Pharmacological Research

In vitro antibacterial and antifungal activities of *Cassia fistula* Linn. fruit pulp extracts

Bhalodia N. R., Nariya P. B.¹, Acharya R. N.², Shukla V. J.³

Research Associate, ADL, Zydus Cadila Healthcare, Ahmedabad, ¹Research Scientist, ADL, Ramniklal Manikchand Dharwai Research Centre, Valsad, ²Associate Professor, Department of Dravyaguna, ³Head, Pharmaceutical Chemistry Laboratory, Institute for Post Graduate Teaching and Research in Ayurveda, Gujarat Ayurved University, Jamnagar, Gujarat, India

Abstract

Aim of the study is to assess the antimicrobial activity *Cassia fistula* fruit pulp extracts on some bacterial and fungal strains. Hydro alcohol and chloroform extracts of *Cassia fistula* fruit pulp were evaluated for the potential antimicrobial activity. The antimicrobial activity was determined in both the extracts using the agar disc diffusion method. Extracts were effective on tested microorganisms. The antibacterial and antifungal activities of solvent extracts (5, 25, 50, 100, 250 μg/mL) of *C. fistula* were tested against two gram positive, two gram negative human pathogenic bacteria and three fungi, respectively. Crude extracts of *C. fistula* exhibited moderate to strong activity against most of the bacteria tested. The tested bacterial strains were *Staphylococcus aureus*, *Streptococcus pyogenes*, *Escherichia coil*, *Pseudomonas aeruginosa*, and fungal strains were *Aspergillus niger*, *Aspergillus clavatus*, *Candida albicans*. The antibacterial potential of the extracts were found to be dose dependent. The antibacterial activities of the *C. fistula* were due to the presence of various secondary metabolites. Hence, these plants can be used to discover bioactive natural products that may serve as leads in the development of new pharmaceuticals research activities.

Key words: Antibacterial activity, Antifungal activity, Bacteria, *Cassia fistula*

Introduction

Antibiotics are one of the most important weapons in fighting bacterial infections and have greatly benefited the health-related quality of human life since their introduction. Drugs derived from natural sources play a significant role in the prevention and treatment of diseases. In many countries, traditional medicine is one of the primary health care system.¹,² Herbs are widely exploited in the traditional medicine and their curative potentials are well documented.³ About 61% of new drugs developed between 1981 and 2002 were based on natural products and they have been very successful especially in the areas of infectious disease and cancer.⁴ Recent trends, however, shows that the discovery rate of active novel chemical entities is declining.⁵ Natural products of higher plants, may give a new source of antimicrobial agents with possibly novel mechanisms of action.⁶,⁷ The effects of plant extracts on bacteria have been studied by a very large number of researchers in different parts of the world.⁸ Much work has been done on ethno medicinal plants in India.⁹

In the recent years, researches on medicinal plants have attracted a lot of attention globally. Evidences have been accumulated to demonstrate the promising potential of medicinal plants used in various traditional, complementary, and alternative systems of treatment of human diseases. Plants are rich in a wide variety of secondary metabolites such as tannins, terpenoids, alkaloids, flavonoids, glycosides etc, which have been found in vitro to have antimicrobial properties.¹⁰,¹¹

Herbal medicines have been known to man since centuries. Therapeutic efficacy of many indigenous plants for several disorders have been described by practitioners of traditional medicine.¹² Antimicrobial properties of medicinal plants are being increasingly reported from different parts of the world. Traditional medicine continues to be a valuable source of remedies that have been used by millions of people around the world to secure their health.¹³ The pharmaceutical industries have produced a number of new antibiotics, resistance to these drugs by microorganisms has increased. In general, bacteria has the genetic ability to transmit and acquire resistance to synthetic drugs that are utilized as therapeutic agents.¹⁴

Therefore, actions must be taken to reduce this problem, such

Address for correspondence: Mr. Nayan R. Bhalodia, Panchavati Gaushala, 95-B, Saru Section Road, Opposite Gita Bungalow, Jamnagar, Gujarat, India. E-mail: nayanbhalodia@gmail.com
as to minimize the use of antibiotics and to continue studies to develop new drugs, either synthetic or natural to control pathogenic microorganism. In an effort to expand the spectrum of antibacterial agents from natural resources, Aragvadha (Cassia fistula) belongs to Caesalpiniaaceae, which is subfamily of Leguminosae[11-18] has been attempted in this study.

Cassia fistula Linn, a semi wild Indian Laburnum also known as the Golden Shower, is distributed in various countries including Asia, Mauritius, South Africa, Mexico, China, West Indies, East Africa, and Brazil as an ornamental tree for its beautiful branches of yellow flowers. Recognize by the British pharmacopoeia.[19] It is widely used for its medicinal properties, its main property being that of a mild laxative suitable for children and pregnant women. It is also a purgative due to the wax aloin and a tonic.[20] It has been reported to treat many other intestinal disorders like healing ulcers.[21,22] The plant has a high therapeutic value and it exerts an antipyretic and analgesic effect.[23]

In the Indian literature, this plant has been described to be useful against skin diseases, liver troubles, tuberous glands and its use in the treatment of haematemesis, pruritus, leucoderma, and diabetes has been suggested.[24,25] C. fistula extract is used as an antiperiodic agent and in the treatment of rheumatism. It has been concluded that plant parts could be used as a therapeutic agent in the treatment of hypercholesterolemia partially due to their fiber and mucilage content.[26] Beside its pharmacological uses, the plant extract is also recommended as a pest and disease control agents in India.[27-29] This plant is widely used by tribal people to treat various ailments including ringworm and other fungal skin infections.[30] It is used by Malalais tribe in India to treat nasal infection.[31] The pulp of the ripe fruits has mild, pleasant purgative action and is also used as an antifungal drug.[32] The whole plant is used to treat diarrhea; fruits are used to treat skin diseases, fever, abdominal pain, leprosy by traditional people. C. fistula plant organs are known to be an important source of secondary metabolites, notably phenolic compounds.[33] C. fistula possesses pharmacological activities such as hypoglycemic, anticancer, abortifacient, anticoagulant, antifertility, estrogenic, laxative, antibacterial, antipyretic, anti-inflammatory, smooth muscle stimulant, antiatheritic, antiinflamatory, purgative, analgesic, antifungal, antiviral, hepatoprotective, anti-implantation.[34-36] C. fistula exhibited significant antimicrobial activity and showed properties that support folklore use in the treatment of some diseases as broad-spectrum antimicrobial agents.[37] Thus C. fistula is well anchored in its traditional uses and has now found widespread acceptance across the world.

The current investigation carried out a screening of hydro alcoholic and chloroform extracts of C. fistula against pathogenic bacteria and fungi in order to detect new sources of antimicrobial agents. Chloroform extracts were obtained after successive separation from hydro alcoholic extracts which were used for further isolation study. So, the microbial activity of this pure chloroform extracts also needed to check the better results against bacterial and fungal strains. This paper reports the results of the antibacterial and antifungal activity of hydro alcoholic and chloroform extracts of fruit pulp.

**Materials and Methods**

**Collection of plant materials**

The fresh and healthy pods of plants C. fistula were collected in June–Aug 2009 from various areas of Jamnagar district, Gujarat, India. Fruit pulp of plant has been collected according to the Ayurvedic references.[19,34,37,38] The plant specimens were identified in the Pharmacognosy Laboratory of I.P.G.T and R.A, Jamnagar. Plant parts were collected on the basis of the information provided in the ethnobotanical Survey of India. Each specimen/plant material was labeled, numbered, annotated, with the date of collection, locality, and their medicinal uses were recorded.

**Extraction**

The extraction of the C. fistula fruit pulp was carried out using known standard procedures.[39] The fresh pulp of C. fistula pod (25.0 g) was extracted with 900 ml of diluted methanol, filtered and evaporated to dryness on a hot water bath to yield a hydro alcoholic crude extract (9.7 g). In addition, further with chloroform in a Soxhlet apparatus, where chloroform is a hydrolysate extract and used for further isolation purpose. The extraction was continued until the extraction was exhausted. The extracts were then combined, filtered, and evaporated to dryness.

Further, the crude hydro alcoholic and chloroform extracts were cooled and filtered. The concentrated hydro alcoholic and chloroform extracts were further subjected for its antimicrobial studies. The residence was dissolved with dimethyl sulfoxide (DMSO) with different concentrations and checked it for the antimicrobial activity.

**Preliminary phytochemical screening**

The extracts were subjected to preliminary phytochemical testing to detect the presence of different chemical groups of compounds. Air-dried and powdered plant materials were screened for the presence of saponins, tannins, alkaloids, flavonoids, triterpenoids, steroids, glycosides, anthraquinones, cumarin, saponins, gum, mucilage, carbohydrates, reducing sugars, starch, protein, and amino acids.[37,40-45]

**Test microorganisms and growth media**

The following microorganisms: S. aureus (MTCC 96), S. pyogenes (MTCC 442), E. coli (MTCC 443), P. aeruginosa (MTCC 424) and fungal strains A. niger (MTCC 282), A. clavatus (MTCC 1323), C. albicans (MTCC 227) were chosen based on their clinical and pharmacological importance.[44] The bacterial strains obtained from Institute of Microbial Technology, Chandigarh were used for evaluating the antimicrobial activity. The bacterial and fungal stock cultures were incubated for 24 h at 37°C on Nutrient Agar and Potato Dextrose Agar medium (Microcare lab., Surat, India) following refrigeration storage at 4°C. The bacterial strains were grown in Mueller–Hilton agar (MHA) plates at 37°C (the bacteria were grown in the nutrient broth at 37°C and maintained on nutrient agar slants at 4°C) whereas the yeasts and molds were grown in sabouraud dextrose agar (SDA) and potato dextrose agar (PDA) media, respectively, at 28°C. The stock cultures were maintained at 4°C.

**Materials and Methods**

**Collection of plant materials**

The fresh and healthy pods of plants C. fistula were collected in June–Aug 2009 from various areas of Jamnagar district, Gujarat, India. Fruit pulp of plant has been collected according to the Ayurvedic references.[19,34,37,38] The plant specimens were identified in the Pharmacognosy Laboratory of I.P.G.T and R.A, Jamnagar. Plant parts were collected on the basis of the information provided in the ethnobotanical Survey of India. Each specimen/plant material was labeled, numbered, annotated, with the date of collection, locality, and their medicinal uses were recorded.

**Extraction**

The extraction of the C. fistula fruit pulp was carried out using known standard procedures.[39] The fresh pulp of C. fistula pod (25.0 g) was extracted with 900 ml of diluted methanol, filtered and evaporated to dryness on a hot water bath to yield a hydro alcoholic crude extract (9.7 g). In addition, further with chloroform in a Soxhlet apparatus, where chloroform is a hydrolysate extract and used for further isolation purpose. The extraction was continued until the extraction was exhausted. The extracts were then combined, filtered, and evaporated to dryness.

Further, the crude hydro alcoholic and chloroform extracts were cooled and filtered. The concentrated hydro alcoholic and chloroform extracts were further subjected for its antimicrobial studies. The residence was dissolved with dimethyl sulfoxide (DMSO) with different concentrations and checked it for the antimicrobial activity.

**Preliminary phytochemical screening**

The extracts were subjected to preliminary phytochemical testing to detect the presence of different chemical groups of compounds. Air-dried and powdered plant materials were screened for the presence of saponins, tannins, alkaloids, flavonoids, triterpenoids, steroids, glycosides, anthraquinones, cumarin, saponins, gum, mucilage, carbohydrates, reducing sugars, starch, protein, and amino acids.[37,40-45]

**Test microorganisms and growth media**

The following microorganisms: S. aureus (MTCC 96), S. pyogenes (MTCC 442), E. coli (MTCC 443), P. aeruginosa (MTCC 424) and fungal strains A. niger (MTCC 282), A. clavatus (MTCC 1323), C. albicans (MTCC 227) were chosen based on their clinical and pharmacological importance.[44] The bacterial strains obtained from Institute of Microbial Technology, Chandigarh were used for evaluating the antimicrobial activity. The bacterial and fungal stock cultures were incubated for 24 h at 37°C on Nutrient Agar and Potato Dextrose Agar medium (Microcare lab., Surat, India) following refrigeration storage at 4°C. The bacterial strains were grown in Mueller–Hilton agar (MHA) plates at 37°C (the bacteria were grown in the nutrient broth at 37°C and maintained on nutrient agar slants at 4°C) whereas the yeasts and molds were grown in sabouraud dextrose agar (SDA) and potato dextrose agar (PDA) media, respectively, at 28°C. The stock cultures were maintained at 4°C.
Antimicrobial activity

Determination of zone of inhibition method

In vitro antibacterial and antifungal activity was examined for hydro alcoholic and chloroform extracts. Antibacterial and antifungal activities of plant extracts against four pathogenic bacteria (two gram positive and two gram negative) and three pathogenic fungi were investigated by the agar disc diffusion method.[45–47] Antimicrobial activity testing was carried out by using the Agar cup method. Each purified extracts were dissolved in DMSO, sterilized by filtration using sintered glass filter and stored at 4°C. For the determination of zone of inhibition (ZOI), two gram positive, two gram negative and three fungal strains were taken as a standard antibiotic for comparison of the results. All the extracts were screened for their antibacterial and antifungal activities against the E. coli, P. aeruginosa, S. aureus, S. pyogenes, and the fungi C. albicans, A. niger, and A. clavatus. The sets of five dilutions (5, 25, 50, 100, and 250 μg/mL) of C. fistula extract and standard drugs were prepared in double distilled water using nutrient agar tubes. Muller Hinton sterile agar plates were seeded with indicator bacterial strains (10^6 cfu) and allowed to stay at 37°C for 3 h. The zones of growth inhibition around the disks were measured after 18 to 24 h of incubation at 37°C for bacteria and 48 to 96 h for fungi at 28°C, respectively. The sensitivity of the microorganism species to the plant extracts were determined by measuring the sizes of inhibitory zones (including the diameter of disk) on the agar surface around the disks, and values <8 mm were considered as not active against microorganisms.

Results

Preliminary phytochemical screening

It was found that hydro alcoholic extracts of C. fistula fruit pulp contained tannins, flavonoids, saponins, triterpenoids, steroids, glycosides, anthraquinones, gum and mucilage, reducing sugars, carbohydrates, protein and amino acids, and chloroform extracts contained glycosides, phenolic compounds, tannins, and anthraquinones-type compounds in higher amount [Table 1].

Microbial activity

The antimicrobial activity of both the extracts of C. fistula were studied in different concentrations (5 μg/ml, 25 μg/ml, 50 μg/ml, 100 μg/ml, 250 μg/ml) against four pathogenic bacterial strains two Gram positive (S. aureus MTCC 96, S. pyogenes MTCC 442), two Gram negative (E. coli MTCC 443, P. aeruginosa MTCC 424) and three fungal strains (A. niger MTCC 282, A. clavatus MTCC 1323, C. albicans MTCC 227).

Antibacterial and antifungal potential of extracts were assessed in terms of zone of inhibition of bacterial growth. The results of the antibacterial activities are presented in Tables 2-5.

The antibacterial and antifungal activity of the extracts increased linearly with increase in concentration of extracts (μg/ml). As compared to standard drugs, the results revealed that in both the extracts for the bacterial activity, S. pyogenes were more sensitive as compared to S. aureus, E. coli, and P. aeruginosa, and for fungal activity C. albicans shows good result as compare to A. niger and A. clavatus. The growth inhibition zone measured ranged from 10–20 mm for all the sensitive bacteria, and ranged from 12–21 mm for fungal strains [Figure 1-5].

The results show that both the extracts of C. fistula were found to be more effective against all the microbes tested.

Discussion

Antimicrobial properties of medicinal plants are being increasingly reported from different parts of the world. The World Health Organization estimates that plant extract or their active constituents are used as folk medicine in traditional therapies of 80% of the words population. In this work, both the extracts obtained from C. fistula fruit pulp shows strong activity against most of the tested bacteria and fungal strains. The results were compared with standard antibiotic drugs. In this screening work, no extracts of C. fistula were found to be inactive against any organism, such as Gram positive, Gram negative and fungal strains were resistant to all the extracts of C. fistula.

This study shows the presence of different phytochemicals with the biological activity that can be of the valuable therapeutic index. The result of phytochemicals in the present investigation showed that the plant contains more

---

Table 1: Phytochemical screening of Cassia fistula fruit pulp

| Functional groups | Metabolite (Hydro alcoholic extract of Cassia fistula) |
|-------------------|------------------------------------------------------|
| Alkaloids         | +                                                    |
| Tanins            | +                                                    |
| Flavonoids        | +                                                    |
| Saponins          | +                                                    |
| Triterpenoids     | +                                                    |
| Steroids          | +                                                    |
| Glycosides        | +                                                    |
| Coumarin          | –                                                    |
| Anthraquinones    | +                                                    |
| Reducing sugars   | +                                                    |
| Carbohydrates     | +                                                    |
| Gum and mucilage  | +                                                    |
| Starch            | –                                                    |
| Proteins          | +                                                    |
| Amino acids       | +                                                    |

(+) present and (–) absent

Table 2: Antibacterial activity of hydro alcoholic and chloroform extracts of Cassia fistula [zone of inhibition]

| Microorganism | Zone of inhibition (mm) | Concentration in μg/mL |
|---------------|-------------------------|------------------------|
|               | Hydroalcoholic extracts (μg/ml) | Chloroform extracts (μg/ml) |
| E. coli      |                          |                        |
| P. aeruginosa|                          |                        |
| S. Pyogenes  |                          |                        |
| S. aureus    |                          |                        |

- = No zone of inhibition
or less same components such as saponin, triterpenoids, steroids, glycosides, anthraquinone, flavonoids, gum, mucilage, proteins and aminoacids. From the above results, the activity of hydroalcoholic extracts of *C. fistula* shows significant antibacterial and antifungal activity. The chloroform extracts, whose purity level is good as compared to hydroalcoholic extract and porosity is lower than hydroalcoholic extract, found to be more active against fungal strains. Results shows that this plant is rich in tannin and phenolic compounds have been shown to possess antimicrobial activities against a number of microorganisms.

### Conclusion

Antimicrobial resistance is a global problem. Emergence of multidrug resistance has limited the therapeutic options. Hence,
monitoring resistance is of paramount importance. Hence, this study was aimed to focus the antimicrobial properties of *C. fistula* on gram positive, gram negative, and fungal organisms. In the current investigations, the hydroalcoholic and chloroform extracts of *C. fistula* were found to be active on some isolated microorganism and fungi as compared to standard drugs. This study has justified the traditional use of fruit pulp in infectious conditions. However, before use in human being isolation of pure compound, toxicological study, and pharmacological activity should be carried out thereafter. However, further studies are needed to better evaluate the potential effectiveness of the crude extracts as the antimicrobial agents.

**Acknowledgement**

The Authors are thankful to the Director, I.P.G.T. and R.A., Gujarat Ayurved University, Jamnagar, Gujarat, India for invaluable support and for providing of research facilities. We are also thankful to the Mycrocare laboratory, Surat, Gujarat, India for helping and providing necessary facilities for this research work.

**References**

1. Farnsworth NR. Ethno pharmacology and future drug development: The North American experience. J Ethnopharmacol 1993;38:145-52.
Bhalodia, et al.: Anti-microbial activity of Cassia fistula Linn.

2. Houghton PJ. The role of plants in traditional medicine and current therapy. Altern Complement Med 1995;1:131-43.
3. Dubey NK. Kumar R, Tripathi P. Global promotion of herbal medicines: India's opportunity. Curr Sci 2004;86:37-41.
4. Cragg GM, Newman DJ. Biodiversity: A continuing source of novel drug leads. Pure Appl Chem 2005;77:7-24.
5. Lam KS. New aspects of natural products in drug discovery. Trends Microbiol 2007;15:279-89.
6. Runyoro D, Matee M, Olipa N, Joseph C, Mbwanbo H. Screening of Tanzanian medicinal plants for anti-Candida activity. BMC Complement Altern Med 2006;6:11.
7. Shahidi BH. Evaluation of antimicrobial properties of Iranian medicinal plants against Micrococcus luteus, Serratia marcescens, Klebsiella pneumonia and Bordetella bronchoseptica. Asian J Plant Sci 2004;3:82-6.
8. Reddy PS, Jamil K, Madhusudhan P. Antibacterial activity of isolates from Piper longum and Taxus baccata. Pharm Biol 2001;39:236-8.
9. Maheshwari JK, Singh KK, Saha S. Ethno-botany of tribes of Mirzapur District, Uttar Pradesh. Economic Botany Information Service, NBRI, Lucknow, 1986.
10. Dahanukar SA, Kulkarni RA, Rege NN. Pharmacology of Medicinal Plants and Natural Products. Indian J Pharmacol 2000;32:581-118.
11. Cowan MM. Plant products as anti-microbial agents. Clin Microbiol Rev 1999;12:564-82.
12. Ramasamy S, Charles MA. Antimicrobial eVect of volatile components of selected medicinal plants against human pathogens. Asian J Microbiol Biotech Environ, 2004;6:10-9.
13. Anonymous. Traditional Herbal Remedies for Primary Health Care. Foreword by Dr Samlee Plangbangchang, New Delhi: World Health Organization (WHO), Regional Ofce for South-East Asia, 2010; p.viii.
14. Towers GH, Lopez A, Hudson JB. Antiviral and antimicrobial activities of medicinal plants. J Ethnopharmacol 2001;77:189-96.
15. Col. Sir Henry collett, Flora Simlensis (Flowering plants of Simla), Bengal Army, Third impression. New Connaught Place, Dehliadun: Bishen Singh Mahendarpal Singh; 1971:p.108.
16. Dutta AC. Botany for degree students 6.ed. Calcutta: Oxford University Press; 2002;p. 558.
17. Cooke T. The Flora of the Presidency of Bombay. Vol. 1. Dehliadun: Bishen Singh Mahendarpal Singh; 1976; p. 447.
18. Duttie JF. Flora of the Upper Gangetic Plain. Reprint Edition, Vol.2. Dehliadun: Bishen Singh Mahendarpal Singh; 1994;p.289.
19. Anonymous. The Wealth of India. 1st ed. Vol. 3. New Delhi: Council of ScientiEe and Industrial Research; 1976. p. 337.
20. Satyavati GV, Sharma M. Medicinal Plant in India. New Delhi: ICMR; 1989.
21. Bawas K, Ghosh AB. In: Bharata Banawasadi. Advancement of learning, Vol. 2. Calcutta, India: Calcutta University; 1973;p. 336.
22. Kirtikar KR, Basu BD. Indian Medicinal Plants. 3rd ed. Vol. 4. New Delhi: Jayved Press; 1975.
23. Patel D, Karbhari D, Gulati D, Gokhale D. Antipyretic and analgesic activities of Aconitum spicatum and Cassia fistula. Pharm Biol,1965;157:22-9.
24. Alam MM, Siddiqui MB, Hussain W. Treatment of diabetes throuherbal drugs in rural India. Fitoterapia 1990;61:240-2.
25. Asolkar LV, Kakkar KK, Chakre OJ. Second supplement to glossary of Indian medicinal plant with active principles. New Delhi: Publication and Information Directorate, CSIR; 1992:p. 177.
26. El-Saadany SS, El-Masry RA, Labib SM, Stoty MZ. The biochemical role and hypocholesterolaemic potential of the legume Cassia fistula in hypercholesterolaemic rats. Die Nahrung 1991;35:807-15.
27. Jaiyal S, Sing Z, Chauhan R. Juvenile hormone like activity in extracts of some common Indian plants. Indian J Agric Sci 1983;53:730-3.
28. Sharma BK, Basandrai AK. E'cacy of some plant extracts for the management of Karnal bunt [Neovossia (Tilletia) indica] of wheat Triticum aestivum. Indian J Agric Sci 1999;69:837-9.
29. Raja N, Albert S, Ignacimuthu S. Effect of solvent residues of Vetex negundo Linn. And Cassia fistula Linn. On pulse beetle, Callosobruchus maculates Fab. And its larval parasitoid, Dinarmus vagabundus (Timberlake). Indian J Exp Biol 2000;38:290-2.
30. Rajan S, Baburaj DS, Sethuraman M, Parimala S. Stem and stembark used medicinally by the TribalSrilukas and Paniyas ofNilgiriDistrict,Tamilnadu. Etnobotany 2001;6:19-24.
31. Perumal Samy R, Ignacimuthu S, Sen A. Screening of 34 medicinal plants for antibacterial properties. J Ethnopharmacol 1998;62:173-82.
32. Kasuko I, Nagayo O. E'cfects of vegetable drugs on pathogenic fungi I. E'cfect of antithrombine-glycoside containing crude drugs upon the growth of pathogenic fungi. Bull Pharm Res Inst Jpn 1951:2:23-9.
33. Morimoto S, Nonaka G, Chen R. Studies on leaves of Cassia fistula Linn. ChemPharmac Bull 1998;36:39-47.
34. Dr. Pandey G. DravagynaVijnana. 3rd ed. Reprint, Part3,Varanasi: Chowkhamba Krishna Academy;2005. p. 167.
35. Prof. Lavekar GS. Database on Medicinal Plants used in Ayurveda and Siddha. Vol. 6. Central Council for Research in Ayurveda and Siddha, Department of ASUSH, Ministry of Health and Family Welfare, Government of India; 2009; p. 29.
36. Kumar PV, Chauhan NS, Padh H, Rajani M. Search for antibacterial antifungal agents from selected Indian medicinal plants. J Ethnopharmacol 2006;107:182-8.
37. Anonymous. The Ayurvidc Pharmacopoeia of India.1st ed. Part 1, Vol.3. Govt. of India. Ministry of Health and Family welfare,Dept. of I.S.M. and H., New Delhi, 1990, Reprint 2001.
38. Dhirak AK, Medicinal Plants of Uttaranchal State.1st ed. Varanasi: Chowkhamba Sanskrit Series office, 2004. p.170.
39. Harborne JB. Phytochemical Methods: A Guide to Modern Techniques of Plant Analysis, Chapman Hall, 1984.
40. Khandelwal KR. Practical Pharmacognosy.2nd.ed. Pune: Nirali Prakashan;2009;p.149-56.
41. Kakkanakar BS, Practical Pharmacognosy. Delhi: New Gyan Offset Printers; 2000; p.107-9.
42. Kumar A, Ilavarasan R, Jayachandran, Decaraman M, Aravindhan P Phytochemical Investigation on a tropical plant in South India. Pak J Nutr 2009;8:3-5.
43. Baxi AJ, Shukla VJ, Bhatt UB. Methods of Qualitative Testing of some Ayurvedic Formulation. Jamnagar: Gujarat Ayurved University;2001.
44. McCracken WA, Cowan RA. Clinical and Oral Microbiology. New York: Hemisphere Publishing Corporation; 1983. p.512.
45. Alzoreky NS, Nakahara K. Antibacterial activity of extracts from some edible plants commonly consumed in Asia. Int J Food Microbiol 2003;82:233-20.
46. Bauer AW, Kirby WM, Sherris JC, Turck M. Antibiotic susceptibility testing by standardized single disc method. Am J Clin Pathol 1966;36:493-6.
47. Rios JL, Recio MC, Villar A. Screening methods for natural products with antimicrobial activity: A review of the literature. J Ethnopharmacol 1988;23:127-49.
हिन्दी सारांश
आर्ग्वाथ की फलमज्जा के विभिन्न सारों का इन-विट्रो प्रति सूक्षमजैविक एवं प्रतिक्रियाकृति कर्मों का अध्ययन
भालोडिया एन. आर., नारिया दी. बी., आचार्य आर. एन., शुक्ला दी. जे.
इस अध्ययन का उद्देश्य औषधीय सार के प्रति सूक्षमजैविक कर्म का मूल्यांकन एवं कुछ सूक्षमजैविक व fungal strains पर ‘जोन ऑफ इंटीरियोन’ का निरीक्षण करना है। प्रस्तुत अध्ययन में आर्ग्वाथफलमज्जा के हाइड्रोअल्कोहोलिक सार एवं क्लोरोफिल सार का विकिरण रूप से महत्वपूर्ण सूक्षमजैवाणुओं एवं कर्मों (fungal strains) के प्रति रोधी कर्म का आध्ययन किया गया। यह अध्ययन अगार डिस्क डिस्पुजन विधि द्वारा दो Gram +ve जीवाणु प्रजातियाँ S.aureus, S.pyogenes, दो Gram - ve जीवाणु प्रजातियाँ E.coli, P.aeruginosa तथा तीनों कर्म प्रजातियाँ A.niger, A.clavatus एवं C.albicans में किया गया। दोनों प्रकार के सार मात्रानुसंधार प्रभावी पाये गये। फाइटोकेमिकल विश्लेषण द्वारा उपायकारी तत्त्वों की उपस्थिति में प्रतिसूक्षम जैविक कर्म पाया गया है। इस प्रकार वनस्पतियों संक्रिय प्राकृतिक जैव उत्पाद शोध में कार्यरत नवीन फार्मास्यूटिकल शोध कार्यों के विकास में महत्वपूर्ण योगदान दे सकती हैं।