

ANTIMICROBIAL ACTIVITY OF NYCTANTHES ARBORTRISTIS

DIVYA PAIKARA^{a1}, SHEETAL SINGH^b AND BHAWANA PANDEY^c

^{abc}Department of Biotechnology and Microbiology, Bhilai Mahila Mahavidyalaya, Hospital Sector, Bhilai, Chhattisgarh, India

ABSTRACT

Nyctanthes arbor-tristis is commonly known as Night-flowering Jasmine, Coral Jasmine and Parijat. In the present study, an attempt was made to investigate the anti-bacterial activity of *Nyctanthes arbortristis*. The crude drug powder extracts of the leaves of the above plants were taken for the study. The antibacterial activity was performed by using gram positive and gram negative organism viz., *Staphylococcus aureus* and *Escherichia coli*. The antifungal activity was performed by using both *Aspergillus niger* and *Aspergillus flavus*. Antimicrobial activity was performed by agar well diffusion method. The different solvent extracts of test material showed marked antimicrobial activity against pathogenic microorganism. The results showed that the Chloroform extract of *Nyctanthes arbor-tristis* shows highest antimicrobial activity and petroleum ether shows no antimicrobial activity.

KEYWORDS: Antimicrobial activity, *Nyctanthes arbor-tristis*

Ayurveda system of medicine use plants to cure the ailments and diseases. Despite the availability of different approaches for the discovery of therapeutically, natural products still remain as one of the best reservoir of new structural types (Cowan 1999). Medicinal plants besides therapeutic agents are also a big source of information for a wide variety of chemical constituents which could be developed as drugs with precise selectivity (Vijyalakshmi R, Ravindran R., 2012). *Nyctanthes arbor-tristis* is small sacred ornamental tree known across the country for its fragment white flowers. Plant is commonly known as night jasmine or parijata (Evans WC). Different parts of *N. arbortristis* are known to possess various ailments by rural mainly tribal people of India (Orissa and Bihar) along with its use in Ayurveda, Sidha and Unani systems of medicines (A.K. Nadkarni, 1982). The plants used in the traditional system of medicine of India and china are now receiving much scientific attention. Plants produce certain chemicals which are normally toxic to bacteria (Singh B, Bhat 2003). Traditionally the powdered stem bark is given in rheumatic joint pain, in treatment of malaria (Gupta, 2005) and also used as an expectorant. The bark is used for the treatment of snakebite and bronchitis. Juice of the leaves is used as digestives, antidote to reptile venoms, mild bitter tonic, laxative, diaphoretic and diuretic. (Hukkeri, 2006). It has also been reported to possess hepatoprotective, anti-leishmanial, anti-viral and anti-fungal activities (Puri, 1994) and analgesic, antipyretic and ulcerogenic activities (Saxena, 1987). The plant also possess anti-allergic (Gupta et al., 1993), anti-malarial (Badam et al., 1988), antihelminthic (Lal, 1976), activities and

recently reported hepatoprotective (Hukkeri Kusum et al., 2006), anti-spermatogenic (Gupta, 2006) and antioxidant activities (Rathee, 2007). The aim of the present study was to evaluate the susceptibility of various disease causing bacterial and fungal strains to different Solvent extracts of leaves of *Nyctanthes arbor-tristis* for Antimicrobial activity.

MATERIALS & METHODS

Collection of the plant samples: Fresh plant parts were collected randomly from Durg district of Chhattisgarh. The plants were identified and studied according to their families Fresh plant materials were collected and washed under tap water, shade dried and then homogenized to fine powder and stored in airtight bottles.

Preparation of plant extract: Ten grams of air dried powder was taken in 100 ml of petroleum ether in a conical flask, plugged with cotton wool and then kept on a rotary shaker at 190-220 rpm for 24 hours. After 24h, the supernatant was discarded and petroleum ether was evaporated from the powder. This dry powder was then taken in 100 ml of solvent (methanol or acetone) in a conical flask, plugged with cotton wool and then kept on a rotary shaker at 190-220 rpm for 24 h. After 24 h, the extracts were centrifuged at 5000 g for 10 min, the supernatant was collected, solvents were evaporated and the dry extract was weighed and stored at 4C in airtight bottles. The extraction was done at least three times for each plant. The preliminary qualitative phytochemical analysis was carried out in crude dry powder of selected plants.

Soxhlet extraction method: Leaves of selected plants were collected locally. Leaves were washed; air dried under shade and powdered with the help of Grinder. Powdered leaves were weighed and packed in soxhlet. Solvent used for soxhletion was petroleum ether and ethanol. Extraction was continued at the temperature of 35°C till clear solvent was observed in thimble. Extract was concentrated in water bath at 40°C. Concentrated extract was concentrated at 40°C in hot air oven. Concentrated extract was packed in an air tight container.

Antimicrobial activity: The antimicrobial activities of different plants were evaluated by Agar well diffusion test technique.

Antimicrobial Screening Test microorganisms: Antimicrobial activity was evaluated against common pathogenic microorganisms, Gram positive bacteria- *Staphylococcus aureus*, Gram negative- *Escherichia coli* and fungal strains *Aspergillus niger* and *Aspergillus flavus*. The bacterial cultures were grown and maintained on Nutrient Broth medium at 37°C for 24h while the fungal culture were maintained on Potato Dextrose Agar slants and incubated at 27°C for 48h.

Antibacterial assay: Fresh microbial culture of 0.1ml was spread on nutrient agar plate with glass spreader. A well of 6 mm diameter was punched off into agar medium with sterile cork borer and filled with 50 µg of ethanol, acetone, chloroform and petroleum ether extracts by using micropipette in each well in aseptic condition. The petriplates were then kept in a refrigerator to allow pre-diffusion of extract for 30 minutes and further incubated in an incubator at 37 °C for 24 h. The

antibacterial screening was evaluated by measuring the zone of inhibition.

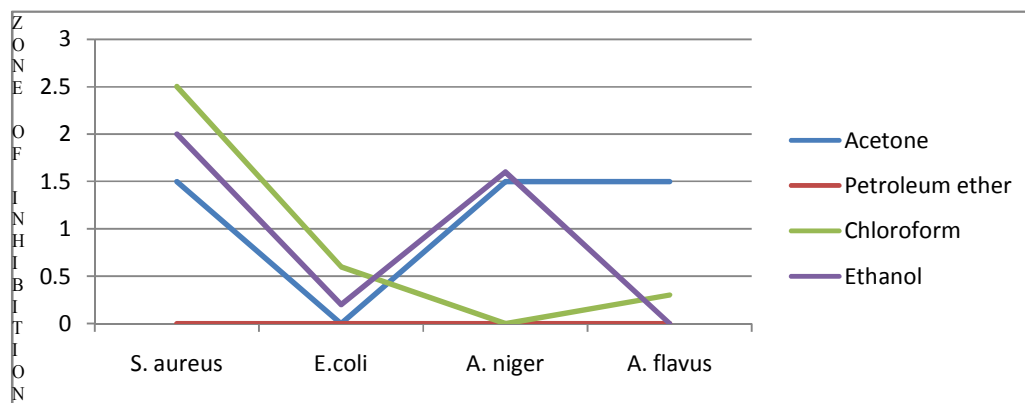
Antifungal assay: The antifungal activity of the leaf extracts was determined using agar well diffusion method. Small amount of diluted fungal suspension were poured over the media to spread uniformly on the surface. Later when the surface was little dried wells of 8mm were punched in the agar with stainless steel borer and filled with 300µl of plant extracts. Control wells containing neat solvents (negative control) were also run parallel in the same plate. The plates were incubated at 28°C for 72 hours and the antifungal activity was assessed by measuring the diameter of the zone of inhibition at the interval of every 24hrs.

RESULTS AND DISCUSSION

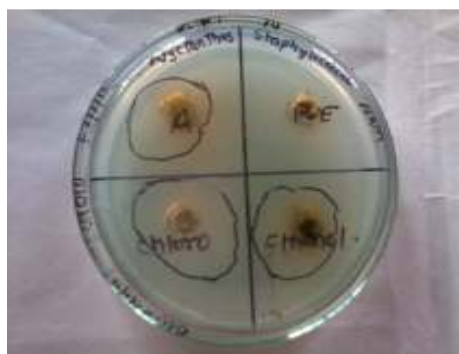
In the present investigation, antimicrobial efficacy of the crude extract of *Nyctanthes arbor-tristis* was quantitatively assessed on the basis of inhibition zone. The ethanol, chloroform, petroleum ether and acetone of *Nyctanthes arbor-tristis* exhibited varying degree of inhibitory effect against all tested pathogenic strains (Table). The most susceptible bacterium and fungi are *S. aureus* and *A.niger*, respectively. The inhibition zone were in the range of 2.5 to 1.6 mm for most of the tested strains. Chloroform extract of *Nyctathes arbor-tristis* shows highest antimicrobial activity and petroleum ether shows no antimicrobial activity shown in graph. According to the graph and table shows that the Chloroform leaves extract exhibited highest zone of inhibition against *S. aureus* and ethanol leaves extracted exhibited highest zone of inhibition *A. niger* (2.5mm and 1.6mm, respectively).

Table: Antimicrobial activity of leaf of *Nyctanthes arbor-tristis* (inhibition zone)

S.No.	Solvents	Zone of inhibition (mm)			
		Bacteria		Fungus	
		<i>S. aureus</i>	<i>E.coli</i>	<i>A.niger</i>	<i>A. flavus</i>
1	Acetone	1.5	-	1.5	1.5
2	Petroleum ether	-	-	-	-
3	Chloroform	2.5	0.6	-	0.3
4	Ethanol	2	0.2	1.6	-



Graph 1: Antimicrobial activity of leaf extract of *Nyctanthes arbor-tristis*



Nyctanthes arbor-tristis against *S. aureus*.



Nyctanthes arbor-tristis against *E. coli*



Nyctanthes arbor-tristis against *A. niger*.



Nyctanthes arbor-tristis against *A. flavus*

CONCLUSION

Many plants were being investigated for their antimicrobial activity by many scientists of different parts world. Leaves extract of *Nyctanthes arbor-tristis* with different solvents posses antimicrobial activity. From the above result it can be concluded that the leaf extract of selected plants have potential as antimicrobial compounds against microorganisms and they can be used in the treatment of infectious diseases caused by many microorganisms. By the above experimentation and result we can conclude that the selected leaf

extracts were showing highest antimicrobial activity with chloroform and ethanol extract.

REFERENCES

- Badam L, Deolankar RP, Rojatkark SR, Nagsampgi BA and. Wagh UV, 1988. In vitro antimalarial activity of medicinal plants of India. J. Med. Res., 87: 379-383.
- Cowan MM, 1999. Plant products as antimicrobial agents. Clin. Microbial. Rev., 12:564-582.
- Evans WC. "Trease & Evan's", Pharmacognosy. Harcourt Brace Asia, 48.

- Gupta P, Bajaj SK, Chandra K, Singh KL, Tandon JS, 2005 "Antiviral Profile of Nyctanthes arbor-tristis against encephalitis causing viruses". Indian Journal of Experimental Biology , 43:1156-1160.
- Gupta PP, Srimal KL, and Tondon JS, 1993. Anti-allergic activity of some traditional Indian medicinal plants. Int. J. Pharmacog., 31: 15-18.
- Hukkeri V, Akki K, Sureban RR, Gopalkrishnan B, Byahatti VV, Rajendra SV, 2006.
- Hepatoprotective activity of the leaves Nyctanthes arbor-tristis Linn". Indian Journal of Pharmaceutical Sciences 4:542-543.
- Khatune NA, Mosaddik MA and Haque ME, 2001. Antibacterial activity and cytotoxicity of Nyctanthes arbor-tristis flowers. Fitoterapia, 72: 412-414.
- Lal J, Chandra S, Ravi Prakashand V, and Sabir M, 1976. In vitro anthelmintic action of some indigenous medicinal plants on Ascaridia galli worms. Ind. J. Physiol. Pharmacol., 20: 64-68.
- Nadkarni AK, 1982, Indian Materia Medica, Vol.I, 3rd ed. (Popular Prakashan Pvt. Ltd.), 857-858.
- Rathee, JS, Hassarajani SA and Chattopadhyay S, 2007. Antioxidant activity of Nyctanthes arbor-trisr-tis leaf extract. Food Chem., 103: 1350-1357.
- Singh B, Bhat TK, Bhupender S, 2003. Potential Therapeutic Application of Some Antinutritional Plants Secondary Metabolite. Journal of Agriculture Food Chemical; 51:5579-5597
- Saxena RS, Gupta B, Saxena KK, Srivastava VK and Prasad DN, 1987. Analgesic, antipyretic and ulcerogenic activity of Nyctanthes arbor-tristis leaf extract J. Ethno pharmacol., 19; 193-200.
- Vijyalakshmi R, Ravindran R, 2012. Preliminary comparative phytochemical screening of root bextracts of *Diospyrus ferrea* (Wild.) Bakh and *Arva lanata* (L.) Juss. Ex Schultes. Asian J Plant Sci Res 2:581-587