

## PHYTO-PHARMACOLOGICAL REVIEW OF *ARGYREIA NERVOSA*

A. Krishnaveni\*<sup>1</sup> and T. Sant Rani<sup>2</sup>

<sup>1</sup>College of Pharmacy, Madurai Medical College, Madurai, Tamil Nadu, India

<sup>2</sup>Institute of Pharmaceutical Technology, Sri Padmavathi Viswavidhyam, Tirupathi, Andhra Pradesh, India

\*A.Krishnaveni, M.Pharm,(Ph.D), College of Pharmacy, Madurai Medical College, Madurai 625020, Tamilnadu, India. Email: [akrishnaveni72@rediffmail.com](mailto:akrishnaveni72@rediffmail.com)

Article Received on: 29/12/10 Revised on: 23/01/11 Approved for publication: 12/02/11

### ABSTRACT

Herbal medicines are the significant and reliable sources for treating various diseases. *Argyrea nervosa* is traditionally used in wound healing, syphilis, diuretics, rheumatic affections, leucorrhoea, cerebral disorders, ulcers, as anti-tumour and to prevent contraception. Phytoconstituents such as flavanoids, steroids, ergoline alkaloids and triterpenoids were identified. Pharmacological studies proved its anticonvulsant, immunomodulatory, hypotensive, anti-inflammatory and nootropic effect. The present form of article highlights the phytochemical and pharmacological studies including traditional practice of *Argyrea nervosa* have been carried out so far.

**KEY WORDS:** *Argyrea nervosa*, ergoline alkaloids, Convolvulaceae

### INTRODUCTION

Complementary system of medicine includes namely Ayurveda, Siddha, Unani, Kaempo and Chinese medicine have gained its popularity in recent years<sup>1</sup>. Herb is said to be biosynthetic lab for the active principles like alkaloids, glycosides, resins, oleo-resins, tannins, phenylpropanoids etc, which exhibits definite pharmacological and therapeutic effect based on the dose<sup>2</sup>.

*Argyrea nervosa* belongs to Convolvulaceae commonly known as Elephant creeper. It is a climbing shrub with hard woody stalk bearing heart shaped leaves of 9-12 cm long and 8-10 cm breadth often cultivated throughout India. It is grown at an elevation of 900m. Leaves are larger, acute apex, cordate base, glabrous above and tomentose beneath. Flowers are large, purple silky pubescent with peduncled infundibula. Petals are purple, silky pubescent outside and wooly glabrous inside. Calyxes are white, tomentose outside with glabrous ovary. Fruits are globose and indehiscent berry<sup>3</sup>. The present article review the traditional uses, phytochemical studies and modern pharmacological studies evaluated so far.

### TRADITIONAL USES

*Argyrea nervosa* has been used widely by the tribals of Rajasthan to prevent conception while in Assam and Bihar leaves are used as vegetable. In Uttar Pradesh folklore practice the young leaves are used for healing the wounds<sup>4</sup>. Young leaves are used to treat wounds and skin infections. According to Yuani medicine, roots are used in rheumatic affections, alternative tonic to cerebral disorders and as diuretic<sup>5</sup>. Roots preparations are used to treat syphilis, synovitis, rheumatism, obesity, wound, ulcers, skin infections. Roots are used as an appetitizer, aphrodisiac, brain tonic, cardiotonic, expectorant, and in anti-inflammatory. In Hindu medicine, root is used externally for to reduce obesity<sup>6,7</sup>.

### PHYTOCHEMISTRY

Among the phytoconstituents ergoline alkaloids are the major constituents in *Argyrea nervosa*. It is one of the essential phytoprinciple of *Argyrea nervosa* from the historical and quantitative point of view. A group of nineteen indole alkaloids were identified and isolated by thin layer chromatography and paper chromatography. Among those constituents ergoline alkaloids, lysergic acid and isolysergic acid were analysed by thin layer chromatography, m.p, ultraviolet and infrared spectral analysis. Seeds are found to

possess hypotensive and spasmolytic activity which were due to the mixture of ergot alkaloids, isolated and analysed by ultraviolet. Due to instability only one constituent was identified as ergometrine. Other constituents such as caffeic acid and ethyl caffeate were identified<sup>8,9</sup>. Apart from ergoline alkaloids, N-formyllooline alkaloids, flavonoidal sulphates steroids and triterpenoids were isolated from other parts of *Argyrea nervosa*<sup>10,11</sup>. Para-hydroxy-cinnmate, scopelitin and argyroside<sup>12,13</sup>.

## **PHARMACOLOGICAL ACTIVITIES**

### **Antimicrobial activity**

According to Mishra et al., isolated oil from the seed of *Argyrea nervosa* and evaluated the antibacterial effect<sup>14</sup>.

### **Antifungal and phytotoxic activity**

A study was undertaken by Shukla et al., isolated the constituents p-hydroxy cinnmate and scopelitin from *Argyrea nervosa*, gallic acid from *Oenothera biennis*. The isolated phytochemicals were evaluated for antifungal activity and phytotoxicity. Para-hydroxy-cinnmate and scopelitin showed potent activity against the fungi *Alternaria alternate* and showed significant moderate inhibition of root growth of germinated seeds of wheat at 250ppm levels whereas 1000 ppm completely inhibited the root growth<sup>12</sup>.

### **Immunomodulatory effect**

A study was undertaken by Gokhale et al., proved that ethanolic extract of *Argyrea nervosa* root produced the immunomodulator effect on cellular and humoral immunity when administered at the dose of 50,100 and 200 mg/kg orally to rats potentiated the delayed type of hypersensitivity reaction induced by sheep red blood cells (SRBC) and oxazolone. It enhanced the production of circulating antibody titre in mice in response to SRBC. The above dose failed to show any effect on macrophage phagocytosis. Chronic administration of *Argyrea nervosa* significantly ameliorated the total WBC and restored the myelosuppressive effects induced by cyclophosphamide<sup>15</sup>.

### **Anti-inflammatory effect**

A pharmacological study by Bacchav et al., proved the anti-inflammation effect of *Argyrea nervosa*. Ethanolic extracts of various doses 50,100,200, mg/kg p.o were administered to acute and chronic models of inflammation induced by carrageenan. *Argyrea nervosa* significantly reduced the paw edema and prevented the accumulation of inflammatory cells in carrageenan induced peritonitis. The above study supported and adds credibility to the traditional use of this plant in rheumatism<sup>16</sup>.

### **Hallucinogenic effect**

The plant contains ergoline alkaloids showed hallucinogenic effect and used for recreation purposes<sup>17</sup>.

### **Anti diarrhoeal activity**

An investigation was taken by Ran et al., established the antidiarrhoeal effect of *Argyrea nervosa*. Hydroalcoholic extracts of *Argyrea nervosa* was undertaken to evaluate the anti diarrhoeal activity against experimental models of diarrhoea in rats. Administration of 50,100 and 150 mg/kg orally showed dose dependent decrease in the intestinal propulsion from 61.54 %-41.36 % in comparison with control and standard Atropine used. The extract appears to be as inhibitory as atropine and reduced the intestinal propulsion to greater extent. The extract also showed dose dependent decrease in the number of faecal matter passed by rats. In addition the extract also exhibited concentration-dependent reduction in the intestinal fluid accumulation from 9.97-39.52 when compared to control group<sup>18</sup>.

### **Aphrodisiac property**

Subramoniam et al., evaluated this property by mounting behavior, mating performance of male mice and its fertility effect on litter size and sex ratio of pups were observed. Aqueous extract of roots, flowers and leaves were used for screening the aphrodisiac effect. Alcoholic extract and n-hexane extract of leaves were used for the screening.

### **Mounting behavior**

Repeated administration of suspension 1g/kg of fresh parts of the plant (leaf and fruit) exhibited excessive mounting behavior in comparison with control. Therefore leaves treated mice showed only marginal activity when compared to root and flower. Single dose of 200mg/kg of ethanol extract of root was found to be effective, about 400% increases in mounting behavior to control group.

### **Mating performance**

Daily administration of root and flower suspension for six days to male mice resulted in increase in mating behavior of mice. Leaf suspension caused only moderate effect when compared to root and flower.

### **Effect of herbal drug on litter size (sex ratio of pups)**

The pups of dams of control treated group when inseminated found that all females were pregnant. As a outcome of results male/female ratio of pups were significant increase in the herbal drug treated. Thus, the root and flower of *Argyreia nervosa* are safe and effective medicine for stimulating sexual activity with preferential influence on sex ratio. Therefore, the sexual behavior could be elevated testosterone levels and other drugs, such effect can be achieved by *Argyreia nervosa* without any known toxic symptoms which adds credibility to ethnomedical belief that increase the chances of male offspring<sup>19</sup>.

### **Nootropic effect**

Joshi et al., performed nootropic activity by using memory models. Aqueous extract of *Argyreia nervosa* prepared by simple maceration. Two different doses of 100mg/kg and 200mg/kg were administered in the form of suspension by using Tween 80. Nootropic effect was evaluated by

1. **Memory model** such as exteroceptive behaviour model and passive shock avoidance paradigm. A dose of 200mg/kg was administered to rats orally increased step down latency significantly as compared to control. It also reversed effects of diazepam, scopolamine induced amnesia, natural aging and decreased the transfer latency on elevated the plus maze model.

2. **Estimation of brain acetyl cholinesterase (ACh -E) activity.** The whole brain AChE activity was measured using Ellman method. A dose of 100 and 200mg/kg were administered to rats showed significant reduction in ACh -E activity in comparison with piracetam and control<sup>20</sup>.

### **Effect of central nervous system**

A pharmacological study by Galani et al., evaluated the hydroalcoholic extract of *Argyreia nervosa* when fractionated with n-hexane, chloroform, ethyl acetate and water were concentrated under vacuum. The above derived fractions were evaluated for neuropharmacological activity including spontaneous motor activity and pentobarbitone induced hypnosis at different dose such as 100,200 and 500 mg/kg to various groups along with chlorpromazine (2mg/kg) and control. All the extracts produced significant and dose dependent reduction in spontaneous motor activity and potentiation of pentobarbital sodium induced sleep duration with hexane, chloroform, ethyl acetate and water fraction. Reduced onset of sleep and prolonged duration of sleep was observed in positive control. The results of above activity confirmed the ability of *Argyreia nervosa* in potentiating the pentobarbitone induced sleep and decrease in spontaneous locomotor activity highly recommends the effect of *Argyreia nervosa* in rheumatism<sup>21</sup>.

### **Anticonvulsant activity**

A pharmacological study by Vyawahare et al., evaluated the anticonvulsant effect of *Argyreia nervosa* roots. Different doses of 100,200 and 400 mg/kg p.o were administered to rats. Convulsions were induced by pentylenetetrazole (PTZ) and maximal electric shock (MES) method. Pretreatment of *Argyreia nervosa* caused significant delay in the onset of convulsion as well as lethality. At the dose of 100 mg no effect was observed whereas the other two dose produced significant prolonged onset of lethality after pentylene tetrazole administration and exhibited protection of 16.66 % and 33.33% in maximal electric shock. Pretreatment of *Argyreia nervosa* significantly reduced the hind limb extension<sup>22</sup>.

### **CONCLUSION**

The plant *Argyreia nervosa* is exclusively identified. The leaves are used internally and externally to treat wounds, skin infections by the local traditional practitioners for longer time. The research study may further be continued in identifying the phytoconstituents, toxicity and other useful pharmacological studies to explore its utilization.

**REFERENCES**

1. Eisenberg DM, Kessler RC, Foster CNorlock, CECalkins, DRD Delbanco TL. Unconventional medicine in the United States- Prevalence, costs and Pattern of use NEJM 1993; 328: 246-252
2. Montavale New Jersey, PDR for herbal medicines 1998; 1:1177-1178.
3. Varier's PS Indian Medicinal plants Editor's Warriar P.K.Nambiar, V.P.K and Ramankutty C.OrientLongman 1996; I Edtion:Volume 1: 191-195.
4. Anonymous, Wealth of India,CSIR, Govt of India, New Delhi 86-87, .& 1985 418
5. Nandkarni KM. Indian Materia Medica,Popular PrakashanPvt Ltd, Bombay, 1995; Vol.I: 136-137.
6. Kirthkar KR and Basu.BD Indian Medicinal Plants, International Book Distributors Dehradun,1993; volume I : 2<sup>nd</sup> edition:1707-1708.
7. Singh MP and Panda. H.Medicinal Herbs with Their Formulations Volume I, Daya Publishing House Delhi 2005 115-116
8. Miller MD. Isolation and Identification of Lysergic acid and isolysergic acid as the Principle ergoline alakoids in *Argyreia nervosa*, a tropical wood rose.Association of analytiacal Chemist, 1970; Jan: Volume 53: issue 3: 123-127.
9. Chao JM and Mardersian AH. Ergoline alkaloidal constituents of Hawaiian Baby Wood Rose. *Argyreia nervosa*.Bojer. Journal of Pharmaceutical Sciences, 1973; Issue April:Volume 62:588-591.
10. Agarwal SK, Rastogi RP. Ergometrine andother constituents of *Argyeia speciosa*. Indian Journal of Pharmacolgy, 1974 ;Volume 36: Sept October:118-119.
11. Petra Mann, Britta Tofern, Macki Kaloga and Eckart Eich. Flavonoid sulfates from the Convolvulaceae: Phytochemistry, 1999; January: Volume 50: Issue 2:26: 267-271.
12. YNShukla, Anil Srivastav, Sunil Kumar and Sushil Kumar. Phytotoxic activity and antimicrobial constituents of *Argyreia speciosa* and *Oenthera biennis*. Journal of Ethnopharmacolgy 1999; 67 (2): 41-245.
13. Rahman A, Ali M, Khan NZ. Argyroside from *Argyeia nervosa* seeds. Pharmazie 2003 Jan; 58 (1): 60-62.
14. Mishra SH and Chaturvedi.SC.Antibacterial and antifungal of the oil and unsaponifiable matter of *Argyreia nervosa*. Indian drugs Pharmaceutical Industry, 1978; 13 (5): 29-31.
15. Gokhlae AB, Damre AS and Saraf HN. Investigations into the immunomodulatory activity of *Argyreia nervosa*. Journal of Ethnopharmacolgy , 2003; Jan 84 1: 109-114
16. Shaw Cross WE. Recreational use of ergoline alkaloids from *Argyreia nervosa*. Journal Psychoactive drugs. 1983; Oct-Dec 15(4): 251-259.
17. ChV Rao, SKOjha, G Dreddy, AKS Rawat, GMM Rao and P Pushpangadan. Antidiarrhoeal Activity of *Argyreia speciosa* Flower: an Ethnopharmacological Study. Acta Pharmaceutia Turica, 2004; 46: 149-159.
18. Subramoniam AV, Madhavachandran K, Ravi and VS Anuja. Aphrodisiac property of Elephant creeper, *Argyeia nervosa* .Journal of Endocrinology Reproduction, 2007; 2: 282-85.
19. Joshi H. Habbu PV ,Mahadenen KM, Naveet K, Chauhan J, Krupa M. Evaluation of nootropic effect of *Argyreia nervosa* in mice. Journal of Health Sciences.2007;53(4): 382-388.
20. VJ Galani and BG Patel. Central Nervous System Activity of *Argvreia speciosa* Roots in Mice.Research J.Pharm and Tech, 2009 April; 2(2): 331-334.
21. NS Vyawahare and SL Bodhankar. Anticonvulsant Activity of *Argyreia speciosa* in Mice. Indian Journal of Pharmaceutical Sciences, 2009; March –April: 131-133.
22. AS Bacchav, VS Gulache and CD Upasain. Analgesic and Anti inflammatory activity of *Argyreia nervosa* root. Indian Journal of Pharmacology, 2009; 41(4):158-161.