

Anthelmintic Activities of Roots of *Cocos nucifera* and Aerial Parts of *Jasminum multiflorum*

DILIPKUMAR PAL*, SANDIP KUMAR PAHARI and ABHISHEK KUMAR MISHRA
Department of Pharmaceutical Chemistry, Seemanta Institute of Pharmaceutical
Sciences, Jharpokharia, Mayurbhanj-757 086, India
Fax: (91)(6791)222238; Tel: (91)(3244)243265 E-mail: drdilip2003@yahoo.co.in

The anthelmintic activities of different extract of roots of *Cocos nucifera* and aerial parts of *Jasminum multiflorum* were evaluated separately on adult Indian earthworm (*Pheretima posthuma*) and the activities were compared with standard drug piperazine citrate and albendazole. It was found that petroleum ether, benzene, chloroform and ethyl acetate extract of *Cocos nucifera* (PECN, BCN, CCN and EACN, respectively) and petroleum ether extract of *Jasminum multiflorum* (PEJM) exhibited dose dependant action and inhibition of spontaneous motility (paralysis) and death of earthworms. The results indicated that the PECN, BCN and CCN were more potent than EACN and PEJM.

Key Words: *Cocos nucifera*, *Jasminum multiflorum*, Piperazine citrate, Albendazole.

INTRODUCTION

Cocos nucifera (Nariyal in Hindi; Narikel, Dab in Bengali; Nadia, Paido in Oriya; Family : Palmae) is a tall tree, distributed throughout tropical islands, coasts, South America, Florida, Bahamas in North America, Southern India and Sri Lanka¹. Every part of the coconut palm is used by man. Seed is cooling, tonic, laxative, cardio tonic and useful in the treatment of leprosy, tuberculosis, liver complaints, piles, *etc.* Bark is good for teeth and scables. Flower is useful in diabetes, dysentery, urinary discharges, *etc.*^{2,3}.

Jasminum multiflorum (Kunda in Sanskrit, kundaphul in Bengali, Chemeli in Hindi; Family: Oleaceae) is a large scandent, tomentose shrub with young branches clothed with velvety pubescence, distributed throughout India. Dried leaves of the plant are good for indolent ulcer⁴. The flowers are useful in vitiated condition of pitta, inflammation, rheumatism and cephalalgia. The root is antidote to cobra venom^{5,6}. From literature survey, it was found that no detailed study has yet been done regarding the anthelmintic properties of roots of *Cocos nucifera* and aerial parts of

Jasminum multiflorum Andr. On preliminary testing, it was found that petroleum ether, benzene, chloroform and ethyl acetate extract of *Cocos nucifera* (PECN, BCN, CCN and EACN, respectively) and petroleum ether extract of *Jasminum multiflorum* (PEJM) showed significant anthelmintic activity compared to other extracts of them. Hence, in the present study, we have evaluated the anthelmintic activities of petroleum ether, benzene, chloroform and ethyl acetate extracts of *Cocos nucifera* (PECN, BCN, CCN, EACN) and petroleum ether extracts of *Jasminum multiflorum* (PEJM) to substantiate the folkloric claims.

EXPERIMENTAL

The roots of *Cocos nucifera* were collected from Jharpokharia, Orissa during the month of July and aerial parts of *Jasminum multiflorum* were obtained from various places of Midnapore district of W.B. during the month of October-November. The samples were authenticated by Dr. H.J. Chaudhury, Joint Director, Central National Herbarium, Botanical Survey of India, Howrah, W.B. The voucher specimens have been preserved in our laboratory for further references (DAS1 and DPS1).

Extraction: Shade-dried, powdered and sieved (40 mesh size) roots were extracted in succession with petroleum ether (40-60°C), benzene, chloroform and ethyl acetate using soxhlet apparatus. Similarly, the powdered aerial parts of *Jasminum multiflorum* were extracted first with petroleum ether (40-60°C) and then with ethanol. The extracts were evaporated to dryness. The trace amount of solvent which may be present within the solid mass of respective extracts, was removed under vacuum. The yield of PECN, BCN, CCN, EACN and PEJM were 0.33, 0.16, 0.35, 0.77 and 2.10 % w/w, respectively with respect to dry starting materials.

Evaluation of anthelmintic activities: Anthelmintic activities were evaluated for PECN, BCN, CCN, EACN and PEJM separately. The activity was tested according to method discussed in detail by Kalesaraj and Kurup⁷. *Pheretima posthuma* (earthworm obtained from Horticulture Department) of nearly equal size (8 ± 1 cm) were selected for present study due to its anatomical and physiological resemblance with the intestinal roundworm parasites of human beings^{8,9}.

Each group was treated with one of the followings: vehicle (3 % Tween 80 in normal saline), piperazine citrate (15 mg/mL), albendazole (10 mg/mL) and PECN, BCN, CCN (5, 10 mg/mL) and EACN, PEJM (5, 10, 25 mg/mL) in normal saline containing 3 % Tween 80. Observations were made for the time taken to paralyse and/or death of individual worm up to 4 h of test period. Paralysis was said to occur when the worms did not revive even in normal saline. Death was concluded when the worm lost their motility followed with fading away of their body colour¹⁰.

RESULTS AND DISCUSSION

The anthelmintic activities^{11,12} of the title compound on *P. posthuma* are exhibited in Table-1. The perusal of the data reveals that PECN, BCN, CCN and EACN at the dose level of 5 and 10 mg/mL showed significant anthelmintic activities compared to reference standards albendazole and piperazine citrate. However, PEJM showed the effect at 25 mg/mL concentrations that is comparable with the reference standards.

TABLE- 1
ANTHELMINTIC ACTIVITIES OF ROOTS OF *C. nucifera* AND
AERIAL PARTS OF *J. multiflorum*

Compound	Concentration (mg/mL)	Time (min)	
		For paralysis	For death
Control (3 % Tween 80 in normal saline)	–	–	–
Albendazole	10	34.62 ± 0.61	63.78 ± 0.65
Piperazine citrate	15	18.57 ± 0.32	79.20 ± 1.00
PECN	5	11.30 ± 0.25	16.85 ± 0.50
	10	9.06 ± 0.45	14.10 ± 0.39
BCN	5	7.53 ± 0.44	22.69 ± 0.71
	10	5.03 ± 0.32	14.0 ± 0.30
CCN	5	6.16 ± 0.34	9.90 ± 0.40
	10	3.70 ± 0.25	7.05 ± 0.35
EACN	5	34.00 ± 0.55	56.10 ± 1.10
	10	20.20 ± 0.42	39.45 ± 0.80
PEJM	5	100.25 ± 1.40	130.55 ± 1.70
	10	75.99 ± 0.99	104.30 ± 1.39
	25	34.28 ± 0.50	63.00 ± 0.58

Results are expressed as mean ± SEM from six observations. PECN = petroleum ether extract of *C. nucifera*, BCN = benzene extract of *C. nucifera*, CCN = chloroform extract of *C. nucifera*, EACN = ethyl acetate extract of *C. nucifera*, PEJM = petroleum ether extract of *J. multiflorum*.

The present study therefore reveals that CCN, BCN and PECN are more potent than EACN and PEJM, even though all the five extracts were endowed with anthelmintic properties. The activity reveals concentration dependant nature of the different extracts. Potency of the extracts was found to be inversely proportional to the time taken for paralysis/death of the worms.

The above findings justify the anthelmintic properties of the above plants, which augment their use by the tribes of the particular areas. Further studies regarding the isolation and characterization of the active principle(s) responsible for anthelmintic activities and their mode of action are currently under progress.

ACKNOWLEDGEMENTS

The authors are thankful to Principal and Management of Seemanta Institute of Pharmaceutical Sciences, Jharpokharia, Mayurbhanj, Orissa for providing necessary facilities to carry out the present research work.

REFERENCES

1. K.R. Kartikar and B.D. Basu, *Indian Medicinal Plants*, Sri Satguru Publication, New Delhi, Vol. 10, p. 3557 (2000).
2. K.M. Nadkarni and A.K. Nadkarni, *Indian Meteria Medica*, Popular Prakashan, New Delhi, Vol. 1, p. 363 (2000).
3. R.N. Chopra, S.L. Nayer and I.C. Chopra, *Glossary of Indian Madicinal Plants*, CSIR, New Delhi, edn. 3, p. 73 (1992).
4. Anonymous, *The Wealth of India: A Dictionary of Indian Raw Materials and Industrial Product*, Publication and Information Directorate, CSIR, New Delhi, Vol. 5, p. 284 (1997).
5. A.V. Sala, *Indian Medicinal Plants*, Orient Longman Pvt. Ltd., Chennai, Vol. 3, p. 254 (2002).
6. M. Abraham, N. Sarada Devi and R. Sheela, *Indian J. Med. Res.*, **69**, 88 (1979).
7. R. Kalesaraj and A. Kurup, *Indian J. Pharm.*, **74**, 64 (1962).
8. G.W. Thorn, R.D. Adams, E. Braunwald, K.J. Isselleacher and R.G. Petersdrof, *Harrison's Principle of Internal Medicine*, McGraw Hill Co., New York, p. 1088 (1977).
9. Z. Vigar, *Atlas of Medicinal Parasitology*, P.G. Publishing House, Singapore, p. 216 (1984).
10. V.M. Shivkar and V.L. Kumar, *Pharm. Biol.*, **41**, 263 (2003).
11. G.K. Dash, B. Mishra, A. Panda, P.C. Patro and G. Ganapathy, *Ind. J. Nat. Prod.*, **19**, 24 (2003).
12. I.J. Kuppasta and V. Nayak, *Ind. J. Nat. Prod.*, **19**, 27 (2003).

(Received: 5 August 2006; Accepted: 7 June 2007) AJC-5678

**POLYMER FOAM 2007
INTERNATIONAL BUSINESS CONFERENCE ON POLYMER
FOAM TECHNOLOGY AND MARKETS**

2—3 OCTOBER 2007

DOUBLETREE HOTEL, NEWARK, NJ, USA

Contact:

Mrs. Margit Korsak,
Conference Director,
Tel: +1 610 478 0800
Email: mk@researchami.com