Short Communication

Antidiarrheal Activity of Methanolic Extract of Leaves of Gymnosporia Emerginata

A. Suvarchala Kiranmai, Hemamalini. K*, P. Sumalatha, G Shashi Priya and Uma Vasireddy

1M. Pharmacy Student, Dept. of Pharmacology, Teegala Ram Reddy College of Pharmacy, Meerpet, Hyderabad,
2IPA Treasurer, A.P, Hyderabad.

ABSTRACT

The methanolic leaf extract of Gymnosporia emerginata showed significant antidiarrheal activity on castor oil induced diarrheal model in rats. These results obtained revealed that the leaf extract possess pharmacological activity against diarrhea and may possibly explain the use of the plants in traditional medicine.

Key words: Atropine, Antidiarrheal, Gymnosporia emerginata.

INTRODUCTION

In developing countries, the majority of people living in rural areas almost exclusively use traditional medicines in treating all sorts of disease including diarrhea. There are large number of epidemiological and experimental evidence pertaining to worldwide acute-diarrheal disease, which is one of the principal causes of death in the infants, particularly in malnourished and which is of critical importance in developing countries (Syder et al, 1982). Thus it becomes important to identify and evaluate commonly available natural drugs as alternative to currently used anti-diarrheal drugs, which are not completely free from adverse effects (Hardman, et al., 1992). Several studies have evaluated the effectiveness of some traditional medicines in treating diarrhea in all different continents (Mukherjee et al., 1998). India has a great environmental and biological diversity compared with the rest of the world. A range of medicinal plants with anti-diarrheal properties has been widely used by the traditional healers; however, the effectiveness of many of these anti-diarrheal traditional medicines has not been scientifically evaluated.

The present study was carried out in rats with methanolic leaf extracts of Gymnosporia emerginata. The purpose of the present study was to find out the antidiarrheal activity of methanolic extract of Gymnosporia emerginata in experimentally induced acute diarrhea in rats.

MATERIALS AND METHODS

Plant material

Fresh leaves of Gymnosporia emerginata (Celastraceae) collected from Tirupathi S.V. University; the plant was identified confirmed and authenticated by comparing with an authentic specimen by a botanist Dr. Madhavachetty. K., A voucher specimen has been deposited at S.V. University Tirupathi, India. Dried and coarsely powdered Gymnosporia emerginata leaves (500g) was repeatedly extracted with methanol. The extract was concentrated under vacuum and the residue was used in the experiments. The dried plant extracts were freshly re-dissolved in normal saline and given to adult albino rats fed a standard animal diet.

Animals

Male albino rats weighing 150-180g were used for castor oil induced anti-diarrheal activity. The animals were acclimatized to room temperature (28±5°C) with relative humidity of 55±5% in a standard wire meshed plastic cages for 4 to 5 days prior to
commencement of the experiment. All animals were fed standard animal feed and tap water ad libitum before the experiments. The animals were maintained as per the norms of CPCSEA (Regd.No.1447/PO/a/11/CPCSEA) and cleared by CPCSEA and institutional ethics committee (Teegala Ram Reddy College of Pharmacy) each experimental group consisted of six animals housed in separate cages.

**Castor oil induced diarrhea**

Rats were divided into three groups of six animals each, diarrhea was induced by administering 1ml of castor oil orally to rats Group-I served as control (2ml / kg, i.p saline), Group-II received atropine (3mg / kg, i.p.) served as standard and Group-III received (300 mg / kg, i.p.) of Gymnosporia emerginata 1hour before castor oil administration. The animals were housed individually in a metal cage lined with white paper. Fecal output was assessed by collecting the fecal material for 8hours after drug administration, and the total weight in each group was recorded. The percentage protection was calculated as follows (Wouters et al. 1978).

\[
\text{Total weight of stool in control animal} - \frac{\text{Total weight of stool in drug treated animal}}{\text{Total weight of stool in control animals}} \times 100
\]

**Statistical analysis**

The experimental results are represented as mean ±SEM (Standard error of the mean). ANOVA test was used for the evaluation of data and ***p<0.001 and **p<0.02 accepted as significant.

**RESULT**

**Castor oil induced diarrhea**

30 min after administration of castor oil the diarrhea was clinically apparent in all the animals of control group, for the next 8 hours was markedly reduced by the intra-peritoneal injection of atropine, 3 mg / kg (57.67) (Table-1).

**Table 1:** Effect Gymnosporia emerginata extract on castor oil induced diarrhea in rats.

<table>
<thead>
<tr>
<th>S.No</th>
<th>Treatment</th>
<th>Frequency of defection (times / 8hr)</th>
<th>Total weight stools mg for 8h</th>
<th>% inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control-Castor oil(1ml/p.o) ± saline (2mg/kg,i.p)</td>
<td>9.66 ±1.42</td>
<td>3.19 ±6.2</td>
<td>-</td>
</tr>
<tr>
<td>2</td>
<td>Standard-Castor oil ± Atropine (3mg/kg,i.p)</td>
<td>2.5 ±5.248</td>
<td>1.35 ±0.27***</td>
<td>57.6</td>
</tr>
<tr>
<td>3</td>
<td>Castor oil± M.E Gymnosporia emerginata(300 mg/kg,i.p)</td>
<td>2.96 ±0.516</td>
<td>2.12 ±0.31**</td>
<td>33.54</td>
</tr>
</tbody>
</table>

Values are expressed as mean ±SEM from the experiments. ***p<0.02, **p<0.001 when compared with Castor oil ± saline treated group.

**DISCUSSION AND CONCLUSION**

Diarrhea results from an imbalance between the absorptive and secretory mechanisms in the intestinal tract, resulting in an excess loss of fluid in the feces. In some diarrheas, the secretory component predominates, while other diarrheas are characterized by hypermotility. The use of castor oil induced diarrhea model in our study is logical because the autocoids and prostaglandins are involved these have been implicated in the causation of diarrheas in man (Horton, et al., 1968). The liberation of ricinoleic acid from castor oil results in irritation and inflammation of intestinal mucosa, leading to release of prostaglandins, which stimulates motility and secretion (Pierce, et al., 1971). The results of the present study showed that the extract of Gymnosporia emerginata produced a statistically significant reduction in the severity and frequency of diarrhea produced by castor oil. It is also noted that the extract significantly inhibited castor oil induced intestinal fluid accumulation and the volume of intestinal content, dose dependently more than Atropine. The extracts significantly reduced the castor oil induced diarrhea in rats. In this study, atropine produced a significant reduction in the number of stools possibly due to its anti-cholinergic effect (Brown et al., 2000). Castor oil is also reported to induce diarrhea by increasing the volume of intestinal content by prevention of the reabsorption of water. The liberation of ricinoleic acid results in irritation and inflammation of intestinal mucosa, leading to release of prostaglandins which results in stimulation of secretion. Thereby prevents the reabsorption of NaCl and H₂O (Galveg et al, 1993).

The antidiarrheal activity of the extract may also be due to the presence of denature proteins forming protein tannates. Protein tannates make the intestinal mucosa more resistant and reduce secretion (Tripathi, 1994). The secretory diarrhea is associated with an activation of CI channels, causing CI efflux from the cell, the efflux of CI results in massive secretion of water into the intestinal lumen and profuse watery diarrhea (Ammon et al., 1985).

The extract may inhibit the secretion of water into the lumen by reverting this mechanism.

Anti-dysenteric and anti-diarrheal properties of medicinal plants were found to be due to tannins, alkaloids, saponins, flavonoids, sterols and/or triterpene and reducing sugar (Longanga-otshudiet et al., 2000). The phytochemical analysis of the extracts revealed the presence of triterpinoids and may mediate the antidiarrheal property of Gymnosporia emerginata extracts. The extract resulted in a marked reduction in the number of diarrhea stools. This signifies the usefulness of this model and the clinical effect of the extract.

**REFERENCES**


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