered to be less significant, significant and more significant respectively when all groups were compared with control group.

**RESULTS**

Percentage yield of ethanolic extract of EA was 12.68% (w/w). Generally AChE activity is raised chronically with the administration of scopolamine. The tested extracts were effective in reducing the activity of AChE in comparison to the control group. Piracetam showed significant activity i.e. 15.17 units/L. Results from Table 1, ethanolic extract of EA showed a protective effect against scopolamine-induced degradation of ACh. Study revealed that extract has anti-inflammatory activity in the form of heat hemolysis was significantly reduced at 50 and 100 µg/ml while hypotonic solution induced inhibition was shown significant at 50, 100 and 200 µg/ml. Extract was effective in inhibition of hemolysis in a dose-dependent manner (Figure 1).

**DISCUSSION**

The role of ACh is very important in peripheral and central nervous systems both; In peripheral nervous system, it performs the locomotor activity while in central nervous systems, it help in the formation of memory. In brain, deficiency of ACh or depletion of ACh causes various neurodegenerative diseases in which AD is one of them.21 AD is an irreversible neurodegenerative disorder primarily targeting elderly populations.22 The level of ACh is decrease due to AChE which affects the cognition and progressed with locomoter disfunctioning. Dementia is major symptom of AD patients. ACh also affects the locomotor activity due to nicotinic acetylcholine receptors.23,24 Rotarod and actophotometer experimental models are used to determine the motor coordination of rodents. This method is sensitive to evaluate cerebellar dysfunction.25,26

Inflammation is a biological defense mechanism and secondary response of brain. During inflammation, pro-inflammatory mediators i.e interleukin-1β, IL-6, IL-8 are activated. Some other like tumor necrosis factor-α, reactive oxygen species, nitric oxide and prostaglandins are also activated. Over production in long time cause degenerative diseases i.e. asthma, cancer, arthritis, atherosclerosis and AD etc.27 Generally plants are the factory of metabolites in which phenolic components play vital role. Phenolic components have multiple activities in which anti-inflammatory activity is very important. They inhibit either the production or the action of pro-inflammatory mediator. Some other component can up/down regulated transcriptional factors like nuclear factor-KB or Nrf-2 in inflammation and antioxidant pathway. Study reveals that a high intake of phenolic rich food decreased rate of chronic disease such as cardiovascular disease, diabetes, Parkinson disease and AD etc.28

EA is traditionally claimed for memory enhancement. Its activity showed their scavenging properties in brain. However, the EA showed more significant results at 200 mg/kg in comparison to other doses. Results were found that ethanolic extract of EA were effective in inhibition of hemolysis in both heat induced and hypotonic solution experimental models. The anti-inflammatory activity of plant at various doses shows new criteria for determining the effective dose.

**CONCLUSION**

Ethanolic extract of EA showed a protective effect against scopolamine-induced degradation of ACh. Study revealed that extract has anti-
inflammatory activity which is very important for treatment of neurodegenerative diseases. It can be concluded from the study that EA possess highly significant anti-inflammatory activity in dose of 200 mg/kg so that it can be used as therapeutics against inflammation in several CNS diseases.

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CONFLICT OF INTEREST

The authors declare no Conflict of interest.

REFERENCES