

Research Journal of Pharmaceutical, Biological and Chemical Sciences

Evaluation of antimicrobial activity of *Solanum trilobatum* linn. Roots

V Mihira^{*}, K V Ramana, S Ramakrishna, B Praveen Kumar

*Pharmacology Division, A.S.N.Pharmacy College, Burrupalem Road, Tenali-522201, Guntur Dist., A.P.

ABSTRACT

The *Solanum trilobatum* Linn. (Family: Solanaceae) root powder was extracted with chloroform, methanol and screened for its anti-microbial activity against various gram (+) ve, gram (-) ve bacteria and fungal organisms using cup plate agar diffusion method. The results revealed that the chloroform and methanolic extracts shown significant anti-microbial activity at concentrations of 100 mg/ml and 200 mg/ml against tested organisms, particularly these were effective against gram (+)ve bacteria *Bacillus subtilis*, *Staphylococcus aureus*, gram (-)ve bacteria *Escherichia coli* and fungi *Candida albicans*, *Saccharomyces cerevisicea*.

Keywords: *Solanum trilobatum* Linn., anti-bacterial activity, anti-fungal activity.

***Corresponding author**

INTRODUCTION

The plant *Solanum trilobatum* Linn. (Family: Solanaceae) grows as a climbing undershrub and is widely distributed throughout the state of Andhra Pradesh and Tamilnadu. This plant is well known in Ayurved and Siddha systems. In Sanskrit it is known as 'Alarka', in Telugu 'Alarkapatramu', in Tamil 'Tuduvalai' and in Malayalam 'Tutuvalam'. The roots, berries and flowers are used for cough [1]. The decoction of entire plant is used to treat acute and chronic bronchitis [2]. The review of literature revealed that some chemical constituents like solasodine and β -solamarine have been isolated from whole plant [3]. The *Solanum trilobatum* Linn. posses antioxidant, hepatoprotective, anti-inflammatory, analgesic, antidiabetic and antimicrobial activities [4, 5,6,7,8,9,10]. In this present study the antimicrobial activity of the *Solanum trilobatum* Linn. roots chloroform and methanolic extracts were have been investigated.

MATERIALS AND METHODS

Preparation of extracts

The roots of *Solanum trilobatum* Linn. were collected from the coastal area of Andhra Pradesh. They were dried, powdered and extracted in soxhlet with chloroform and methanol (2 liters each) and concentrated to a small volume. The concentrated extracts were tested for anti-microbial activity.

Procedure

Evaluation of anti-microbial activity:

The anti-bacterial activity of extracts was screened against *Bacillus subtilis*, *B.pumilis*, *B.cereus*, *Staphylococcus aureus* gram (+)ve and *Escherichia coli*, *Pseudomonas aurgenosa*, *Proteus vulgaris*, *Serratia marceseanis* gram (-)ve using agar diffusion cup-plate method [11]. The extracts were tested at 100 mg/ml and 200 mg/ml levels. The results were compared with Neomycin sulphate (10 μ g/ml in DMSO). The results are recorded in Table-1.

Table no: 1 Evaluation of Anti-Bacterial Activity

| BACTERIA | Chloroform Extract (100 mg/ml) | Methanolic Extract (100 mg/ml) | Chloroform Extract (200 mg/ml) | Methanolic Extract (200 mg/ml) | Solvent Control DMSO | Standard (10 μ g/ml) |
|------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|----------------------|--------------------------|
| <i>Bacillus subtilis</i> | 18 | 11 | 19 | 15 | 4 | 19 |
| <i>Bacillus cereus</i> | 11 | 10 | 12 | 13 | 3 | 20 |
| <i>Bacillus pumilis</i> | 17 | 11 | 18 | 13 | 3 | 19 |
| <i>Staphylococcus aureus</i> | 12 | 11 | 14 | 12 | 2 | 20 |
| <i>Escherichia coli</i> | 19 | 12 | 20 | 14 | 3 | 18 |
| <i>Pseudomonas aurgenosa</i> | 10 | 10 | 12 | 10 | 2 | 20 |
| <i>Proteus vulgaris</i> | 11 | 10 | 12 | 11 | 3 | 20 |
| <i>Serratia marceseanis</i> | 10 | 10 | 11 | 11 | 3 | 20 |

Standard Drug: Neomycin Sulphate – 10 μ g/ml

The anti-fungal activity of extracts was screened against *Aspergillus niger*, *Candida albicans*, *Penicillium notatum*, *Saccharomycis cerevisicea* using agar diffusion cup-plate method. The extracts were tested at 100 mg/ml and 200 mg/ml levels. The results were compared with Nystatin (10 µg/ml in DMSO). The results are recorded in Table-2.

Table: 2 Evaluation of Anti-Fungal Activity

| FUNGI | Chloroform Extract (100 mg/ml) | Methanolic Extract (100 mg/ml) | Chloroform Extract (200 mg/ml) | Methanolic Extract (200 mg/ml) | Solvent Control DMSO | Standard (10 µg/ml) |
|----------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|----------------------|---------------------|
| <i>Aspergillus niger</i> | 10 | 11 | 12 | 13 | 3 | 20 |
| <i>Candida albicans</i> | 18 | 12 | 19 | 15 | 4 | 20 |
| <i>Penicillium notatum</i> | 12 | 11 | 13 | 13 | 3 | 18 |
| <i>Saccharomycis cerevisicea</i> | 17 | 15 | 18 | 16 | 2 | 18 |

Standard Drug: Nystatin – 10 µg/ml

RESULTS AND DISCUSSION

The tested bacteria and fungi have shown significant susceptibility to the chloroform and methanol extracts of *solanum trilobatum* Linn. roots. Three of the tested bacteria *Bacillus subtilis*, *Staphylococcus aureus* gram (+)ve, *Escherichia coli* gram (-)ve and two of the tested fungi *Candida albicans*, *Saccharomycis cerevisicea* were found to be more sensitive to the chloroform and methanol extracts of *solanum trilobatum* Linn. roots. The observed activities were further revealed that the amount of activity is increased with concentration of the extract and also the chloroform extract has shown more degree of anti-microbial activity than the methanolic extract.

REFERENCES

- [1] Anonymous, The Wealth of India-A Dictionary of Indian Raw Materials & Industrial Product, Vol-IX, CSIR, 1972: 395-396.
- [2] Nadkarni KM. Indian Material Medica, Popular Prakasan Pvt. Ltd.1976, 1153-1154.
- [3] Purushothaman KK, Balakrishana K, Sarada A, Bhema Rao R. Ind Drugs 1987; 24: 214-215.
- [4] Sini H & Devi KS. Pharm Biol 2004; 42: 462 – 466
- [5] Shahjahan M, Sabitha KE, Mallika Devi R, Shyamala CS. Ind J Med Res 2004; 123: 23-27.
- [6] Emmanuel S, Ignacimuthu S, Perumalsamy R, Amalraj T. Fitoterapia 2006; 77: 611–612.
- [7] Pandurangan A, Khosa RL, Hemalatha S. Oriental Pharmacy and Exp Medicine 2008; 8(4): 416-422.
- [8] Pandurangan A, Khosa, RL, Hemalatha S. Iranian J Pharmaceutical Research 2008; 7(3): 217-221.



- [9] Doss A, Palaniswamy M, Angayarkanni J, Dhanabalan R. African J Biotech 2009; 8(20): 5562-5564.
- [10] Swapna Latha P, Kannabiran K. African J Biotech 2006; 5(23): 2402-2404.
- [11] The Indian Pharmacopoeia, Vol-II, 3rd edition, Publication and Information Directorate, CSIR, New Delhi, 1984, 90.