EFFECT OF SALACIA OBLONGA ROOT EXTRACT AGAINST CLINICAL ISOLATES STAPHYLOCOCCUS AUREUS

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INTRODUCTION

Antimicrobial compounds are very important in reducing the worldwide burden of infectious diseases [1]. However, multidrug-resistant bacteria have become a significant public health risk and sometimes available antimicrobial agents are not effective toward infection caused by pathogenic bacteria. This has led to the search of new antibacterial compounds. From the past many years, plants have been used to treat infectious diseases and are considered as major source of new antimicrobial agents [2,3]. Numerous works have been done to study the antimicrobial effects of herbal plant extracts, including roots, stem, leaves, or flowers [4,5]. Many countries in the world have continued to encourage screening programs of plants used in traditional medicine and identification of their mode of action is important to authenticate their antimicrobial activities and possible inclusion in primary health care.

Salacia oblonga is an important medicinal plant that has been extensively used in traditional Indian Ayurvedic medicine as a liver tonic, anti-inflammatory agent, anodyne ameno-rhoea, diabetes, and treatment of wounds. The root bark extracts are used for itches, asthma, thirst, and ear diseases [6,7]. The important phytoconstituents of S. oblonga include salacinol, mangiferin, and kolutanol, with proven biological activities, namely, nephroprotection, antimutagenic, anti-inflammatory, and antimicrobial [6,8,9]. The present study was undertaken to evaluate the effect of antibacterial activity of S. oblonga root extracts against clinical isolate Staphylococcus aureus.

METHODS

Chemical and reagents

The analytical and high-performance liquid chromatography grade chemicals and reagents used in the study were procured from HiMedia and Merck. Amikacin antibiotic was obtained from Sigma-Aldrich (USA).

RESULTS

In the treated cells, enzyme activity was low compared to the control cells. Extract displayed good growth inhibition and MIC of the extract was 80 µg/ml. DNA fragmentation assay was carried out, in treated and control cells. Extract determined by DNA fragmentation and respiratory dehydrogenase enzyme activity assays.

ABSTRACT

Objective: Salacia oblonga Wall. is an important medicinal plant belonging to the family Celastraceae. The study reports the effect of S. oblonga root extracts against clinical isolate Staphylococcus aureus.

Methods: Antibacterial activity was evaluated by agar diffusion method and assay for minimum inhibitory concentration (MIC) of extract. Further, the effect of S. oblonga extract determined by DNA fragmentation and respiratory dehydrogenase enzyme activity assays.

Results: S. oblonga ethyl acetate root extract was evaluated for antibacterial activity towards clinical isolate S. aureus. Bacterial growth was determined in treated and control cells. Extract displayed good growth inhibition and MIC of the extract was 80 µg/ml. DNA fragmentation assay was carried out, this result has shown that treated bacterial cell has DNA damage compared to the control cell. Further, respiratory dehydrogenase enzyme activity was determined. In the treated cells, enzyme activity was low compared to the control cells.

Conclusion: Salacia oblonga root extract inhibiting the growth of S. aureus by different modes of action.

Keywords: Clinical isolate, Staphylococcus aureus, DNA fragmentation assay, Respiratory dehydrogenase, Salacia oblonga.

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DNA fragmentation assay
DNA fragmentation experiment was done using standard DNA isolation procedure. Inoculate the culture into the LB broth incubated at 37°C overnight. Culture was centrifuged at 10,000 rpm for 10 min and dissolved the pellet in the tris-ethylenediaminetetraacetic acid (EDTA) buffer. Added lysozyme (10 mg/ml) enzyme and 10% sodium dodecyl sulfate to the above suspension and kept it for the incubator at 37°C for 10 min. Added RNase (5 mg/ml) and proteinase K (5 mg/ml) of 5 µl to the above suspension and incubate at 55°C water bath for 1 h. Added equal volume of phenol-chloroform to the above suspension then gently vortex and tubes were centrifuge at 10,000 rpm for 5 min. Transferred top aqueous layer into the fresh Eppendorf repeated this step twice. To this added 100 µl of 3 M sodium acetate and added 0.1 ml volume chilled ethanol kept at −20°C for 30 min. To the pellet added 75% ethanol and kept at 20°C for 10 min. Centrifuge the above suspension at 10,000 rpm for 10 min and dry the pellet. Dissolve the pellet in tris-EDTA buffer. Isolated DNA was resolved on 1% agarose gel electrophoresis.

Effect of extract on enzymatic activity of respiratory chain dehydrogenases in S. aureus
Minimum inhibitory concentration (MIC) of extract was treated to the bacterial cells and incubated to 48 h. Experiment was conducted in the absence of the extract as the control. Cultures were grown at 37°C with shaking at 200 rpm. One milliliter culture was sampled separately from the cultures and centrifuged at 12,000 rpm; then, the supernatants were discarded and the bacteria washed by phosphate-buffered saline (PBS) twice and added 0.9 ml PBS to suspend the bacteria [11]. INT solution (0.1 ml 0.5%) was added, the culture was incubated at 37°C in dark for 2 h, and then 50 µl formaldehyde was added to terminate the reaction. Centrifuged the culture to collect the bacteria, and 250 µl solutions of acetone and ethanol 1:1 in volume were used to distill the iodonitrotetrazolium formazan twice. The supernatants were finally combined. The dehydrogenase activity was then calculated according to the maximum spectrophotometrical absorbance of INF at 490 nm by spectrophotometer (UV–2450 UV–visible spectrophotometer SHIMADZU).

RESULTS
Antibacterial assay
In the present study, the antibacterial activity of S. oblonga EtOAc root extract (1 mg/ml) against clinical isolate S. aureus was evaluated by agar well diffusion method. The extract has shown significant growth inhibition towards pathogen (Fig. 1).

Growth of S. aureus exposed to S. oblonga extract
The growth curves of S. aureus treated with root extract are shown in Fig. 2 by measuring optical density at 540 nm. Under the absence of extract, S. aureus reached exponential phase rapidly. Then the culture was treated with 20, 40, 60 and 80 µg/ml of extract, bacterial cells were lagged at different time point with 20, 40 and 60 µg/ml concentrations. When the concentration of extract was 80 µg/ml, no growth of S. aureus could be detected within 3 days (Fig. 2), indicating that MIC of extract to clinical isolate pathogen S. aureus was 80 µg/ml.

DNA fragmentation assay
DNA fragmentation assay was done, to check the effect of S. oblonga extract against clinical isolate S. aureus DNA. Treated the pathogen with MIC concentration of the extract and resolved on 1% agarose gel (Fig. 3). We have seen smear and fragmented in the treated DNA sample; on the other hand, no fragmentation was observed in control cells without treatment of the extract. This indicated our extract is acting on DNA and inducing the damage that leads to cell death of the pathogen.

Effect of S. oblonga extract on respiratory chain dehydrogenase of S. aureus
The effect of extract on respiratory chain dehydrogenase of S. aureus is shown in Fig. 4. Activity of respiratory chain dehydrogenase in control cells increased slightly with the incubation time. Interestingly, enzyme activity was decreased in treated cells with MIC concentration at 24 h.

Fig. 1: Antibacterial activity of Salacia oblonga root extract against pathogen by agar diffusion method
Fig. 2: Growth curve of Staphylococcus aureus exposed to different concentrations of Salacia oblonga extract
Fig. 3: Effect of Salacia oblonga extract by DNA fragmentation assay, (1) control, (2) treated
Fig. 4: Activity of respiratory chain dehydrogenase in control and treated cells

DISCUSSION
The search for novel antimicrobial became the major goal of various research groups oriented toward medicinal chemistry and pharmacology [12,13]. In the present study, S. oblonga root extract has...
displayed good antibacterial activity by agar diffusion method against clinical isolate S. aureus. Further, bacterial growth was determined by treating the extract at different concentrations and identified that the MIC concentration was 80 µg/ml. To evaluate the effect of extract on DNA, fragmentation assay was carried out. DNA fragmentation was observed in treated cells with MIC concentration of the extract compared with the control cells without the treatment of extract. *Gaesalpinia coriaria* ethanolic extract induces DNA fragmentation toward different pathogenic bacteria [14]. Our *S. oblonga* root extract also induces DNA damage in clinical isolates S. aureus. Hence, the *S. oblonga* root extract has shown antibacterial potential. Respiratory chain dehydrogenase enzyme activity was determined in treated and control cells. Enzyme activity was less compared to the control cells. Diterpene isolated from the medicinal plant *Pseudognaphalium cheiranthifolium* inhibited the bacterial respiratory chain enzyme and displayed antibacterial activity [15]. Phytochemicals from the crude extract of *S. oblonga* may break on cell wall and destroy respiratory chain dehydrogenase, further inhibiting respiration of cells. Phytochemicals present in the crude extract of *S. oblonga* might be inhibited the dehydrogenase enzyme activity and shown antibacterial activity.

**CONCLUSION**

*S. oblonga* root extract displayed good growth inhibition against clinical isolate *S. aureus*; this was confirmed by a agar diffusion method and growth of the bacteria after treatment with extract. The extract was inducing DNA damage and also inhibiting the respiratory chain dehydrogenase enzyme activity. Phytochemicals from the crude extract might be showing different modes of action and inhibiting the growth of the pathogen.

**AUTHORS’ CONTRIBUTIONS**

Dr. Musini and Dr. Zahoorullah, both are designed the work, involved in the experiments and writing the manuscript. Other two authors involved in the experimental work.

**CONFLICTS OF INTEREST**

The authors have no conflicts of interest.

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