

IN-VITRO EVALUATION OF ANTIMICROBIAL POTENTIAL OF AYURVEDIC POLY-HERBAL FORMULATION: SUDARSHAN CHURNA

**Baljinder Singh^{*1}, Dharmendra Kumar², Neerja Jindal³, Vikas Gupta⁴,
Parveen Bansal⁴**

¹Swami Vivekanand College of Pharmacy, Banur, India

²Laureate Institute of Pharmacy, Kangra, India

³Department of Microbiology, GGSMCH, Faridkot, India

⁴University Centre of Excellence in Research, BFUHS, Faridkot, India

Article Received on
10 January 2014

Revised on 25 January 2014,

Accepted on 22 February
2014

***Correspondence for
Author**

Dr. Baljinder Singh

Swami Vivekanand College
of Pharmacy, Banur, India.

ABSTRACT

Ayurvedic medicine plays a crucial role in healthcare and serves the health need of a vast majority of people in developing countries. *Sudarshan Churna* (SC) is very potent ayurvedic medicine; composed of 42 medicinal plants, which is used traditionally in treatment of malaria, viral fever, and bacterial infection. In this study the activity of this formulation was compared with the standard antibiotics like Amikacin and Norfloxacin. Ethanol, methanol and acetone extract of *Sudarshan Churna* demonstrated good antimicrobial activity and thus can form the basis for the development of a novel antibacterial formulation.

KEY WORDS: *Sudarshan Churna*, antimicrobial activity, ayurvedic medicine.

INTRODUCTION

In the last few decades, there has been an exponential growth in the field of ayurvedic medicine (Indian Traditional System of Medicine (1). Herbal medicines are being used increasingly as dietary supplements to fight or prevent common diseases (2). The search and use of drugs and dietary supplements of plant origin have accelerated in recent years. Ethnobotanists, pharmacologists, microbiologists and natural product chemists are combing the earth for phytochemicals which could be developed for treatment of infectious diseases.

Out of 25 to 50% of current pharmaceuticals derived from plants, very few are used as antimicrobials. Since time immemorial, the traditional physicians have been using plants for prevention or cure of infectious conditions. Western medicine is trying to duplicate their success (3). Today 80% of the world's population in African, Asian, Latin American and Middle Eastern countries is using plants as traditional health remedies due to minimal side effects (4-6). In present scenario, pharmaceutical companies are investing significant amount of time and money for development of therapeutics based upon natural products obtained from plants (7-8). It is well known that use of most of the modern antimicrobials is fraught with adverse effects. The problem gets complicated since these antimicrobials are used for an extended period of time. Therefore, there is need to explore antibacterial activity of certain herbs and to create evidences for their efficacy. In case of *Sudarshan Churna* the key ingredient is *Swertia chirata* which constitutes 50% of its total composition (9). Ayurvedic Churnas are solid dosage form of medicaments meant for internal use. The dose is 1-2 tea spoonful which may be increased or decreased according to age and severity of disease. Churnas can be administered with water, milk, fruit juices or any other suitable liquid depending on the nature of disease. They may be given by mixing with honey in equal quantity, with sugar twice the quantity and with the milk four times the quantity as that of the drug (10). The *Sudarshan Churna* was procured from local market of Faridkot and subjected to study the *in-vitro* antimicrobial activity to justify the traditional claims of the Polyherbal formulation.

The dose of *Sudarshan Churna* is 2-4 gm.

Therapeutic uses

The *Sudarshan Churna* is used to treat Yakrt (Liver), Pliha vrddhi (Splenomegaly), Jvara (Fever), Visama jvara (Intermittent fever), Jirna jvara (Chronic fever), and Gulma (Abdominal Lump) (11).

MATERIAL & METHODS

Drugs and chemicals

The present study was conducted in the Department of Microbiology, Government Medical College, Faridkot. *Sudarshan Churna* was procured from the local market of Faridkot.

Microbial strains

The various different strains of microorganisms were used and were purchased from M.T.C.C. Institute of Microbial Technology (IMTECH) Chandigarh in lyophilized form with

complete detail of growth media required for converting these to the active culture. These are as follows *Pseudomonas aeruginosa* (MTCC 424), *Staphylococcus aureus* (MTCC 3160), *Klebsiella pneumonia* (MTCC 3384), *Bacillus subtilis* (MTCC *121), *Escherichia coli* (MTCC 739).

Extraction of drug material

Method of Parekh *et al.* (2005) was used for the extraction of drug material (after some modifications). The aqueous extract was prepared by adding 20 g of herbal preparations in 200 ml distilled water, heated at 60° C for 2 h, filtered and the filtrate was evaporated on sand bath. The dry mass (3.6%w/w) was then stored at 4°C. The organic solvent extract was prepared by adding 20 g herbal preparation (powder) in 200 ml of organic solvent (acetone, ethanol and methanol) in screw-capped bottles and was put at 190-220 rpm on a rotary shaker. After 24 h of shaking, the extract was filtered, evaporated in vacuum and dried by rotary evaporator at 60°C (12-13). Dried extracts (2.9%, 3.1%, 3.7% w/w respectively) were stored in labeled sterile screw capped bottles at 4°C and later used for the *in vitro* study.

Formulation of extract

A known amount of powder was suspended in corresponding solvent to get desired concentration of suspension for the study of antimicrobial activity on the day of experimentation (14).

Antimicrobial activity

Antibacterial activity was determined by using Disc Diffusion Method. The impregnated filter paper discs were employed to determine the antibacterial activity of both aqueous and organic solvent extracts of herbal preparation (15). For antibacterial properties, 0.1 ml bacterial suspension of 10^5 CFU ml⁻¹ was swabbed on Nutrient Agar plate to form lawn culture. The aqueous, acetone, ethanol and methanol extracts were prepared in their respective solvents. The filter paper discs (6mm in diameter) were separately impregnated with different concentrations of extract and then placed on the agar plates which had previously been inoculated with the test microorganisms. Discs were soaked in various organic solvents, dried and were placed on lawns as negative control. After incubation of 24 h at 37° C, zone of inhibition of growth was measured in mm. The % inhibitory concentrations of the different extracts were measured and compared with the antibiotics (16-17) like Amikacin (30 µg) and Norfloxacin (10 µg) as standard.

RESULTS

From table it is evident that Ethanolic extracts of formulation showed more activity against *Pseudomonas aeruginosa* as compared to other extracts of the formulation, Acetone extract showed more significant activity against *Staphylococcus aureus* which is comparable with standard antibiotics, while ethanol extract also showed significant antimicrobial activity against *Staphylococcus aureus* which is comparable with norfloxacin (as shown in the table). *Klebsiella pneumoniae* showed sensitivity against ethanol and acetone extract which is comparable to norfloxacin while in case of *Bacillus subtilis* the methanol extract shows significant antimicrobial activity which is nearby the activity shown by standard antibiotics norfloxacin on the other hand *Escherichia coli* was found sensitive to methanol, ethanol and acetone extracts of formulation which is comparable with standard antibiotics.

ANTIMICROBIAL ACTIVITY

Table Antimicrobial activity of different extract of Sudarshan churna

Sr. no.	Microbial strains	Type of Extract	Zone of inhibition at different concentration (%) of extracts (mm)							
			100	80	60	40	20	C	Standard Antibiotics	
									Amikacin (30 µg)	Norfloxacin (10 µg)
1.	<i>Pseudomonas aeruginosa</i> (MTCC 424)	Methanol	15	13	12	8	-	7	33	34
		Ethanol	20	17	14	12	-	7		
		Acetone	17	13	12	7	-	7		
		Water	-	-	-	-	-	-		
2.	<i>Staphylococcus aureus</i> (MTCC 3160)	Methanol	18	16	10	8	-	6	23	17
		Ethanol	18	12	11	9	-	7		
		Acetone	21	16	11	9	-	7		
		Water	-	-	-	-	-	-		
3.	<i>Klebsiella pneumonia</i> (MTCC 3384)	Methanol	17	12	8	-	-	6	27	19
		Ethanol	20	13	12	9	-	7		
		Acetone	20	18	16	12	-	9		
		Water	-	-	-	-	-	-		
4.	<i>Bacillus subtilis</i> (MTCC *121)	Methanol	21	17	14	10	-	8	34	25
		Ethanol	18	13	11	8	-	-		
		Acetone	18	15	10	7	-	8		
		Water	-	-	-	-	-	-		
5.	<i>Escherichia coli</i> (MTCC 739)	Methanol	19	17	14	9	-	6	0	0
		Ethanol	21	20	17	13	8	8		
		Acetone	15	13	12	10	-	7		
		Water	-	-	-	-	-	-		

MTCC- Microbial Type Culture Collection C- Control

DISCUSSION

Sudarshan Churna contains 42 different constituents including 50% of *Swertia chirata* Buch Ham and the formulation is described in the ancient ayurvedic literature. A survey on the activities of the constituents revealed that *Swertia chirata*, *Ureria picta*, *Curcuma longa*, *Terminalia chebula*, *Asparagus racemosus*, *Acorus calamus*, *Zingiber officinale*, *Azadiracta indica*, *Glycerrhyza glabra* are reported to be effective as antimicrobial herbs (18-23). The *Sudarshan Churna* contains flavonoids and sterol, which may responsible for antimicrobial activity (17, 24-25).

The reported antimicrobial activity in the present study seems to be the outcome of antimicrobial action of its active components like flavonoids and sterols.

CONCLUSION

Our findings suggest that the Ayurvedic herbal preparation *Sudarshan Churna* extracts have antimicrobial properties and they can be used in the treatment of infectious diseases. On comparing the zone of inhibition of *Sudarshan Churna* extract to that of standard antibiotics (Amikacin and Norfloxacin) it is concluded that, the formulation in question shows activity against *E. coli*. whereas the standard antibiotics shows no activity i.e. the bacterial strain is resistant towards these two antibiotics. The most active extract can be further evaluated pharmacologically as well as for its chemically active components in the formulation.

ACKNOWLEDGMENT

The authors are very grateful to Hon'ble Vice Chancellor, Dean (College Development) Baba Farid University of Health Sciences, Faridkot and Principal, GGSMC, Faridkot for providing support and facilities for completing this work.

REFERENCES

1. Govind D. Bhaishajya Ratnavali, New Delhi; Motilal Banarasidas publishers: 2002, pp. 461.
2. Eisenberg DM, Davis RB, Ettner SL. Trends in alternative medicine use in the United State 1990-1997: results of a fallow up national survey. JAMA, 1998; 280(18): 1569-1575.
3. Bansal R, Bansal P, Gupta V, Kumar S, Sharma S, Rao MM. Drug designing through antimicrobial potentials of Indian herbs. J Pharmacy Research, 2010; 3(2): 364-370.

4. Bibitha B, Jisha VK, Salitha CV, Mohan S, Valsa AK. Antibacterial activity of different plant extracts. Short communication. Indian J Microbiol, 2002; 42: 361-363.
5. Maghrani M, Zeggwah N, Michel J, Eddouks M. Antihypertensive effect of *Lepidium sativum* in spontaneously hypertensive rats. J Ethnopharm, 2005; 102(1-2): 193-197.
6. Doughari JH. Antimicrobial activity of *Tamarindus indica* Linn. Tropical J Pharm. Res. 2006; 5(2): 597-603.
7. Ben SA, Barzallah SF, Aouni M. Investigation of some medicinal plants from Tunisia for antimicrobial activities. Pharmaceut Biol., 2007; 15(5): 421-428.
8. Coruh I, Gornez AA, Ercisli S. Total phenolics, mineral elements, antioxidant and antibacterial activities of some edible wild plants in Turkey. Asian J Chem, 2007; 19(7): 5755-5762.
9. Bhargava S, Bhargava P, Saraf S, Pandey R, Sukla SS, Garg R. Evaluation of Antipyretic activity of *Sudarshan Churna*: an Ayurvedic formulation. Journal of Research Education of Indian Medicines, 2008; XIV(2): 11-14.
10. Gupta AK. Introduction to Pharmaceutics –I. 3rd ed., New Delhi; CBS Publishers: 2002.
11. The Ayurvedic Formulary of India Part- I, Government of India, Ministry of Health and Family Welfare, Department of Indian System of Medicine and Homeopathy, New Delhi. 2nd ed., Delhi; Controller of Publications Civil Lines: 2003, pp.116-117,461-480.
12. Parekh J, Jadeja D, Chanda S. Efficacy of Aqueous and Methanol extract of Some Medicinal Plants for Potential antibacterial activity. Turk J Biol, 2005; 29: 203-310.
13. Tambekar DH, Dahikar SB, Lahare MD. Antibacterial Potentials of Some Herbal Preparations Available in India. Research J of Medicine and Medical Sci, 2009; 4(2): 224-227.
14. Sharma S, Sharma MC. Studies of antimicrobial activity of ethanolic plant extract of *Mollugo pentaphylla* Linn. Archives of Applied Sciences Research, 2010; 2(1): 242-246.
15. NCCLS (National Committee for Clinical Laboratory Standards) Performance Standards for antimicrobial susceptibility testing. 8th Informational Supplement. M 100 S12. National Committee for Clinical Laboratory Standards, Villanova, Paris; 2002.
16. Kumar M, Agarwal RC, Dey S, Rai VK, Johnson B. Antimicrobial activity of aqueous extract of *Terminalia chebula* Retz. on Gram-Positive and Gram- Negative micro-organisms. International J Curr Pharmaceutical Res, 2009; 1(1): 56-60.
17. Sharma PV. Alkaloids of *Swertia chirata* Buch-Ham. Indian J Pharm Sciences, 1982; 2: 36.

18. Chopra RN, Nayar SL, Chopra IC. Glossary of Indian Medicinal Plants (Including the Supplement). New Delhi; Council of Scientific and Industrial Research: 1986.
19. Gurudeva MR. Botanical and Vernacular Names of South Indian Plants, Bangalore, India; Divyachandra Prakashana: 2001.
20. Indian Herbal Pharmacopoeia. Mumbai; Indian Drug Manufacturer Association: 2002, pp.165-66.
21. Kokate CK. Textbook of Pharmacognosy. 3rd ed., India; Nirali Prakashan: 1995.
22. Singh B, Gupta V, Bansal P, Kumar D, Murali KC. Pharmacological Potential of Polyherbal Formulation, Sudarshan Churna – A Review. International Journal of Ayurvedic Medicine, 2011; 2(2): 52-61.
23. The Wealth of India– A dictionary of Indian raw materials and industrial products. Council of Scientific and Industrial Research Publishers, Government of India: 1962.
24. Bhargava S. Evaluation of Antibacterial activity of aqueous extract of *Swertia chirata* Buch. Ham. Root. International J. Green Pharmacy, 2007; 1(2): 51-52.
25. Kirtikar KR, Basu BD. Indian Medicinal Plants. II ed(3): Allahabad; 1935: 1664-1666.