ABSTRACT

Objective: To screen the antimicrobial activity Of Callyspongia Diffusa (Marine Sponge) Associated Endophytic Bacterial Strains.

Methods: We have isolated endophytic bacterias CDB-1 and CDB-2 from marine sponge Callyspongia diffusa and identified as Pseudomonas taiwanensis strain and Lysinibacillus sphaericus strain respectively by the phylogenetic analysis. Fractions of CDB-1 and CDB-2 were screened for in vitro and in vivo antimicrobial activities against pathogenic bacteria and mycobacterium tuberculosis H37 RV strain by Minimum Inhibitory Concentration (MIC) method.

Results: The lowest MIC against Klebsiella pneumoniae, Escherichia coli and Enterococcus faecalis was found to be 0.2 µg/ml and 0.4 µg/ml respectively for CDB-2. A significant antifungal activity was observed against Candida albicans (0.2-0.8 µg/ml) and Aspergillus niger (0.2-0.4 µg/ml). Further, Chloroform fraction of CDB-1 and ethyl acetate fraction of CDB-2 have shown significant anti-tubercular activity against the tested organism with MIC of 6.25µg/ml. This was supported by in vivo antimicrobial activity against K. Pneumoniae infection in mice and least haemolytic activity against erythrocytes was observed. Compared to chloramphenicol.

Conclusion: In this study, we have reported the marine species offer a rich source of bioactive metabolites that can exploit to develop novel, useful and potential therapeutic agents.

Keywords: Antimicrobial, Anti-tubercular, Pseudomonas taiwanensis, Lysinibacillus sphaericus, Callyspongia diffusa, Endophytic bacteria

INTRODUCTION

For the treatment of the treatment of many infectious diseases, resistances of antimicrobial compounds by the micro-organisms have become a universal problem. Several infections caused by multi-drug-resistant organism threaten the human health. In developed and developing countries tuberculosis (TB) is a major infectious disease. It is projected that out of 3 persons one will be infected with TB. Particularly in Asia and sub-Saharan Africa. The occurrence of TB in India is around 30%, whereas in China around 4% [1]. Nearly around two million people will die due to TB, with a world case fatality rate of 24% and it is reaching more than 50% in African countries. In India because of more rates of co-existing HIV infection. A person infected with HIV is more susceptible to TB [2]. Even though various treatment regimens for tuberculosis, still it is major problem in so many areas of the world because of drug resistance strains. Thus there is a great need in discovering effective and alternative antimicrobial compounds. In biomedical research, the formulation and improvement of the new antimicrobial drug from a natural source is of great importance. Many antimicrobial agents have been invented by continuous efforts [3].

In the last two decades, research work on natural products obtained from marine plants and animals are the principal focus. Marine environment is the rich source of bioactive constituents with numerous health beneficial effects. Among these organisms, marine sponge and algae species have been pointed as on under-exploited production of secondary metabolites, either by associated microflora or by themselves, to stay alive in their intense habitat [6]. Throughout the years, wide screening programs were developed worldwide and immense efforts have been dedicated aiming of the isolation of new bioactive metabolites from marine microorganisms. Cyclic tetrapeptides isolated from Pseudoolteromonas sp and Pseudomonas sp. of seaweed Diginea sp [12], α-campholeine aldehyde and Lucenin-2 from endophytic fungus aspergillus ochraceus MP2 of marine sponge [13], Artukeminc A and B from endophytic bacteria Rapidithrix sp of marine source [14], 1-(2, 6-dihydroxyphenyl) pentan-1-one from endophytic bacterium Cryptosporiopsis of Clidamiahirta plant [15], 1, 3,8-Trihydroxy-6-methoxy anthraquinone from endophytic bacteria Curvularilalanata of the marine sponge Niphatesolemda [16]. These are some important antimicrobial secondary metabolites obtained from marine sponge derived endophytic microbes. Callyspongia diffusa is a member of a marine sponge, reported to possess antioxidant and antimicrobial activities [17, 18]. Metabolites and vidarabine, marine natural products have gained much research interest [6, 7]. So far, over 20,000 natural products have been isolated and identified from various marine organisms [8].

In general, sponge research is yielding more than 200 new pharmacologically active metabolites per year at present, and several compounds that are derived from sponges or from sponge-associated bacteria are in different phases of clinical trials [9]. Sponges are a rich source of steroids, sulphates and alkaloids. From Micronesian sponge, new sesquiterpene quinols were isolated which possess potent antimicrobial activity [10].

Marine endophytes are of huge interest as rich and novel sources of biologically active compounds. They exist in close relationship with soft-bodied marine organisms, which not have clear structural protection mechanisms, and thus rely on chemical protection by the production of secondary metabolites, either by associated microflora or by themselves, to stay alive in their intense habitat [11]. Throughout the years, wide screening programs were developed worldwide and immense efforts have been dedicated aiming of the isolation of new bioactive metabolites from marine microorganisms.
were isolated from *Virgibacillus* sp. associated with *Callyspongia diffusa* reported the antimicrobial and antioxidant activity [17]. The bacterial strain (CSD 1-CSD10) of *Callyspongia diffusa* showed antimicrobial and immunomodulatory activity [17]. Isolation of bioactive proteins from bacterial species (*Pseudomonas aeruginosa*, *K. pneumoniae*) associated in *Callyspongia diffusa*, and also isolated proteins showed antibacterial activity [20]. The aim of this study was to isolate the microorganisms (endophytes) from the marine natural species which are good candidates for studying, associated microbes and the production of novel secondary metabolites, and which is a great need in finding new classes of natural products that may be effective against resistant strains of microbes, especially from bioactive endophytes from marine source due to their rich source of secondary metabolites and least toxic effects.

**MATERIALS AND METHODS**

**Chemicals used**

Sodium hypochlorite, Distilled water, Tetracycline, Almir blue reagent, Alcohol procured by Sigma-aldrich Chemicals Pvt Ltd, Bangalore, India, and remaining all the chemicals purchased from good companies.

**Instruments used**

Laminar air flow (Reschler), Autoclave (Natsteel), Incubator (Tempo), Refrigerator or deep freezer (Remi), the Rotary flash evaporator (Superfit) etc.

**Collection and authentication of marine sponge**

Sponge sample was collected with the help of fisherman during the lowest low tide from the intertidal zone of Majaaahalli beach, Karwar, Karnataka, India in October 2016. It was authenticated as *Callyspongia diffusa* by Dr. Pratibha Rao, Scientist-in-charge, Central Marine Fisheries research institute, Mangalore, Karnataka, India. The voucher specimen of the algae (Voucher Nos. 14 to 15) was deposited at Pharmacognosy Department, SET’s Pharmacy College, Dharwad, Karnataka, India.

**Isolation of endophytes from Callyspongia diffusa**

The samples were washed with sterile seawater after collection, to remove linked debris. Epiphytes were detached of the sponge. The cleaned sponge material was then surface dried out by pressing it for a short time between paper sheets of towelling and air dried at 30 °C in the shade during 24 h. The microflora present in the surface was detached by washing the sponge samples for 10 to 15 min with 30% ethanol [21]. Then the sample of algae was surface sterilized by 4% sodium hypochlorite for 5 min, 1 min with 70% ethanol and 1 min 2–3 times with distilled water. The sterilized pieces were transferred to alcohol sterilized mortar and triturated into suspension by using distilled water and made the serial dilutions. The diluted suspensions were transferred on sterilized nutrient agar plate. After incubation for 4–5 d at 30 °C, picked up predominant isolates of bacteria and purified. The chosen isolates were experimental their morphological characteristics [22, 23].

**Fermentation and preparation of crude fractions**

Pure isolated bacteria are grown in nutrient agar plates for 4–5 d at 30 °C. These isolated bacteria are identified and given a name as CDB-1 and CDB-2 respectively. CDB-1 and CDB-2 endophytes used for fermentation and extraction process. Purified bacterial isolates were inoculated and fermented individually in Erlenmeyer flask having nutrient broth for 7–14 d at 30 °C under immobile condition. Each broth of bacteria was filtered through 4 layers of cheesecloth and homogenized at 4,000 rpm to squeeze the bacterial cells from broth. The filtrate was solvent extracted with chloroform and ethyl acetate fraction, respectively. Then organic part was separated and dried under reduced pressure using rotary evaporator (SuperfitRotavap, PBU-6) and weighed the crude extract. In this study, CDB-1 and CDB-2 were taken for *in vitro* and *in vivo* antimicrobial activity and further endophytes characterization was studied by sequential analysis. The yield of every extract ranged from 60-80 mg/l of fermented medium [23]. Chloroform and ethyl acetate fractions of CDB-1 and CDB-2 were also subjected to preliminary phytochemical investigations to identify the presence of secondary metabolites.

**Study design**

**Test organisms**

Aspergillus niger, Candida albicans, Enterococcus fecalis, Klebsiella pneumoniae, Bacillus subtilis, Escherichia coli. Which are obtained from Microbiology Laboratory, KMC Mangalore, Karnataka, India.

**In vitro antimicrobial activity**

Nine dilutions of each test samples have made with BHI (Brain Heart Infusion) broth for MIC. Into the 380 µl of BHI broth, 20 µl of the drug was added in the initial tube. 200 µl of BHI broth was added for dilutions into the remaining next 9 tubes individually. Then transferred 200 µl from the initial tube to the first tube consisting 200 µl of BHI broth. It was referred as 10⁻¹ dilution. From 10⁻¹ diluted tube to make 10⁻² dilution. The above serial dilution was repetitive up to 10⁻⁶ dilution for every drug. 5 µl was taken from the maintained stock cultures of necessary organisms and transferred into 2 ml of BHI broth. In every serially diluted tube, 200 µl of above suspension culture was added. Then tubes were incubated for 24 h and turbidity was observed [24].

**In vitro anti-tuberculosis activity**

The anti-TB activity of test fractions was evaluated against *Mycobacterium tuberculosis* H37Rv by using Microplate Alamar Blue assay (MABA). This method is non-toxic and thermally stable reagent is using, and it shows better correlation with BACTEC radiometric method. To minimize the evaporation of the medium in the test walls during incubation 200 µl of sterile deionized water was added to all outer perimeter wells of sterile 96 wells plate. 100 µl of the Middle brook 7H9 broth was added in 96 wells plate, and directly on plate serial dilution of compounds was done. The final drug concentrations tested were 100 to 0.2 µg/ml. Plates were sealed with Parafilm and it was incubated for five days at 37 °C. After incubation time freshly prepared 25 µl of 1:1 mixture of Almar Blue reagent and 10% tween 80 was added to the plate and incubated for 24 h. A blue color in the well was indicated as no bacterial growth and pink color indicates as the growth of bacteria. The MIC was defined as the lowest concentration of the drug that prevents the colour change from blue to pink [25].

**Analysis of cellular toxicity to human erythrocytes**

Chloroform and ethyl acetate endophytic fractions of CDB-1 and CDB-2 were further investigated for cellular toxicity of RBC’s. Blood was obtained from blood bank of KMC hospital, Hubli, Karnataka, India. Human erythrocytes were isolated from the blood by removing theuffy coat and suspended in PBS (10 mmol phosphates, 150 mmol sodium chloride, PH 7.4) which were dispensed in sugar tubes (10[10]cells/500 µl/tube). The serial dilutions of endophytic fractions were made and mixed with erythrocytes keeping the final volume of 1 ml. The cells were incubated for 1hr at 37 °C and finally centrifuged at 1500 g for 10 min. Lysis of the cells was observed by determining absorbance at 600 nm using a colorimeter. The respective dilution of test compounds (without erythrocytes) was used as blank for determination of absorbance. The erythrocytes were completely lysed by treatment with 1% Triton-X100 and absorbance of the released haemoglobin was taken as 100% lysis [26].

**In vivo antitumor activity**

Swiss mice of both male and female (20-22 gms) were used in the study. Standard pellet diet and water ad libitum were given to all the animals. Were given a. Daily animals were checked for their mortality and morbidity till the commencement of the study and only healthy animals were used in the experiment. Techniques used in the experiment like injection, bleeding and sacrifice of the animal were approved by the Animals Ethics Committee as per CPCSEA guidelines (SDCP/AIBC/2014-2). Each animal was challenged by 5 X 10⁶ viable *K. pneumoniae* bacteria in 0.9% of normal saline (200 µl) through intravenous route. After 24 h post infection, the drug treatment was started, suspensions of ethyl acetate and chloroform were administered to all the animals. The study design and test organisms were similar to the *in vitro* antibacterial assay. The test fractions were used for *in vivo* activity. The bioassay was done on human K562 cell line and experimental their morphological characteristics [22, 23].
fractions of CDB-1 was prepared in Tween 80 and orally administered at a dose of 100 mg/kg body weight, normal saline was given to the control group. Administered the test samples for 7 days and necessary precautions were taken to administer a specified dose of the drug to the experimental animals.

Statistical analysis
The values were expressed in mean±SEM (n=6). The results were analyzed by using one-way analysis of variance followed by Anovatukey’s multiple comparison tests to determine the statistical significance between the groups.

PCR sequential analysis of 16s rRNA gene of CDB-1 and CDB-2
The genomic DNA was isolated from the given organism using genomic DNA extraction kit (Bhat Biotech Ltd. Bangalore, India). Amplification of the 16s rDNA gene was performed using the primers. Forward primer: 5'-AGAGTTTGATCCTGGCTCAG-3', Reverse Primer: 5'-ACGGCTACCTTGTTACGACTT-3' PCR was performed as follows in a Master cycler® (DNA-AMP Bhat Biotech) using the following program. Initial Denaturation was carried out at 94 °C for 2 min followed by 40 cycles of denaturation at 94 °C for a minute, annealing at 55 °C for a minute and extension at 72 °C for 1 min. Final extension was carried out at 72 °C for 10 min. The ~1500 bp PCR product was purified to remove unincorporated dNTPs and Primers before sequencing using PCR purification kit (GENEASY PCR PRODUCT PURIFICATION KIT). Both strands of the rDNA region amplified by PCR were sequenced by automated DNA sequence-3037xl DNA analyzer from Applied Biosystems using BigDye® Terminator v3.1 cycle sequencing Kit (Applied Biosystems). Sequence data were aligned and dendograms were generated using sequence analysis software version 5.2 from applied biosystems.

The sequences obtained for plus and minus strands were aligned using the appropriate software before performing bioinformatics. Sequences were compared to the non-redundant NCBI database using BLASTN, with the default settings used to find sequences closest to each other. The Expected value ande values were noted for the most similar sequences. Ten similar neighbours were aligned using CLUSTAL W2. The multiple alignment files thus obtained was then used to create a phylogram using the MEGAS software.

RESULTS
The yield of chloroform and ethyl acetate fractions were found to be 0.6 gm/l and 0.7 gm/l for CDB-1 and 0.8 gm/l and 0.7 gm/l for CDB-2 respectively. Preliminary phytochemical investigations of CDB-1 and CDB-2 fractions showed the presence of alkaloids, steroids, triterpenoids and flavonoids as important constituents (table 1).

Table 1: Qualitative phytochemical analysis of endophytic fractions of Callyspongia diffusa

| S. No. | Endophytic fractions | Carbohydrates | Flavonoids | Alkaloids | Steroids | Tannins | Triterpenoids |
|--------|----------------------|---------------|------------|-----------|----------|---------|--------------|
| 1      | CDB-1 (Chloroform)   | +             | +          | +         | +        | _       | _            |
| 2      | CDB-2 (Chloroform)   | +             | _          | +         | +        | _       | _            |
| 3      | CDB-1 (Ethylacetate) | +             | +          | +         | +        | _       | +            |
| 4      | CDB-2 (Ethylacetate) | +             | _          | +         | _        | _       | _            |

Endophytic fractions: (+) indicates the presence of the component, (-) indicates complete absence of the component.

Table 2: Efficacy of different endophytic fractions of Callyspongia diffusa against pathogenic bacteria and fungi

| S. No. | Test organism | CDB-1(MIC-µg/ml) | CDB-2(MIC-µg/ml) |
|--------|---------------|------------------|------------------|
|        |               | C | E | C | E |
| 1      | Bacillus subtilis | 0.2 | 100 | 3.12 | 100 |
| 2      | Escherichia coli | >100 | 100 | 100 | 0.4 |
| 3      | Enterococcus fecalis | 1.6 | 100 | 0.4 | 0.4 |
| 4      | Klebsiella Pnumoniae | >100 | 50 | 0.2 | 0.2 |
| 5      | Candida albicans | 0.2 | 0.4 | 3.2 | 0.8 |
| 6      | Aspergillus niger | 0.4 | 6.25 | 0.2 | 0.2 |

C-Chloroform fraction, E-Ethyl acetate fraction

The results of antimicrobial activities (MIC) of the compounds against the selected two Gram-positive, two Gram-negative bacteria and two fungi are shown in table (2). The activity of ciprofloxacin and norfloxacin are used for comparison. All the compounds showed moderate to significant microbial inhibition. Chloroform fractions of CDB-1 and CDB-2 have shown MIC of 0.2-3.12 µg/ml against Bacillus subtilis respectively, whereas MIC of ethyl acetate fractions was more than 100 µg/ml. The MIC of chloroform and ethyl acetate fractions of CDB-2 against Escherichia coli was found to be 0.4 µg/ml respectively. The lowest MIC against Enterococcus fecalis among the fraction tested was 0.4 µg/ml for CDB-2 followed by 1.6 µg/ml for chloroform fraction of CDB-1. Chloroform and ethyl acetate fraction of CDB-2 has shown MIC of 0.2 µg/ml against Klebsiella pneumoniae. A significant antifungal activity was observed against Candida albicans (MIC range 0.2-3.2 µg/ml) and Aspergillus niger (MIC range 0.2-6.25 µg/ml).

In antitubercular assay against Mycobacterium tuberculosis H37RV strain, chloroform fraction of CDB-1 and ethyl acetate fraction of CDB-2 has shown MIC of 6.25 µg/ml respectively. However, the MIC for ethyl acetate fractions of CDB-1 and chloroform fraction of CDB-2 was found to be 12.5 µg/ml. The results are shown in (table 3).

Table 3: Antitubercular activity of different endophytic fractions of Callyspongia diffusa against Mycobacterium tuberculosis H37 sensitive strain

| Endophytic fractions | CDB-1(MIC-µg/ml) | CDB-2(MIC-µg/ml) |
|----------------------|------------------|------------------|
|                      | C | E | C | E |
| CDB-1                | 6.25 | 12.5 |
| CDB-2                | 12.5 | 6.25 |

C-Chloroform fraction, E-Ethyl acetate fraction

The activity of ciprofloxacin and norfloxacin are used for comparison. All the compounds showed moderate to significant microbial inhibition. Chloroform fractions of CDB-1 and CDB-2 have shown MIC of 0.2-3.12 µg/ml against Bacillus subtilis respectively, whereas MIC of ethyl acetate fractions was more than 100 µg/ml. The MIC of chloroform and ethyl acetate fractions of CDB-2 against Escherichia coli was found to be 0.4 µg/ml respectively. The lowest MIC against Enterococcus fecalis among the fraction tested was 0.4 µg/ml for CDB-2 followed by 1.6 µg/ml for chloroform fraction of CDB-1. Chloroform and ethyl acetate fraction of CDB-2 has shown MIC of 0.2 µg/ml against Klebsiella pneumoniae. A significant antifungal activity was observed against Candida albicans (MIC range 0.2-3.2 µg/ml) and Aspergillus niger (MIC range 0.2-6.25 µg/ml).

In antitubercular assay against Mycobacterium tuberculosis H37RV strain, chloroform fraction of CDB-1 and ethyl acetate fraction of CDB-2 has shown MIC of 6.25 µg/ml respectively. However, the MIC for ethyl acetate fractions of CDB-1 and chloroform fraction of CDB-2 was found to be 12.5 µg/ml. The results are shown in (table 3).

Results of haemolysis assay suggested that ethyl acetate fractions of CDB-land CDB-2 caused least haemolysis erythrocytes as compared to chloramphenicol (fig. 1).
In vivo studies demonstrated the survival data on day 10th of post-treatment of CDB-1 endophytic fractions. The results showed about 40% of animals treated with chloroform fraction and 60% of animals treated with ethyl acetate fraction of CDB-1 were survived respectively. All the control animals died within 6 d. The results are shown in (table 4).

**Table 4: Efficacy of endophytic fractions of* Callyspongia diffusa* on* K. Pneumonia* infection in mice**

| Test compounds | No. of days/percentage of protection |
|----------------|-------------------------------------|
| Control        | 100/100/100/85/75/25/NS/NS/NS/NS/NS |
| CDB-1C         | 100/100/100/100/100/85/65/45/45/45 |
| CDB-1E         | 100/100/100/100/100/100/65/65/65/65 |

Swiss albino mice (n=30) were challenged with 5x10^5efuof*K. Pnumoniae. The animals were treated with CDB-1 endophytic fractions (100 mg/kg body wt) daily for 7 d., (CDB-1C-chloroform fraction; CDB-1 E-Ethyl acetate fraction)

The sequence of the 16s gene from VBW-1 and that of matching sequences from 10 nucleotide sequences were aligned by using the Maximum Likelihood method based on the Hasegawa-Kishino-Yano model (fig. 2).

**Fig. 2: PCR amplification of the 16s rRNA gene of CDB-1 and CDB-2**

The tree with the highest log likelihood (-2346.2209) is shown. Initial tree(s) for the heuristic search were obtained automatically as follows. When the number of common sites was<100 or less than one-fourth of the total number of sites, the maximum parsimony method was used; otherwise, BIONJ method with MCL distance matrix was used. The analysis involved 11 nucleotide sequences. Codon positions included were 1st+2nd+3rd+Noncoding. All positions containing gaps and missing data were eliminated. There was a total of 1293 positions in the final dataset. Evolutionary analyses were conducted in MEGAS5. The sequences of CDB-1 and CDB-2 were compared with existing sequences in the NCBI database using the Blast N programme (fig. 3 and 4).

A high score and low expected value indicates the greatest homology. Based on these results CDB-1 and CDB-2 were identified as *Pseudomonas taiwanensis strain BCRC 17751and Lysinibacillus sphaericus strain DSM 28* respectively. The microscopic view of the identified organisms is shown in (fig. 5).
**Fig. 3: Pylogenetic tree of CDB-1**

- Pseudomonas montelii strain NBRC 103158 (NR 114224.1)
- Pseudomonas montelii strain CIP 104885 (NR 112073.1)
- Pseudomonas montelii strain CIP 104883 (NR 024910.1)
- Pseudomonas plecoglossicida strain FPC051 (NR 024692.1)
- Pseudomonas plecoglossicida strain NBRC 103162 (NR 114226.1)
- Pseudomonas taiwanensis strain SCRC 17751 (NR 116172.1)
- Pseudomonas helianthidens strain OHA11 (NR 126220.1)
- Pseudomonas mosselli strain CLI 90-43 (NR 024904.1)
- Pseudomonas entomophila strain L14 (NR 102854.1)
- Pseudomonas entomophila strain L48 (NR 115336.1)
- Sample CDB1

**Fig. 4: Pylogenetic tree of CDB-2**

- Lysinibacillus tabaciibii strain K3514 (NR 132691.1)
- Lysinibacillus varians strain GY32 (NR 134074.1)
- Lysinibacillus mangiferae strain M-GX18 (NR 116146.1)
- Lysinibacillus sphaericus strain ATCC 14577 (NR 115724.1)
- Lysinibacillus sphaericus strain DSM 28 (NR 042073.1)
- Lysinibacillus sphaericus strain NRBC 15095 (NR 112627.1)
- Lysinibacillus sphaericus strain IAM 13420 (NR 115529.1)
- Lysinibacillus parabacilli strain NBRC 103144 (NR 114213.1)
- Lysinibacillus fusiformis strain NBRC15717 (NR 112560.1)
- Lysinibacillus boronitolerans strain NBRC 103108 (NR 114207.1)
- Sample CDB2

**Fig. 5: Microscopic view of endophytic bacterial strains**

Gram-stain micrograph of *Pseudomonas* colonies. Courtesy

Gram-stain micrograph of *Bacillus* colonies. Courtesy

Gram-staining picture of colonies from agar slant VBW-1 (CDB1)

Gram-staining picture of colonies from agar slant VBW-2 (CDB2)
DISCUSSION

In the current investigation, we have isolated and identified two endophytic bacteria (CDB-1 and CDB-2) from Callyspongia diffusa. Through phylogenetic/sequential analysis the organisms were identified as Pseudomonas taiwanensis (CDB-1) and Lysinibacillus sphaericus (CDB-2) strains of NBRC respectively. Preliminary phytochemical analysis of CDB-1 and CDB-2 nonpolar fractions showed the presence of carbohydrates, alkaloids, steroids, triterpenoids and flavonoids as main important constituents. Many reports suggested the marine sponge also is also a rich source of bioactive metabolites and showed the various activities. Some of the activities have been reported. i.e, Antifungal and antimicrobial activities have been reported in the tetracyclic furanoderpenes isolated from sponge *S. officinalis*, Spongia-13(16), 14-dien-19-ol acid, sponge-13(16)-14-dien-19-ol [27] and sponge-13(16)-14-diene [28]. Sponge *Luffarella variabilis* contain sesquiterpenoid antibiotics, manoolide which contains alpha, beta-unsaturated gama-lactone function had anti-inflammatory activity and was found to be an inhibitor of phospholipase 2A [29]. Tetracyclic sesquiterpenes have been isolated from *cacospongiacarcissis* namely phyllopholactone and phyllopholactone exhibited cytotoxic activity [30]. Several peptide alkaloids and proteins have been isolated from a marine sponge, purealin, a novel enzyme activator from the okinawan marine sponge, *Psammamypsilapsapurea* modulates enzymatic reaction of ATPases [31]. Eight endophytic and two epiphytic fungi were isolated from *Leucosolenia* sp. sponge with good antimicrobial activity [32]. Diverse ranges of bioactive and potentially novel bacterial endophytes were isolated from the marine sponges *Subertescomnus* and *Leucosolenia* sp. It was observed that *S. carnosus* isolates may be a better source of antifungal compounds, while *Leucosolenia* sp. appear to be a better source of antifungal compounds [33]. Ten marine sponge species from Rameshwara region of southern part of India were studied for fungal diversity and their biological effects. Genera such as *Acremonium*, *Alternaria*, *Aspergillus*, *Cladosporium*, *Fusarium* and *Penicillium* were frequently isolated. Species of *Aspergillus* were dominant and co-dominant in all the sponges screened.

The fungal isolates produced antifungal, antioxidant, antibiotic, anti-oxidant metabolites. A few fungi produced acetylcholinesterase inhibitors [34]. *Callyspongia diffusa* is a member of a marine sponge, reported to possess antioxidant [17] and antimicrobial activities [18, 19]. Metabolites were isolated from *Virgibacillussp* associated with *Callyspongia diffusa* reported the antimicrobial and antioxidant activity. The bacterial strain (CDB 1-CD10) of *Callyspongia diffusa* showed antimicrobial and immunomodulatory activity [17]. Isolation of antimicrobial agents from bioactive marine species (*Porpita porpita*, *Pseudonocardia* sp. and *V. cholerae*) were associated in the sponge *Callyspongia diffusa*, *S. officinalis* and *S. carnosus*. Metabolites from *Callyspongia diffusa* exhibited the lowest MIC against *Klebsiella pneumoniae*, *Escherichia coli* and *Enterococcus faecalis* were found to be 0.2 µg/ml and 0.4 µg/ml respectively for CDB-2. A significant antifungal activity was observed against *Candida albicans* (0.2-0.8 µg/ml) and *Aspergillus niger* (0.2-0.4 µg/ml). Further, Chloroform Fraction of CDB-1 and ethyl acetate fraction of CDB-2 have shown significant anti-tubercular activity against the tested organism with MIC of 6.25µg/ml. This was supported by *in vivo* antimicrobial activity against *K. Pneumoniae* infection in mice and least hemolytic activity against erythrocytes was observed. The mode of action of these fractions may be due to disruption of ETC (Electron Transport Chain) and cell membrane oxidative phosphorylation, inhibition of enzyme activity, and impairment in nutrient uptake which directly leads to lyses of bacterial cells [35]. Drug resistance of *mycobacterium tuberculosis* has increased in existing chemotherapy regimens [36]. There is a huge need in finding novel classes of natural products that may be efficient against resistant strains of microbes, especially from Bioactive endophytes from marine source due to their rich source of secondary metabolites and least toxic effects [35].

Thus, for the biotechnological production of natural compounds endophyte seems to be the most attention-grabbing alternative and they are likely to play an important role in the years to come.

CONCLUSION

In this study, we have reported the marine natural species offers a rich source of bioactive metabolites that can exploit to develop novel, useful and potential therapeutic agents. In this study we have isolated two bacterial endophytes (CDB-1 and CDB-2) from marine sponge *Callyspongia diffusa*, these offers a novel source of bioactive compounds. Chloroform and ethyl acetate fractions of CDB-1 and CDB-2 showed significant antimicrobial action against pathogenic bacteria, fungi and mycobacterium tuberculosis H7Rv stain.

**In vitro** anti-TB activity of chloroform fraction of CDB-1 and ethyl acetate fraction of CDB-2 has shown more significant (MIC of 6.25 µg/ml) respectively. However, the MIC for ethyl acetate fractions of CDB-1 and chloroform fraction of CDB-2 has shown moderately significant (MIC of 12.5 µg/ml). This was further supported by *in vivo* antimicrobial activity against *K. pneumoniae* infection in mice and least haemolytic activity against erythrocytes was observed. Thus, we can conclude that the antimicrobial activity of chloroform and ethyl acetate fractions of CDB-1 and CDB-2 may be attributed due to the presence of carbohydrates, alkaloids, steroids, triterpenoids and flavonoids as major phytoconstituents.

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AUTHOR CONTRIBUTION

I Mr. Vijayanand B Warad done the Complete research work and manuscript preparation was done by me. Prasanna Hubu was the guide of my PhD work, and he guided me in throughout the work and manuscript preparation, and third author Rajesh Shastri was the Co-guide of my PhD work, and he helped me a lot in the isolation of endophytic fractions and also guided me in throughout the work and manuscript preparation.

CONFLICT OF INTERESTS

Declared none

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