

Research Article

Evaluation of the antibacterial and antifungal potential of spider saxifrage plant (*Saxifraga flagellaris* Willd.)

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Abstract

The current research work was performed to find antifungal and antibacterial potential of spider saxifrage plant. The aim was to find a new source against the selected pathogenic bacterial and fungal strains. *Saxifraga flagellaris* extract was fractionated. Five fractions were prepared and tested for antimicrobial potential against the selected fungal and bacterial strains. The highest antifungal activity (35-65%) was displayed by the crude methanolic extract against all the selected fungal stains and the lowest antifungal trend was exhibited by the aqueous fraction. However in antibacterial activity the n-hexane fraction showed maximum antibacterial activity (46-48%) against the selected pathogens and more or less same trend was followed by crude methanolic extract, while the least inhibition was shown by aqueous fraction. The other fractions though remained with moderate zone of inhibition but were significant.

Keywords: Antibacterial; Antifungal; Inhibition; Spider Saxifrage; *Saxifraga flagellaris*

Introduction

Plants' primary or secondary metabolites are natural products. Primary metabolites are necessary for the growth and development while secondary metabolites govern the biochemical pathways [1]. Secondary metabolites are formed from the primary metabolites via biosynthetic pathways and may or may not be linked to the growth of the plant but have significant ecological functions. The commercial applications of these extracts include different flavorings, fragrances and drugs synthesis etc. [2]. There

is enormous diversity in the nature of the plants derived secondary metabolites. Plants have medicinal properties due to the presence of certain secondary metabolites [3]. According to [4] these compounds are specific at different taxonomic ranks like subspecies, species, genera and families. About 50% of the medicines introduced into the market in the last two decades have been obtained from medicinal plants [5]. Modern screening methods have given medicinal plants amazing importance because of the enhanced biological impact, structural

confirmation and isolation of active metabolites. Currently three generations of medicinal plants are identified [6]. Based on the enormous availability of the biochemical variety; medicinal plants provide a limitless resource of the new drugs discovery [7, 8]. About 25% of the medications and 66% of the anti-inflammatory and anticancer drugs in market and laboratory trials are of the medicinal plants origin [9]. Based on the fact that microbes become resistant over time, the antibiotics; which play a fundamental role in the treatment of infectious diseases; become less effective which results in the increasing failures of the therapies. These diseases caused about 50% of the deaths globally and became the 3rd major reason of deaths [10]. For the treatment of bacterial diseases hundreds of plants derived products are used [11]. The use of plant natural products is ever increasing by the local communities. Berberine and quinines are examples of these products [12]. Similarly liposomes can be used to detoxify the toxins produced by *Streptococcus pneumonia* and *S. aureus*. *Azorella cryptantha*'s product Chrysolthol exhibit very strong activities against *S. enteritidis* and *E. coli* [13]. Similarly the leaves extract of *Catharanthus roseus* were active against the species of *Streptococcus*, *Pseudomonas* and *E. coli* [14]. Fungi are one of the infectious agents causing dreadful diseases and posing as a cause among the others in global health problems [15]. Fungi have dual threatening effects as these not only cause severe ailments in economic crops but also allergies and toxicities [16, 17]. Fungicides of biological origin have commonly been used to cure fungal ailments. Due to the continuous arise of the resistant varieties there is an ever growing demand for harmless alternative and useful compounds [18]. The species, e.g. *Aloe vera* containing C-glucoside Barbaloin, with effective antifungal properties can yield biologically active products [19, 20]. Assessment of the

antifungal activities of local Brazilian medicinal plants showed that the most vigorous were the methanolic extracts of *Vernonia polyanthes* [21]. Similarly the growth of *Mycosphaerella eumusae* can be inhibited by different extracts obtained from *Allium sativum* [22]. In this regard the current research work was designed with the aim to evaluate the antibacterial and antifungal activities of *Saxifraga flagellaris* WILLD.

Materials and methods

Antibacterial activity

Media preparation and agar well diffusion method

Following [23] agar well diffusion Method was used for antibacterial activity. 1 liter distilled of water was used to liquefy 25g of Luria Broth, Miller powder at pH adjusted at 7.0. 100ml of this media were autoclaved in 250ml flask. The selected bacterial strains were introduced into the flasks and kept overnight at 150RPM at 37^oC. The agar was allowed to solidify in the petri-dishes and after that 5 holes were tunneled through a sterilized borer. The tunnels were then introduced with the inoculums. The bacterial and fungal strains were selected based on their frequent occurrence in the local hospitals and their resistance to common drugs.

Tested bacterial strains

The selected fungal strains included three gram positive strains i.e. *Staphylococcus aureus*, *Streptococcus mutans* and Methicillin-resistant *Staphylococcus aureus* (MRSA) and a single gram negative strain i.e. *Serratia marcescens*.

Extract preparation and measurement of zones of inhibition

The crude extract of *Saxifraga flagellaris* were dissolved in 20mg/ml of Dimethyl sulfoxide (DMSO). Standard antibiotic; Cefotaxime (2mg/ml) was used as positive control while pure DMSO as negative control. 75µl of the plant fraction was introduced into the wells in petri-dishes. The

samples were incubated for 24 hours at 37°C. After incubation diameter of the transparent zone, around each well was measured. The experiment was repeated thrice and standard deviation was calculated from the data obtained.

Antifungal activity

Media preparation and agar well diffusion method

Potato dextrose agar (39g) was taken in liter distilled water (1 liter). The mixture was sterilized at 15PSI for 15 minutes in the autoclave. It was cooled at room temperature and poured into the petri-dishes to solidify. The method of [24] i.e. agar well diffusion method was followed. 100µl of different strains of the selected fungal strains were placed on the surface of agar plate through a micropipette. The culture was spread by a sterilized inoculation loop. Holes were made in the culture (Plate 1) via a sterile cork borer. Crude extract (75ml) of the *Saxifraga flagellaris* were added to the petri-dishes and incubated at 37°C for 24 hours. Results were calculated by measuring the clear areas around each hole which indicated the extent of antifungal activity of each extract against the tested fungal strain. Each test was repeated thrice and the data was analyzed statistically.

Tested fungal strains

The selected fungal strains included *Alternaria alternate*, *Aspergillus flavus*, *Polysphondylium pallidum* and *Fusarium oxysporum*.

Results and discussion

The current research work was carried out to evaluate the anti-fungal and anti-bacterial potential of the spider saxifrage plant (*Saxifraga flagellaris* WILLD.) plant. The results revealed that this plant is a very effective source for the inhibition of the growth of the tested species of both fungi and bacteria. The plant can be investigated for active metabolites to evolve the anti-bacterial

and anti-fungal drugs through biochemical and biophysical essays.

Antibacterial activities of *Saxifraga flagellaris*

In the current research work five fractions of *Saxifraga flagellaris* extracts were used to know their potential use against the selected bacterial strains that are *Staphylococcus aureus*, *Streptococcus mutans* and Methicillin-resistant *Staphylococcus aureus* (MRSA) and *Serratia marcescens*. These species were selected for the current experiments on the basis of their frequent pathological reports from hospitals of Khyber Pakhtunkhwa. Results of the anti-fungal activity as shown in (Table 1) showed that all the fractions were active against the tested bacterial strains. The growth of *Staphylococcus aureus*, *Streptococcus mutans*, *Serratia marcescens* and Methicillin-resistant *Staphylococcus aureus* (MRSA) was inhibited by the crude methanolic extract with 10.0mm, 12.0mm, 09.0mm and 11.0mm zones of inhibition respectively. The *n*. hexane extracted fractions inhibited the growth of *Staphylococcus aureus*, *Streptococcus mutans*, *Serratia marcescens* and Methicillin-resistant *Staphylococcus aureus* (MRSA) with 12.0mm, 13.0mm, 14.0mm and 10.0mm zones of inhibition respectively. The chloroform fractions were produced 11.0mm, 14.0mm, 07.0mm and 14.0mm zones of inhibition against *Staphylococcus aureus*, *Streptococcus mutans*, *Serratia marcescens* and Methicillin-resistant *Staphylococcus aureus* (MRSA) respectively. The ethyl acetate fraction was most active against *Streptococcus mutans* (ZOI=10.0mm) followed by Methicillin-resistant *Staphylococcus aureus* (ZOI=09.0mm), *Staphylococcus aureus* (ZOI=08.0mm) and *Serratia marcescens* (ZOI=08.0mm). The aqueous extracted fraction inhibited the growth of tested bacterial strains in the order of Methicillin-

resistant *Staphylococcus aureus* (ZOI=08.0mm), *Staphylococcus aureus* (ZOI=05.0mm), *Serratia marcescens* (ZOI=05.0mm) and *Streptococcus mutans* (ZOI=04.0mm) respectively. Globally several hundred genera of medicinal plants are used as the main sources of anti-bacterial drugs [25, 26]. According to [27, 28] in the previous century most of the faculties in the medicine sector switched their concern from natural to synthetic drugs. But for the last few decades this trend is shifting in the reverse direction [29]. In a survey 25% of the drug prescriptions in 35 countries are plants derived in non-modified or little modified [30, 31].

Antifungal activities of *Saxifraga flagellaris*

Results of the anti-fungal activity as shown in (Table 2) showed that the n-hexane extracted sample inhibited the growth of *Fusarium oxysporum* completely. Chloroform and Ethyl acetate inhibited the growth of *Polysphondylium pallidum*. While the aqueous extract showed significant activity against *Aspergillus flavus* and *Alternaria alternate*. Crude methanolic extract showed good results against *Polysphondylium pallidum*. Trend in results revealed that *Fusarium oxysporum* is comparatively more sensitive to the n-hexane extracts, *Polysphondylium pallidum* to chloroform and Ethyl acetate extracts while *Aspergillus flavus* and *Alternaria alternate* to the aqueous extracts of plants. The usage of

medicinal plants as anti-fungals is quite old practice performed by human beings. These result in the production of important compounds that are used to develop drugs against fungal pathogens. One of the positive aspects of these drugs is this that these drugs have very little adverse/side effects on the human health [32]. The demand for the development anti-fungal drugs has enormously increased as it is established that among others fungal pathogens are causing serious losses in the quantity, quality, shelf life and mortality of the crop plants [33]. In regard to humans, fungi as the cause of diseases offer a new aspect of human pathology [34]. In this regard medicinal plants are proved to be the sources for the discovery of new drugs [35]. In the developing countries, which face the problems of the quantity and quality of modern drugs, medicinal plants derived compounds have recently gained much importance and their old traditional uses are valued much more nowadays [36-38]. The current study revealed that antifungal activity of the wild plants proved to be much more significant and can be a very suitable substitute of the modern medicines. Similar findings have been reported by [39] for the fungus *Alternaria alternate*. Similarly [40] revealed that the crude methanolic extract of the medicinal plants showed maximum anti-fungal properties which is in agreement with our work. Similar results with other medicinal plants were reported by [41].



Plate 1. Petri Plates showing the antibacterial and antifungal activities of *Saxifraga flagellaris*

Table 1. Antibacterial activity of the crude extracts of *Saxifraga flagellaris* Willd

Bacterial Species	Standard	Cr. Met. Ext		<i>n</i> - hexane		CHCl ₃		EtOAc		Aqueous	
		ZOI	%	ZOI	%	ZOI	%	ZOI	%	ZOI	%
MRSA	25.0±0.67	11.0±0.93	44.00	12.0±0.90	48.00	10.0±0.57	40.00	09.0±0.87	36.00	08.0±0.23	32.00
<i>S. marcescens</i>	21.0±0.87	09.0±0.56	42.86	10.0±0.88	47.62	07.0±0.45	33.33	08.0±0.44	38.1	05.0±0.22	23.81
<i>S. mutans</i>	28.0±0.60	12.0±0.67	42.86	13.0±0.66	46.43	14.0±0.56	50.00	10.0±0.34	35.71	04.0±0.34	14.29
<i>S. aureus</i>	26.0±0.56	10.0±0.87	38.46	12.0±0.93	46.15	11.0±0.21	42.31	08.0±0.33	30.77	05.0±0.50	19.23

Table 2. Antifungal activity of the crude extracts of *Saxifraga flagellaris* Willd

Fungal Species	Standard	Cr. Met. Ext		<i>n</i> - hexane		CHCl ₃		EtOAc		Aqueous	
		ZOI	%	ZOI	%	ZOI	%	ZOI	%	ZOI	%
<i>A. flavus</i>	100.0±0.00	45.0±0.55	45	45.0±0.78	45	30.0±0.67	30	40.0±0.34	40	0.0±0.00	0
<i>A. alternate</i>	100.0±0.00	65.0±0.54	65	30.0±0.88	30	45.0±0.56	45	40.0±0.10	40	0.0±0.00	0
<i>F. oxysporum</i>	100.0±0.00	57.0±0.23	57	00.0±0.00	0	32.0±0.77	32	30.0±0.22	30	23.0±0.34	23
<i>P. pallidum</i>	100.0±0.00	35.0±0.44	35	29.0±0.67	29	00.0±0.00	0	00.0±0.00	0	30.0±0.21	30

Conclusion

It is evident from the results of anti-bacterial and anti-fungal activities that the selected plant i.e. *Saxifraga flagellaris* possessed very significant properties. The crude methanolic and ethanolic extracts were comparatively more active which clarified that polar solvents dissolved more properly in the polar solvents.

Authors' contributions

Conceived and designed the experiments: M Hamayun, KU Rehman & S Wali, Performed the experiments: KU Rehman, S Wali & I Ahmad, Analyzed the data: N Akhtar, B Ullah & S Afzal, Contributed materials/ analysis/ tools: M Hamayun, N Akhtar, B Ullah & S Afzal, Wrote the paper: KU Rehman & S Wali.

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