“ANALGESIC ACTIVITY OF HYDRO ALCOHOLIC EXTRACTS OF STEMS AND ROOTS OF EUPATORIUM TRIPLINERVE VAHL.”

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KEYWORDS:

Eupatorium triplinerve, Ayapana, Stem and root extracts, Analgesic activity.

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ABSTRACT

Eupatorium triplinerve commonly known as ayapana is a most important plant used in traditional medicine. The plant is considered to be a therapeutic agent for the treatment of various diseases. The main aim of this research was to study the analgesic activity of the stem and root extracts. The hydroalcoholic extract of stem and roots of E.triplinerve was subjected to preliminary phytochemical screening. Acute toxicity studies were carried out in Swiss albino mice. Analgesic activity was evaluated by well established models like acetic acid induced writhing. Acute toxicity studies showed that the stem and root extracts was non-toxic upto a maximum dose of 2000 mg/kg body weight. The hydroalcoholic extract of the stem and root exhibited significant inhibition of acetic acid induced writhing. The present study indicates that the stem and root extracts of Eupatorium triplinerve has potential analgesic activity.
1. INTRODUCTION:

*E. triplinerve* is a member of large Asteraceae plant family. Ayapana is an ornamental erect Perennial herb with aromatic leaves. The leaves of *Eupatorium triplinerve* are reported to be useful in pain and inflammatory disorder. The essential oil from the plant has been reported to possess a number of medicinal properties such as central nervous system (CNS) depressant, analgesic and sedative effects. The methanol extract of ayapana leaves shows antifungal activity and potent antibacterial activity. Ayapana is a rich source of naturally occurring coumarin chemicals. Pain is a complex unpleasant phenomenon composed of sensory and emotional experiences associated with actual and potential tissue damage. Analgesics is defined as the agents which selectively relieve pain by acting in CNS or by peripheral pain mechanisms. Here the acetic acid induces the writhing. Writhing means the twisting body movements as in pain or struggle or abdominal constrictions. The present study has been designed to investigate the hydro alcoholic extracts of the stem and root, exhibits significant inhibition of acetic acid induced writhing, therefore it have analgesic activity.

2. MATERIALS AND METHODS:

2.1 collection of plant material:

The whole plant of *Eupatorium triplinerve* were collected from Thrissur district in Kerala in month of January 2016.

2.2 Animals

Swiss albino mice (20-25gm) were taken from the institutional animal house. The animals are fed with food and water. All the experiments were conducted according to the CPCSEA.

2.3 Preparation of extract

**Maceration**

The stem and roots of ayapana were collected, shade dried for 7 days. Stem and roots are crushed in grinder then were extracted with 80% aqueous ethanol by double maceration process at room temperature. The extract was filtered and then filtrate was concentrated.

2.4 Preliminary chemical tests

The hydro alcoholic extract of stem and roots of ayapana was subjected to phytochemical screening and plant extract contain alkaloids, carbohydrates, saponins, flavonoids, tannin, and glycoside using appropriate reagents.

2.5 Acute toxicity study

Acute oral toxicity study for the test was carried out according to Organization for Economic
Co operation and Development OECD 423. The test procedure is to minimize the no. of animal required to estimate the oral toxicity. Food was withheld for the study with a dose of 2000mg/kg of body weight. The animals are weighed and the extract was administered in a single dose as 1% suspension in Carboxy methyl cellulose by oral intubation. After dosing periodically during the first 24 hours daily after for a total of the 14 days. The LD 50 of the compound was estimated to be more than 2000mg/kg, so that doses of 100, 200, 400, 1000mg/kg orally were safe for the dose.

2.6 Acetic acid induced abdominal constriction

Acetic acid induced writhing method was adopted for evaluation of analgesic activity. Writhing is defined as a stretch, tension to one side, extension of hind legs, contraction of the abdomen, any writhing is considered as a positive response.

Mice (20-25gm) of either sex were divided to 3 different groups each containing six animals and marked (food, but not water should be withheld). Mice’s were orally treated with hydro alcoholic extract of eupatorium triplinervis, after 60 minutes later acetic acid (0.6% v/v in saline) volume 0.1ml/10g was injected intra peritoneally. The numbers of abdominal constrictions are counted for 15 minutes following acetic acid injection. Any significant reduction in number of abdominal constriction by treatment compared to the vehicle treatment and standard treatment.

Percentage of inhibition was evaluated using following formula: (C-T/C) x100

where C is the number of abdominal constrictions recorded in vehicle treated animals and T is the number of abdominal constrictions in the treatment group

Statistical analysis

All the values of in vivo analgesic studies of hydro alcoholic extract of ayapana were expressed as mean and standard error of mean (S.E.M) and were examined for significance by ANOVA (Analysis of variance) and groups were compared by Dunnet’s test for individual comparison of groups with control. P value were measured moderate significant at P<0.01, <0.001 level

3. RESULTS:

Phytochemical screening

The percentage yield of hydro alcoholic extract of the stems and roots of ayapana was found to be 9.8% w/w. The chemical tests indicate the presence of alkaloids, carbohydrates, saponins, flavonoids, tannin, and glycoside.
Acute toxicity studies
In acute oral toxicity studies no mortality was recorded in these animals up to 14 days. Thus the extract was non toxic up to 2000mg/kg.

Analgesic activity
The writhing was counted. There is a significant reduction in number of writhing between the vehicle treated and standard treated animals.

4. DISCUSSION:
E. triplinerve Vahl (synonyms: Eupatorium ayapana, Ayapana triplinervis) belongs to the Asteraceae family and is commonly known as ayapana in Hindi. The present study was conducted to scientifically validate the traditional claims of Eupatorium triplinerve with particular reference to its analgesic activity. In acute toxicity testing no mortality was observed in mice even in a dose of 2g/kg of petroleum-ether extract of E.triplinerve which indicates the safe nature of the extract. Acetic acid induced abdominal constriction is regarded as a very sensitive method which induces minimal noxious stimulus. The advantage of this method is that even compounds with weaker analgesic property can be detected from the results of this test. The acetic acid causes pain by liberating endogenous substances such as serotonin, histamine, prostaglandins, bradykinins and substance P. Here we conclude that the hydro alcoholic extracts of stems and roots of ayapana shows analgesic activity.

Table no.1 shows there is an decrease in writhing by administration of ayapana extract with reference to the vehicle treated mice and plant extracts shows analgesic activity.

<table>
<thead>
<tr>
<th>Group</th>
<th>Treatment</th>
<th>Dose</th>
<th>No. of wriths in 15 mins (mean±SEM)</th>
<th>Inhibition %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>Vehicle(carboxymethyl cellulose)</td>
<td>1.3ml</td>
<td>20.6±0.33**</td>
<td>-</td>
</tr>
<tr>
<td>Test</td>
<td>Plant extract</td>
<td>4.8mg in cmc orally</td>
<td>13.16±0.30**</td>
<td>36%</td>
</tr>
<tr>
<td>Standard</td>
<td>Diclofenac potassium</td>
<td>1mg orally</td>
<td>5.1±0.30**</td>
<td>75%</td>
</tr>
</tbody>
</table>

Table no.1 Effect of plant extract on acetic acid induced writhing in mice

n = 6. *P < 0.05, **P < 0.01 and ***P < 0.001 show a comparison of group # 2,3 vs. group # 1 (One-way ANOVA followed by Dunnett’s test).
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6. Conclusions
The traditional claims of the usefulness of *Eupatorium triplinerve* in the treatment pain has been scientifically validated by the results of the present study .The stem and root extracts of ayapana plant has analgesic activity. The study indicates that the data obtained will be basis or further studies and applications of this plant.

7. REFERENCES:


