

IN-VIVO ANTI-INFLAMMATORY ACTIVITY OF *STROBILANTHES CILIATUS* NEES

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ABSTRACT

The anti-inflammatory activity of ethanolic extract of *Strobilanthes ciliatus* aerial parts was studied using carrageenan induced paw oedema model in wistar rats. The test extract (200, 400 and 800 mg/kg, p.o) showed significant anti-inflammatory activity when compared to standard drug diclofenac and vehicle control.

Keywords: *Strobilanthes ciliatus*, aerial parts, paw edema, anti-inflammatory, diclofenac.

INTRODUCTION

The composition of the internal environment is maintained within narrow limits, and this fairly constant state is called *homeostasis*. Despite the progress made in medical research during the past decades, treatment of many serious diseases is still problematic. The most common presentation of patient to the doctor is pain and inflammation. Inflammation is a host defence mechanism of the body and it's an essential immune response that enables the body to survive during infection or injury and maintains tissue homeostasis in noxious conditions. But in some conditions appears to be no resolution and a chronic state of inflammation develops that may last the life of the individual. Such conditions include the inflammatory disorders osteoarthritis, rheumatoid arthritis, inflammatory bowel diseases, retinitis, multiple sclerosis psoriasis and atherosclerosis¹. Treatment of inflammation is a debate as the conventional NSAIDs are commonest to cause Adverse Drug Reactions. Hence there is ongoing research to develop safer and more effective drugs for the therapy of inflammation. Traditional plants play a very important role in the discovery of new drugs. *Strobilanthes ciliatus* of Acanthaceae family is a highly potential medicinal plant in ayurveda in the treatment of inflammatory disorders^{2,3}. As per literature survey information, there is no scientific evidence of pharmacological activity of the plant. Hence, the present study has been undertaken to evaluate the *in-vivo* anti-inflammatory activity of *Strobilanthes ciliatus*.

MATERIALS AND METHODS

Plant material

The aerial parts of *Strobilanthes ciliatus* were collected from Nilambur, Mallapuram district, Kerala, India, identified and authenticated. The sample specimen was deposited, at the Department of Pharmacology, Chalapathi Institute of Pharmaceutical Sciences, Lam, Guntur, Andhra Pradesh, India.

Preparation of plant extracts⁴

Air dried pieces of aerial parts of *Strobilanthes ciliatus* were thoroughly cold percolated and extracted with ethanol (60- 80 °C) for 72 hours. The extraction was filtered, concentrated and stored in the dark at +4°C until tested.

Qualitative phytochemical analysis⁵

The preliminary chemical tests were carried out for the ethanolic extract of *Strobilanthes ciliatus* to identify the presence of various phytoconstituents.

IN-VIVO STUDIES

Experimental animals

Adult Wistar albino rats (150-180 g) of either sex were procured from the laboratory animal house, Chalapathi institute of pharmaceutical sciences, Guntur, Andhra Pradesh, India and used in the study. The animals were kept under standard environmental conditions of room temperature (220 ±20C), relative humidity (50% ± 5%) and 12 h light and dark cycle. The animals were housed in the rat cages and provided feed (commercial pellets contain a balanced ration obtained from the

Sri Venkateswara Enterprises, Bangalore) and water *ad libitum*. All the animals were acclimatized to the laboratory environment 5 days prior to the experiment. The animals were fasted overnight just prior to the experiment but allowed free access to drinking water. All the experiments were carried out in accordance with the guidelines of the Institutional Animal Ethics Committee. The study was conducted after obtaining ethical committee clearance from the Institutional Animal Ethics Committee.

I. Acute toxicity study

3 female Wistar rats were selected for the study. The overnight fasted animals (with water *ad libitum*) were administered with ethanolic extract of *Strobilanthes ciliatus* at a single dose of 2000 mg/kg body weight by orally. The dose volume was fixed at 10ml per kg body weight. The animals are observed for 0min, 30 min, 1 hr, 2hr, 4hr, 6hr, and thereafter every day for 14 days. Food was withheld for a further 3-4 hours after administration of test extract and was observed for signs of toxicity. The body weight of the rats before and after administration are noted that changes in skin and fur, eyes, mucous membranes, respiratory, circulatory, autonomic and central nervous system and motor activity and behavior pattern was observed and also sign of tremors, convulsions, salivation, diarrhoea, lethargy, sleep and coma were noted. At the end of the 14th day the animals were sacrificed with excessive ether anesthesia

and dissected for examination of vital organs like brain, liver, kidney, lungs, and heart for pathological changes. For further confirmation the above procedure was repeated on another set of 3 female Wistar rats ⁶.

2. Efficacy studies

Anti-inflammatory activity

Carrageenan induced rat paw edema method⁷

5 groups was formed with 6 wistar rats in each group.

Test agent (Ethanolic extract of aerial parts of *Strobilanthes ciliatus*) and diclofenac (100mg/kg) were administered (p.o) 30 minutes before 1% carrageenan injection (0.1ml) into the sub plantar area of the right hind paw. The volumes of injected and contralateral paws were measured 1, 2 and 3 hours after induction of edema by using digital plethysmometer (Inco, Chennai). The treatment was given in the following manner:

- I Group – Control (Normal saline 10 ml/kg, p.o)
- II Group – *Strobilanthes ciliatus*(200 mg/kg,p.o)
- III Group – *Strobilanthes ciliatus*(400 mg/kg,p.o)
- IV Group – *Strobilanthes ciliatus*(800 mg/kg,p.o)
- V Group – Diclofenac (100 mg/kg,p.o).

Statistical analysis

Results was analysed by one way ANOVA, followed by Turkey's multiple comparison test, 'p' value less than 0.05 was taken as significant (Table 1).

Table 1: Anti-inflammatory action of ethanolic extract of *Strobilanthes ciliates* aerial – parts in carrageenan induced rat paw edema

Group	Dose (mg/kg,p.o)	Swelling volume (ml)		
		1(h)	2(h)	3(h)
Control	10	1.51±0.03	1.56±0.01	1.62±0.17
<i>Strobilanthes ciliatus</i>	200	1.12±0.09*	1.30±0.05*	1.34±0.03*
<i>Strobilanthes ciliatus</i>	400	0.93±0.04*	1.17±0.08*	1.19±0.02*
<i>Strobilanthes ciliatus</i>	800	0.85±0.01*	0.95±0.07**	0.98±0.05*
Diclofenac	100	1.08±0.02**	0.99±0.04**	0.91±0.06*

Values are expressed as Mean ±S.E.M. p* $<$ 0.05, p** $<$ 0.01 considered significant (n=6) when compared to control.

RESULTS AND DISCUSSION

Preliminary phytochemical analysis revealed the presence of carbohydrates, triterpenoids, phytosterols, flavonoids, and tannins in the ethanolic extract of the aerial parts of *Strobilanthes ciliatus*.

The ethanolic extract of the aerial parts of *Strobilanthes ciliatus* was found to be safe and no mortality of the rats was observed at the dose of 2000mg/kg for 14 days in acute toxicity study. So the LD₅₀ of the *Strobilanthes ciliatus* will be $>$ 2000mg/kg body weight.

The ethanolic extract of the aerial parts of *Strobilanthes ciliatus* (200, 400, 800 mg/kg, p.o) showed dose dependent reduction of carrageenan induced paw edema in rats when compared to

control. The lower the rat paws volume the better the anti-inflammatory activity.

The standard drug Diclofenac (100 mg/kg,p.o) produced inhibitory activity which was close to that of ethanolic extract of the aerial parts of *Strobilanthes ciliatus*(Table 1). All the treatments were significant (p $<$ 0.05, p $<$ 0.01).

CONCLUSION

The present study showed that ethanolic extract of aerial parts of *Strobilanthes ciliates* can be effectively used as a anti-inflammatory agent.

ACKNOWLEDGEMENT

The authors are thankful to University Grants Commission (UGC) for financial assistance of this

project. And also thankful to Sri. Y.V.Anjaneyulu, President, Chalapathi educational society, Guntur for providing necessary facilities to carry out the present research work.

REFERENCES

1. Dinarello C. Anti-inflammatory agents: Present and future. *Cell*. 2010;140:935-50.
2. Warriar PK, Nambiar VPK and RamanKutty C. *Indian medicinal plants*. Madras: Orient Longman Ltd. 1994;5:142-5.
3. Thomas J, Joy PP, Mathew S, Skaria BP, Duethi PP and Joseph TS. *Agronomic Practices for aromatic and medicinal plants*. Calicut: Kerala Agricultural University. 2000;124.
4. Mukherjee PK. Quality control of herbal drugs, an approach to evaluation of Botanicals. 1st Ed. New Delhi: Business horizons. 2002;379-401.
5. Raaman N. *Phytochemical techniques*. 1sted. New Delhi: New India publishing agency. 2006;19-24.
6. OECD Guidelines for the testing of chemicals, No.425. Acute oral toxicity-modified up and down procedure. Paris, France : 2001.
7. Winter CA, Risley EA and Nuss CW. Carrageenan induced edema in hind paw of rat as an assay for anti-inflammatory drugs. In: *Proc Soc Exp Biol Med*. 1962;11:544-7.
8. Venu P. *Strobilanthes Blume (Acanthaceae) in peninsular India*. 1st Ed. Kolkata: Botanical Survey of India; 2006;89-93,129-32.