Antibacterial activity of *Ixora coccinea* against selected bacterial pathogens isolated from clinical samples

SOMA BANERJEE¹, NAMITA BHOWMICK², TAPASHI GHOSH³, RESMA SAHA¹, FATIMA NASRIN HAQUE¹, ABHISIKTA DAS¹ AND RIA BANERJEE¹

- ¹Department of Biotechnology, Heritage Institute of Technology, Chowbaga Road, Anandapur, P.O. East Kolkata Township, Kolkata 700107
- ²Department of Bioscience, Sardar Patel University, Vallabh Vidyanagar, Anand, Gujarat 388120.
- ³Department of Microbiology, R.G. Kar Medical College & Hospital, 1 Khudiram Bose Sarani, Kolkata 700004, West Bengal

Received: 06.09.2013

RMS Accepted: 25.03.2014

Published: 28.04.2014

The present work aimed at exploring the antimicrobial activities of flower and leaf extracts of *Ixora coccinea*, belonging to the Rubiaceae family against clinical samples from tertiary health care hospital patients. In this study aqueous, methanolic and ethanolic extracts of both flower and leaf were screened. Antibacterial activity was screened against these samples and also against reference strains by means of agar-well diffusion method. The ethanolic and methanolic extracts of both flowers and leaves of *I. coccinea* showed promising antibacterial activity against both the clinical and reference *Staphylococcus aureus* and *Pseudomonas aeruginosa* strains. The phytochemical screening of the organic plant extracts revealed the presence of alkaloids, steroids, flavonoids and tannins.

Key words: Agar-well diffusion method, anti-bacterial activity, bacterial samples, plant extracts

INTRODUCTION

Drug resistant bacterial strains usually develop due to the over use of a number of antibacterial drugs as a result it arises trouble in controlling the growth of infectious disease, even over usage produces side effects (Tomasz and Tomin, 1986). From ancient times in ayurvedic medicinal plants are used for curative purpose for various ailments as a large number of organic compounds secondary metabolites are present in plants. These act as potent bioactive compounds, for various purposes like chemotherapeutic, bactericidal, and bacteriostatic agents (Bohra and Purohit, 1998). The genus Ixora contains more than 400 species and is one of the low growing evergreen perennial shrubs through Southeast Asia. It is a popular flowering shrub belongs to the Rubiaceae family. Pharmacologically the leaves are reported for their antimicrobial, antidiarrhoeal and antinociceptive activities (Annapurna et al., 2003; Agashikar et al., 2010; Ratnasooriya et al., 2005). Flowers possess antioxidant, antiinflammatory, wound healing activities (Saha et al., 2008; Udupa et al. 1999). The present study has been carried out to evaluate the antibacterial activity and phytochemical analysis of various organic extracts of both the flowers and leaves of *Ixora coccinea* against bacterial pathogens isolated from clinical samples.

MATERIALS AND METHODS

Preparation of plant extracts

The leaves and flowers were shade dried. 20 g of both the samples of leaves and flowers were crushed using a mortar pestle then 70% ethanolic, 70% methanolic and aqueous extract of 200 ml each were prepared and kept at 4°C for 48 h. These extracts were then filtered and were kept in

-18ºC for further use.

Microbial strains and culture media

In the current study, sixty two clinical isolates were used for the assay of the antimicrobial property from *Ixora coccinea*. These isolates were obtained from clinical samples from a tertiary health care hospital, Kolkata. Out of these different clinical samples (wound and pus) 19 samples were found to be positive for *Staphylococcus aureus* and 43 samples for *Pseudomonas aeruginosa*. Apart from that two reference strains of *Staphylococcus aureus* 25923 ATCC and *Pseudomonas aeruginosa* 27853 ATCC were taken. All the isolates were maintained on Nutrient Agar (HiMedia) slants at 5°C.

Antibacterial activity tests

Antibacterial activity tests were performed by standardized agar-well diffusion method (Bazerque et al.1990). For this technique, bacterial pathogens prepared in saline water (0.85% NaCl) and adjusted to a turbidity of 0.5 McFarland standards (108 CFU/ ml) were spread on the solid plates with a sterile swab. Wells (6 mm diam.) were punched and the wells were filled with ethanolic, methanolic and aqueous extracts of both leaves and flowers of Ixora coccinea were used as positive control and the solvents used for preparing extracts were used as negative control following standard protocol. Standard disc of ciprofloxacin (5 mcg/disc), ampicillin/sulbactam (10/10 mcg/disc), levofloxacin (5mcg/disc), chloramphenicol (30 mcg/disc), nitrofurantoin (300 mcg/disc), nalidixic acid (30 mcg/ disc), ceftazidime (30 mcg/disc), piperacillin/ tazobactam (100/10 mcg/disc), co-trimoxazole (25 mcg/disc) and oxacillin (1 mcg/disc) (HiMedia) were used against all isolates. Plates were incubated at 37°C for 24 h. Antibacterial activities were evaluated by measuring the diameter of zone of inhibition.

RESULTS AND DISCUSSION

In this study two reference strains Staphylococcus aureus 25923 ATCC and Pseudomonas aeruginosa 27853 ATCC were taken in parallel to 62 clinically isolated samples. In the study of monitoring antimicrobial activities 10 types of antibiotic discs were tested against individual clinical isolates, i.e., ciprofloxacin, levofloxacin, chloramphenicol,

ampicillin/sulbactam, nalidixic acid, ceftazidime, nitrofurantoin, piperacillin/tazobactam, cotrimoxazole and oxacillin along with aqueous, 70% ethanolic and 70% methanolic exrtracts of both the leaves and flowers.

Table 1 : Sensitivity of reference strains with standard antibiotics discs and plant extracts in (mm)

and extracts (mg)	Pseudomona saeruginosa ATCC 27853	Staphylococcus aureus ATCC 25923
Leaf ethanolic extract (10)	19	21 ·
Leaf methanolic extract (10)	17	18
Leaf aqueous extract (15)	9	8
Flower ethanolic extract (10)	23	22
Flower methanolic extract (10)	17	15
Flower aqueous extract (15)	9	9
Ciprofloxacin (5)	30	33
Levofloxacin (5)	24	33
Chloramphenicol (30)	18	30
Ampicillin/Sulbactam (10/10)	13	31
Nalidixic Acid (30)	24	15
Ceftazidime (30)	27	21
Nitrofurantoin (300)	20	21
Piperacillin/Tazobactam (100/1	0) 28	37
Co-Trimoxazole (25)	38	35
Oxacillin (1)	24	25

All the reference strains showed sensitivity to all the ten antibiotics and in all the plant extracts (Table 1). Among the 43 Pseudomonas aeruginosa clinical isolates they were found to be maximum resistant to 69.8%, 67.4%, and 65.1% with the following antibiotics ampicillin/sulbactam, oxacillin, and nitrofurantoin respectively; in other words 30, 29 and 28 strains were resistance to ampicillin/ sulbactam, oxacillin and nitrofurantoin respectively. Nalidixic acid, chloramphenicol, ceftazidime showed moderate resistance (Table 3). In case of plant extracts, ethanolic extracts of both leaves and flowers showed antibacterial activity of 65.1% and 58% respectively followed by 53.5% and 48.8% of methanolic extracts of both leaves and flowers (Table 2). The nineteen Staphylococcus aureus

Table 2: Resistant patterns of clinical isolates with plant extracts

Leaf and flower extracts		Sensitivity pattern of 62 isolates			
(mg)	Pseudomonas aeruginosa		Staphylococcus aureus		
	No of sensitive isolates	No of resistant isolates	No of sensitive isolates	No of resistant isolates	
Leaf ethanolic extract (10)	28	15	18	1	
Leaf methanolic extract (10)	23	20	17	2	
Leaf aqueous extract (15)	3	40	2	17	
Flower ethanolic extract (10)	25	18	16	3	
Flower methanolic extract (10)	21	22	14	5	
Flower aqueous extract (15)	3	40	2	17	

Table 3: Resistant patterns of clinical isolates with standard antibiotics

	Antibiotic sensitivity pattern of 62 isolates				
Antibiotics (mcg/disc)	Pseudomona	as aeruginosa	Staphylococcus aureus		
, -	Number of sensitive isolates	Number of resistant isolates	Number of sensitive isolates	Number of resistant isolates	
Oxacillin (1)	14	29	10	9	
Ampicillin/Sulbactam (10/10)	13	30	16	3	
Nitrofurantoin (300)	15	28	19	0	
Co-Trimoxazole (25)	21	22	17	2	
Nalidixic Acid (30)	28	15	16	3	
Chloramphenicol (30)	26	17	19	0	
Ceftazidime (30)	29	14	17	2	
Piperacillin/Tazobactam (100/10)	35	8	17	2	
Ciprofloxacin (5)	39	4	19	0	
Levofloxacin (5)	- 39	4	19	0	

Table 4: Phytochemical properties of Ixora coccinea leaf and flower extracts

Test	Ixora coccinea leaf extracts			Ixora coccinea flower extracts		
	n-butanol	ethanol	Acidic methanol	n-butanol	ethanol	Acidic methano
Alkaloids	+	+	+	+	+	+
Tannins	+	+	+	+	+	+
Steroids	+	+	+	+	+	+
Flavonoids	+	+	T-1	# 0	+	+

^{+ =} presence, - = absence

clinical isolates showed moderate resistance to oxacilin that is 9 strains (Table 3). In case of plant extracts, both the leaves and flowers extracts showed very good antibacterial activity, the ethanolic and methanolic extracts of leaves showed

94.7% and 89.5% respectively followed by 84.2% and 73.7% of ethanolic and methanolic extracts flowers (Table 2). Phytochemical analysis revealed the presence of flavonoids, tannins, steroids and alkaloids in all the organic extracts of leaves and

flowers of I. coccinea (Table 4).

The current study states that both ethanolic and methanolic extracts of leaves and flowers show good antibacterial activity against clinical sample than aqueous extracts. Against Staphylococcus aureus both ethanolic and methanolic extracts of leaves and flowers of Ixora coccinea showed higher antibacterial activity and for Pseudomonas aeruginosa it is moderate. So from current study, these plant parts show more antistaphylococcal activity than antipseudomonal activity. The result shows that this plant shows more antibacterial activity against S. aureus which is gram positive bacteria than P.aureginosa which is gram negative bacteria (Table 2). There are numerous workers who found related finding in different medicinal plants like in Ocimum sanctum, Cinnamomum zeylanicum (Joshi et al., 2009) and in leaf extracts of neem (Maragathavalli et al., 2012). Thus, the study suggests this plant can be used in the treatment of wound and pus infections caused by resistant bacteria. Additionally, the potential of these plants must be explored to a greater extent, in turn to develop an alternate therapy for the treatment of infections caused by antibiotic-resistant bacteria.

ACKNOWLEDGEMENT

The authors are grateful to the authorities of the Department of Biotechnology, Heritage Institute of

Technology, Kolkata for providing the necessary facilities for the present study.

REFERENCES

- Agashikar, N. V. Prabhu, B. and Yasmeen, M. 2010, Evaluation of the antidiarrhoeal activity of the leaves of *Ixora coccinea* Linn. In rats. *Jour. Clin. Diag. Res.* **4**: 3298-3303.
- Annapurna J, Amarnath P.V.S., Amar Kumar D., Ramakrishna S.V., Raghavan K.V. 2003, Antimicrobial activity of *Ixora coccinea* leaves. *Fitoterapia*. 74: 291-93.
- Bazerque, P. Paul, M and Perez, C. 1990, Antibiotic assay by agar-well diffusion method. *Acta Biol Medicin Experim*; **15**: 113-115
- Bohra, A. and Purohit, P. 1998, Effect of some plants extracts on conidial germination of some important phytopathogenic fungi. *Geobios New Report.* 17: 183-184.
- Joshi, B. Lekhak, S. and Sharma, A. 2009, Antibacterial Property of Different Medicinal Plants: Ocimum sanctum, Cinnamomum zeylanicum, Xanthoxylum armatum and Origanum majoran. Kat. Uni. Jour. of Sc. Eng. Tech. 5: 143- 150.
- Maragathavalli, S., Brindha, S., Kaviyarasi, N.S., B. Annadurai, B. and Gangwar, S.K. 2012, Antibacterial Activity in Leaf Extract of Neem (Azadirachta indica Linn.). I.J.S.N. 3: 110-113.
- Ratnasooriya W.D., Deraniyagala S.A., Galhena G., Liyanage S.S.P., Bathige S.D.N.K. and Jayakody J.R.A.C. 2005 Antiinflammatory activity of the aqueous leaf extract of *Ixora coccinea*. *Pharmaceutical Biol.* 43: 147 - 152.
- Saha, M. R. Alam A., Akter. R., Jahangir, R.2008. In vitro free radical scavenging activity of Ixora coccinea L. Bang. J Pharm, 3: 90-96
- Tomasz, A. and Tomin, E. 1986. Beta-lactam specific resistant mutants of *Staphylococcus aureus*. Antimicrobiol. Agents Chemother. 30: 377-383.
- Udupa S.L. Nayak B.S, Udupa A.L. 1999, Effect of Ixora coccinea flowers on dead space wound healing in rats. Fitoterapia. 70: 233-236.