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RESEARCH ARTICLE

Evaluation of antimicrobial and phytochemical profile of medicinally important herb Mentha *arvensis* (L) against various microorganism

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Abstract

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Key words: Mentha arvensis , phytochemical and antimicrobial activity. The present study has been carried out to evaluate the Phytochemical analysis and antimicrobial potential of medicinally important herb Mentha *arvensis* belonging to family Lamioceac commonly called as mint and Pudina. The antimicrobial potential of plant is measured against some of the important microorganism such as E.coli , Streptococcus aureus and salmonella typhi. The plant material has been extracted by using the solvent Methenol and chloroform .The present study reveals that the selected herb extracted of methanol found to be more bioactive against salmonella typhi and E-colo rather than streptococcus aureus , Similarlly the chloroform extract is found to be more bioactive against the microorganism E-coli and salmonella typhi rather than streptococcus aureus.

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Introduction

India is rich in rare and useful herbs from which medicines can be prepared. medicinal plant have a long history of use and their use is over the worldwide .in higher aromatic plants have traditionally been used in folk medicine .It also extend the shelf life of foods .The genus Mentha arvensis from family Lamiaceae include large number of species family Lamiacea posses pharmacological and commercial maximum significant . Mentha arvensis found in various countries which is popularly known as Pudina, common house hold remedy in India. It is highly consider as a antiseptic, antipyretic, , antimicrobial, anti-aging properties .The active stimulant with components of herbal remedies have the advantage with many other substance. The effect of the plant extract on bacteria have been studied worldwide by large number of researcher such as Farnz Malik at el.(2012) has evaluated the phytochemical, antiallergic and anti-inflmmatory activity of against in animals . Similarly the same important antimicrobial Mentha arvensis also reported by work of Sunghandhi and Meera Bai (2011) with some microorganisms. Suresh at el .(2012), Gupta et al,.(2010) John et al (2012), and Henrique et al. (2010) has studied the phtotoxic properties of Mentha arvensi.

Londonkar and Poddar (2009) has evaluated activity of extracts of Mentha arvensis Linn against drug induced gastric ulcer in some mammals. Srinivasand Arun has used and investigated the Mentha extract clinically for the treatment of UTI patients and found the improvement.

Material and Methods:

Fresh Mentha *arvensis* were collected aseptically from Marathwada Agricultural University Parbhani and brought to labrotary and only leaves ware collected and washed under running tap water ,washed leaves were dried under room temperature using blander dried leaves were grinding to fine powder. Powder was preserved in air tide container for the further studies

Solvent Extract Preparation:

50 gms of plant material were taken in soxhelt apparatus. For extraction different solvents were taken such as Ethanol and Chloroform were taken for extraction. Extraction process have been carried out 6-8 hrs with 250 ml of each solvent the extract was filtered through the whatman's filter paper No. 1 and clear liquid was collected and the solvent was evaporated to make final volume one fourth of the original volume.

Qualitative Phyto chemical Analysis:

The Methanolic extract and Chloroform extract were used for the preliminary qualitative phytochemial analysis by using the standard biochemical procedure and method. The important phytochemical examination was carried out for the presence of Alkoliods, Polyphenols , Flavonoids , Tannins, Saponins, The Diterpenes were identified by using of Harmoti and Amrani (2008). the methods Roopshree et al.(2008), Ventura et al. (2008) . The Alkolids were detected by using Dragendorffs test and Wagners test, Polyphenols were determined by Ferric chloride test. Flavonoids were identified by using the Alkaline reagent test and lead acetate test .The Tannins was identified by Geletin test .Forth and foam Test was used for the Saphonins, similarly the Diterpenes was detected by using Copper acetate test.

Microorganisms and preparation of bacterial Suspension:

The microorganism, E-coli, Streptococcus aureus and Salmonella typhi were obtained from microbial type culture from the same college Microbiology Department .The bacterial culture strain was cultured in nutrient broth at 37^{0} C and maintained on nutrient agar (Hi-media) slant at 4^{0} C. The previously identified microorganism was inoculated at 35 ± 2^{0} C for 5 hours. A loop of pure slant culture is mixed with sterilised distilled water in a test tube under aseptic condition and content was thoroughly mixed to form a uniform suspension.

Determination of antimicrobial Activity:

For the evaluation of antimicrobial activity standard Agar Disc diffusion method was used described by shanti et al., (2011) in which Muller Helton Agar was used. The plates were swabbed with bacterial suspension of each species used. The sterilised filter paper disc were socked with the test extract and dried and placed on surface of each inoculated plates. The plates were kept for overnight at 37°c. The organism was tested in triplicate .The test Methanolic and Chloroform extract showing the antimicrobial growth activity inhibiting the of selected microorganism such as E-coli, Streptococcus aureus and Salmonella typhi and a clear distinct zone of inhibition was found around the disc. The antimicrobial activity of the extracts was tested by measuring the diameter of zone of inhibition by Himedia scale in mm.

Results and Discussion:

Preliminary phytochemical analysis

The preliminary qualitative phytochemical analysis of the Mentha arvensis leaf extract was carried out and the secondary metabolites like Alkiloidses , Phenolic compound, Flavonoids ,Tannin and Diterpenes was found to be present showing the positive test and the Saponins was found to be absent by the qualitative test. The table no.1 shows the presence and absence of secondary metabolite from the Mentha arvensis .The similar findings were also reported by John De Britto et al., (2012) suresh (2012) and Sugandhi and Meera Bai (2011).Thus indicating the plant extract is a good source of secondary metabolites having a important role in human life.

Antimicrobial activity:

For determination of antimicrobial activity the Mntha arvensisi leaf was used to prepare the crude extract in solvent methanol and chloroform. The methanolic leaf extract was tested against pathogenic organisms like E-coli, Streptococcus areus and salmonella thypi .The methenolic crude extract show the higher degree of antimicrobial activity against salmonella typhi at higher concentration (50%)Showing the 8.0mm zone of inhibition than with E-coli 7.0mm and streptococcus arenus Thus the methenolic leaf extract is found good bioactive compounds to inhibit the growth of salmonella typhi at higher concentration then the lower concentration(Graph -1) . similarly the crude chloroform extract has tested against the above pathogens and found to be more bioactive showing the maximum zone of inhibition 8.0mm against the E-coli with the higher concentration(50%) rather than the other test organisms like salmonella typhi with 7.0mm and S. arenus with 6.0mm zone of inhibition (Table -2,Graph-2)

Thus the present study reveals that the Mentha arvensis is contains the important bioactive compound is secondary metabolite and found to be antimicrobial potential against the selected microorganisms at the higher concentration with the different solvent.

Phytochemical compounds	Test	Results
Alkaloids	Dragenderoffs and Wangners	+ve
Phenolic Compounds	Ferric chloride Test	+ve
Flavonoids	Alkaline reagent test lead acetate test	+ve
Saphonin	Forth Foam Test	-ve
Tannins	Gelatin Test	+ve
Diterpenes	Cooper Test	+ve

Table No. 1 Preliminary phyto-chemical Analysis of the Mentha arvensis (L.)

Table No. 2 Antimicrobial activity of the Mentha arvensis by using various solvents.

Bacteria	Concentration %	Methenolic Extract Zone of Inhibition(mm)	Chloroform Extract Zone of Inhibition(mm)
E-Coli	20%	1.6	2.0
	30%	6.0	7.0
	50%	7.0	8.0
Streptococcus Aureus	20% 30% 50%	2.0 5.2 6.0	104 5.0 6.0
Solmonella Typhi	20%	3.0	1.8
	30%	7.0	6.0
	50%	8.0	7.0





Conclusion:

The antimicrobial potential of the plant may be attributed to the various compound present in the crude extracts. The purified compound may be more potential and significant against the selected microorganism .Further studies require isolation and characterization of individual bioactive compound for the pharmaceutical use.

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