IAMJ

Research Article International Ayurvedic Medical Journal ISSN:2320 5091

ASSESSMENT OF THE ANTI-INFLAMMATORY EFFECT OF A FOLKLORE MEDICINAL PLANT – TILIACORA ACUMINATA MIERS

Vadakkath Anjaly Das¹, Narayanan Manojkumar²

¹MD Scholar, ²Associate professor, Dept. of Dravyaguna Vijnana, VPSV Ayurveda College, Kottakkal, Kerala, India

ABSTRACT

The plant *Tiliacora acuminata* is a Menispermaceae member found all over India. The plant is well known for its folklore uses as an analgesic and anti-inflammatory drug. Diclofenac sodium was used as the standard drug and the test drug used was the *kashaya* (decoction) of the whole plant. Both the test drug and the standard drug in the corresponding doses were administered orally to the respective groups for 5 consecutive days. 24 hours after the 5th dose oedema was induced in the left hind paw of all the animals using 20µl of 1% carageenan in normal saline. The paw thickness measurement was taken using vernier calipers before induction, just after injection of carageenan, then every hour upto the 6th hour and then the 24th hour. The acute antiinflammatory study done in carageenan induced paw oedema model showed significant results in the high dose treated group comparable with that of the standard drug Diclofenac sodium. **Keywords:** *Tiliacora acuminata*, Anti-inflammatory, paw oedema, acute inflammation

INTRODUCTION

The plant *Tiliacora acuminata* Miers is a creeper of the family Menispermaceae. This plant is commonly known as *Vallikanjiram* in Malayalam though it is not in any way related to *kanjiram* (Strychnos nuxvomica of Loganiaceae family) except for



some resemblance in its foliage.¹ This plant has not been described in any of the classical *āyurvedic* texts. *Ācarva* P.V

Sharma gives the botanical identity for a drug *Krşņavetra* as *Tiliacora acuminata*

Miers. This name indicates that the plant has a dark color and is like vetra i.e, bamboocane. It is seen that the branches of Tiliacora acuminata is dark in color and the branches are highly flexible. The long and flexible branches are used for thatching and basket work. The stem is used as rough cordage in South Vietnam.²Krsnavetra has references in Cakraduttam Kustha cikitsa and Vangasēna Samhita Visa pratishedam³ This plant is said to be used by some tribal communities as an antidote for snake venom⁴. In Bangladesh some folk practitioners are using this drug for wound healing⁵. In Kerala many traditional practitioners are using this drug for treatment of conditions like

Vadakkath Anjaly Das L Narayanan Manojkumar: Assessment Of The Anti-Inflammatory Effect Of A Folklore Medicinal Plant – Tiliacora Acuminata Miers

vātakaņţakam which is a very painful condition.

Pharmacological study Collection and processing of drug

The plant Tiliacora acuminata Miers was collected from Puthuparambu, Kottakal. The whole plant collected was washed 3-4 times in water and shade dried. The shade dried plant was crushed and then made into powder. 20g of this powder was taken and soaked in 60 ml of double distilled water. Then the decoction was prepared by standard Avurvedic procedures. i.e 8 times water (160ml) was added and was reduced to one fourth by boiling. This decoction was filtered and centrifuged. The supernatant was collected and further evaporated in a water bath till the extract was dry. The yield was 6.2g. This was reconstituted in 35ml of autoclaved double distilled water and stored in freezing temperature. This prepared drug was used for the whole experiment.

Materials

Reagents and instruments

- Diclofenac sodium
- Ranbaxy
- 1% carageenan
- 0.1g carageenan was dissolved in 10ml normal saline.
- Vernier calipers

Animals

Male Balb/c mice (4-8 weeks old, 20-30g body weight) were obtained from the animal breeding section of Amala Cancer Research Centre, Thrissur. The animals were maintained in well ventilated polypropelene cages under standardized environmental conditions (22 -28⁰ C, 60 – 70% relative humidity, 12 hr dark / light cycle) and fed with standard mouse feed (Lipton India) and water *ad libitum*. All the animal experi-

ments were carried out at Amala Cancer Research Centre by prior permission of Institutional Animal Ethics Committee (IAEC) (No.149/1999/CPCSEA).

Anti-inflammatory study⁶

30 male balb/c mice aged 4-6 weeks weighing 20-25g were selected and grouped as follows with 6 animals in each group Control: which received no treatment? Standard: which received Diclofenac sodium in the dose of 10mg/kg body weight Group1: This received the test drug in a dose of 125mg/kg body weight

Group 2: This received the test drug in a dose of 250mg/kg body weight

Group 3: This received the test drug in a dose of 500mg/kg body weight

Diclofenac sodium and the test drug in the corresponding doses were administered orally to the respective groups for 5 days. 24 hours after the 5th dose oedema was induced in the left hind paw of all the animals using 20 μ l of 1% carageenan in normal saline. The paw thickness measurement was taken using vernier calipers before induction, just after injection of carageenan, then every hour upto the 6th hour and then the 24th hour.

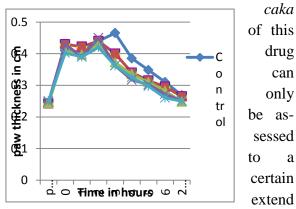
Statistical analysis: The statistical analysis of the data was done by using Instat Graphpad software

Results:Anti-inflammatory study: The anti-inflammatory effect of *Tiliacora acu-minata* Miers decoction was assessed by carageenan induced paw oedema method by measuring the paw thickness using vernier calipers.The drug was assessed for its acute anti-inflammatory effect in carageenan induced acute paw oedema model in Balb/c mice. The standard drug used was Diclofenac sodium and the test drug was administered in 3 different doses- half of therapeutic

dose (125mg/kg body weight), therapeutic dose (250mg/kg body weight) and double therapeutic dose (500mg/kg body weight). The assessment was done by measuring the paw thickness every hour up to the 6th hour and in the 24th hour using vernier calipers. The paw thickness measured up to the 3rd hour did not have any significant difference between the groups. All the groups showed highly significant reduction in paw oedema when compared to the control group (p<0.001). The groups which received the test drug in the therapeutic dose and in double the therapeutic dose showed almost same effect as the standard drug. The half dose group showed significant reduction in paw oedema when compared with control but it was not as effective as the therapeutic dose, double dose and standard drug. In the 4th hour also all the groups showed highly significant reduction in paw oedema when compared to the control group. All the groups which received test drug showed anti-inflammatory effect comparable with the standard drug. In the 5th hour of assessment. also all the groups showed highly significant reduction in paw oedema when compared to the control group. The double dose group and the therapeutic dose group showed antiinflammatory properties almost as same as the standard drug. The half dose group showed a lesser anti-inflammatory effect than the other groups. In the 6^{th} hour, the half dose group did not show any significant reduction in paw oedema when compared with the control group. All the other groups showed significant reduction in paw oedema when compared to the control group. In the 6th hour only the double dose group showed anti-inflammatory activity comparable to the standard drug. In the 24th hour of assess-

ment, the half dose group did not show any significant reduction in paw oedema. All other groups showed significant reduction in paw oedema when compared to the control group. Among those the group which received the therapeutic dose and double dose showed highly significant effect (p<0.001). the anti-inflammatory effects of the therapeutic dose group and the double dose group were comparable to the effect of the standard drug. Thus the statistical analysis showed that the drug in all the 3 doses had significant effect in reducing the paw thickness from the third hour of injection onwards. The double dose group and the therapeutic dose treated groups showed significant, sustained results up to the 24th hour of assessment. These two groups showed antiinflammatory effects which were comparable with that of the standard drug.

Results and discussion: Regarding the plant *Tiliacora acuminata* Miers, the plant is a new one to the $\bar{A}yurvedic$ pharmacopoeia. So the $\bar{A}yurvedic$ pharmacological properties are not mentioned anywhere. The *rasa pan*-



of surety. The *rasa* of the drug when assessed by the *nipāta* method was found as *tikta* by the volunteers without any difference in opinion. So being *tikta* we can assume that the drug may have *laghu*, *rukşa gunas* and hence may cause *pitta śamana* property in the body. The drug was found to have a very good anti-inflammatory property which may be attributed to this. From assessing the type of reaction with distilled water, it was found to be exothermic, so it may be inferred to be an uşna Vīrya drug. Thus it may also have a kapha vāta śamana property. It may be considered as having katu vipaka. Considering the above mentioned findings, we can infer that the drug being tikta rasa, ushna Vīrya and katu vipaka may possess a pācana property. In s'opha samprapti all the 3 dosas have their own roles to play. Pitta causes pāka, kapha causes the puva and vāta causes ruja in s'opha. So this drug alleviates the *paka* in *s'opha* by doing pitta s'amana, alleviates the swelling by reducing kapha thus causing the reduction in paw thickness.

Tiliacora acuminata Miers⁷, a plant belonging to Menispermaceae, is known to be the biggest source of diphenylbisbenzylisoquinoline (DBBI) which are well known for their pharmacological activities such as antitumor (Kupchan etal 1973), antimicrobial (Wu, W.-N. etal 1976) and hypotensive effects (Joshi etal 1974, Wu, W.-N. etal 1976). It⁸ has been shown that prostaglandins, histamine, serotonin, and bradykinin are mediators of different phases of carrageenan-induced oedema. Di Rosa⁹ et al., Capasso ^{10et} al., and Salvemini¹¹ et al. have also reported the involvement of histamine, 5hydroxytrptamine, bradykinin, prostaglandin, and nitric oxide in carrageenan-induced paw oedema. The alkaloids, flavanoids and tannins present in the drug may be contributory to the anti-inflammatory effects.

Another ¹² study indicated that the extracts ethanol (70%), hexane, ethyl acetate and methanol extracts of *Tiliacora acuminata*

showed good antioxidant activity. Among the all the extracts ethyl acetate extract showed better activity. Although the antioxidant activities found invitro experiment were only indicative of the potential health benefit, these results remain important as the first step in screening anti-oxidant activity of *Tiliacora acuminata*. Antioxidant activity can be considered with potential health benefits in many diseases like Parkinsonism, autoimmune disorders and also in diseases where inflammation is considered as the basic pathology. So this is also indicative of its anti-inflammatory effect.

Acknowledgement:

The authors are thankful to DR.Ramadasan Kuttan, Research director of Amala Cancer Research Centre, Thrissur.

Source of support: Nil

Conflict of interest: None Declared

REFERENCES

- V V Sivarajan, Indira Balachandran, Āyurvedic drugs and their plant sources, Newdelhi, Oxford and IBH Publishing co. pvt ltd,1994 pg 218
- 2. The wealth of India- Raw materials, Newdelhi, NISCAIR press, 1976, Vol-X, pg 247,248
- 3. P V Sharma, Classical uses of medicinal plants, Varanasi, Chaukhambha Visvabharati,2004, pg 112
- 4. B. Sandhya Sri and T.V.V. Seetharami Reddi, Traditional phyto-antidotes used for snakebite by Bagata tribe of Eastern Ghats ofVisakhapatnam district, Andhra Pradesh, International Multidisciplinary Research Journal 2011,[cited on 17/3/2013], 1(6):42-45, http://irjs.info/
- Rezwana Afroz, Nabila Islam, Kakoli Rani Biswas, Tasneema Ishika, Mehreen Rahman, Auditi Swarna, Tania Khan, Mirza Nipa Monalisa, Syeda Seraj, Md. Atiqur Rahman, Sadia Moin Mou, Mohammed Rahmatullah. Medicinal Plants

Vadakkath Anjaly Das L Narayanan Manojkumar: Assessment Of The Anti-Inflammatory Effect Of A Folklore Medicinal Plant – Tiliacora Acuminata Miers

Used by Folk Medicinal Practitioners in Three Randomly SurveyedVillages of Rajbari District, Bangladesh, American-Eurasian Journal of Sustainable Agriculture, 5(2): 226-232, 2011[cited on 28/12/2011]

- 6. H. Gerald Voghel, Drug discovery and evaluation, Second edition 2002, Chapter H pg-716
- Arnab Khasnobis, Tapan Seasl, Rajan Vedasiromani, Malaya Gupta, and Biswapati Mukherjee, Some pharmacological studies with Tiliacorine, diphenylbisbenzylisoquinoline alkaloid from Tiliacora racemosa, Natural product sciences 5(3) 142-147 (1999) httpimg.kisti.re.kroriginalVieworiginalV iew.jsp [cited on 9/9/2013]
- K. F. Swingle, "Evaluation for antiinflammatory activity," in Antiinflammatory Agents: Chemistry Pharmacology, R. A. Scherrer and M. W. Whitehouse, Eds., vol. 2, pp. 33–122, Academic Press, New York, NY, USA, 1974.
- M. Di Rosa, J. P. Giroud, and D. A. Willoughby, "Studies on the mediators of the acute inflammatory response induced in rats in different sites by carrageenan and turpentine," Journal of Pathology, vol. 104, no. 1, pp. 15–29, 1971.

- F. Capasso, B. Balestrieri, M. Di Rosa, P. Persico, and L. Sorrentino, "Enhancement of carrageenan foot edema by 1,10-phenanthroline and evidence for bradykinin as endogenous mediator," Agents Actions, vol. 5, no. 4, pp. 359–363, 1975.
- D. Salvemini, Z. Q. Wang, P. S. Wyatt et al., "Nitric oxide: a key mediator in the early and late phase of carrageenaninduced rat paw inflammation," British Journal of Pharmacology, vol. 118, no. 4, pp. 829–838, 1996.
- Narender Prasad Dasari*, Ganga Rao.B., Sambasiva Rao Ethadi, Mallikarjuna Rao.T., V.S.Praneeth.D., Protective effect of phytochemical constituents of *Tiliacora acuminate* against oxidative stress, *Medicinal Chemistry & Drug Discovery* 2012, 3(1) 58-64

CORRESPONDING AUTHOR

Dr. Anjaly Das V

MD Scholar, Dept of Dravyaguna Vijnana, VPSV Ayurveda College , Kottakkal, Kerala, India. **Email:** anjaly.ds@gmail.com

Source of support: Nil Conflict of interest: None Declared