

In vitro bioassay of antibacterial and antifungal activity studies of actinomycetes from soda lakes of Ethiopia

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Abstract

The ever-increasing spectrum of microbial disease and emergence of life-threatening antibiotic-resistant pathogens necessitates more intensive search for microbial antagonistic agents from diverse symbionts and habitats. Based on these hypotheses the current study was initiated to investigate in vitro antibacterial and antifungal activities of both crude and pure compounds isolated from potential actinomycete isolates recovered from soda lakes of Ethiopia. Two soda lakes, Arenguade/Hora Hadho/ and Chitu were included in this study. A total of fifteen 12 water samples were collected and actinomycete cultures were isolated by serial dilution plating technique on various media. Morphological and biochemical characterization of selected isolates was done as described in the International Streptomyces Project Guidelines. Minimum inhibitory concentrations of the crude extract was determined for fungus using the protocol described in National Committee for Clinical Laboratory Standards and for the bacteria disc diffusion assay method as described in the Clinical and Laboratory Standards Institution. Ethanol extracts obtained from Solid State Fermentation a total of 36 isolates of actinomycetes were subjected to primary screening and tested for activity against the bacterial and fungal reference strain. Accordingly, the extracts from 10 isolates that were most active in the primary screen were 6 (16.66%) from Lake Chitu that were inhibited the growth of Gram-positive bacteria and fungal strain; 4 (11.11%) from Lake Arenguade that were inhibited Gram-negative bacteria and fungal. From evaluated the antibacterial and antifungal potential of two soda lakes water-derived actinomycetes; the best ones in terms of activity spectra were actinomycetes species isolated from Lake Chitu. Antibacterial and antifungal metabolite production was detected from isolates of Lake Chitu with maximum zone of inhibition within 5–6 days of incubation against *Candida albicans* ATCC 62376, *Aspergillus niger*, *C. neoformans*, *F. oxysporum* f. sp. *melonis* (CECT 20474), and *Shigella boydii* (clinical). All test microorganisms, except *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical isolate) were sensitive to the culture filtrate starting from 8–12 days of incubation. In terms of the effectiveness of the antibacterial activity, the antibacterial metabolite produced from different carbon sources were more effective on *S. aureus* (clinical isolate), *A. niger*, *C. neoformans*, *S. aureus* ATCC25923, *Fusarium oxysporum* f. sp. *melonis* (CECT 20474), *Shigella boydii* (clinical isolate) respectively. However, the most resistant test isolates with least activity was detected against *Candida albicans* ATCC 62376 and *E. coli* ATCC 25922.

Background Of The Study

The emergence of multi-drug resistant pathogens necessitates continuous search for novel antimicrobial agents. There is the continuous need for novel antibiotics to overcome the serious problem of evolving pathogens, naturally resistant bacteria and fungi, and multidrug resistance among common bacterial pathogens. Also, searching for novel strains from pristine environments is an important approach for obtaining novel bioactive molecules. Ironically, despite the extensive research done on antibiotic-producing *Streptomyces*, interactions of actinomycetes with other bacteria, the ecological rationale and physiological property of the *Streptomyces* the extent to which produce antibiotics is the most striking and poorly understood [1]. Among the microbes, the secondary metabolites recovered from the soda lakes

and untouched forest soil actinomycetes were more added values [2- 6]. New resistant strains emerge more quickly while the rate of discovery of new antibiotics is slowing down. Today, superbugs are estimated to cause 700,000 deaths every year and are expected to rise to 10 million deaths each year by 2050 [7]. This necessitates screening for new and novel drugs producing microorganisms, including actinomycetes [8&9].

In this regard, Ethiopian soda lakes and untouched soil habitats from various climatic zones possessing remarkable biodiversity may harbor new actinomycete strains that have the potential for new antibiotic discovery. Previous studies by a few investigators confirmed the existence of potential antibiotic-producing actinomycetes from different ecosystems of Ethiopia. Accordingly, alkaliphilic actinomycetes capable of producing antimicrobial metabolites active against bacteria and fungi were isolated from Ethiopian soda lakes [10-12], rhizosphere of different plants, farmlands and garden soils [13 &14]. Most microorganisms are still untapped in terms of their capacity to produce secondary metabolites since only a small fraction can be cultured in the laboratory. Thus, improving cultivation techniques to extend the range of secondary metabolite producers accessible under laboratory conditions is an important first step in prospecting underexplored sources for the isolation of novel antibiotics. For industrial application, further, purification, structural elucidation, characterization, and optimization of the culture condition are required [15].

Material And Methods

3.1. DESCRIPTION OF THE STUDY AREA

Two soda lakes, namely, Arenguade/Hora Hadho/ and Chitu were included respectively in this study. As a result of their sensitivity to ecological disturbances, soda lakes are among the ecological systems for which continuous scientific investigations on their microbial diversity should be given high priority [16]. Lake Chitu is located at a distance of about 180 Km South of Addis Ababa at an altitude of 1600 m above sea level [17]. Likewise, Lake Arenguade is located about 50 Km Southeast of Addis Ababa. For this study, Lake Chitu and Lake Arenguade were thus chosen for their peculiar and interesting features to understand the microbial ecology of the soda lakes in the region. These lakes are similar in that both are crater in origin, have closed basins, have very small catchment areas (limited external material input) ,and are highly productive. On the other hand, they have several differences. Lake Chitu is shallow (max. depth 17 m), located inside the rift valley in semiarid lowland area (1600 m.a.s.l.) and is highly alkaline (>600 meq/L) and saline (>6%), while Lake Arenguade is deeper (max. depth 32 m), located outside the rift valley in the highland area around Bishoftu (2000 m.a.s.l) and is less saline (<0.3%) and alkaline (about 50 meq/L). These features are good bases for a comparative microbial community study across sharp environmental gradients [17].

2.2. Water sample collection

From each lakes, a total of 12 water samples were collected in 500 mL sterile screw-capped bottles and sufficient space was provided for aeration and thorough mixing as described in [18]. All samples were labeled and coded as LCHAACC13 (Lake Chitu Alkaliphilic Actinomycetes Culture Collection), LAAACC15 (Lake Arenguade/Hora Hado/ Alkaliphilic Actinomycetes Culture Collection), followed by a specific number to indicate isolate number. Samples were transported to Adama Science and Technology University, Applied Microbiology Laboratory and stored in the refrigerator at 4 °C until analyzed.

2.3. Isolation and screening of alkaliphilic actinomycetes

2.3.1. Media for the cultivation of actinomycetes

For the cultivation of actinomycete, both nutrient limiting and nutrient-rich media were used. The isolation media for the two soda lakes were chosen based on the culturability of general actinobacterial and the geochemical properties of the soda lakes [19]. Actinomycete cultures were isolated by serial dilution plating technique on various media including starch casein agar (SCA) composed of (g/l) soluble starch, 10; Casein ,0.3; KNO₃, 2; NaCl, 2; K₂HPO₄, 2; MgSO₄.7H₂O, 0.05; CaCO₃,0.02; FeSO₄.7H₂O, 0.01; agar, 18; and Na₂CO₃, 10; and distilled water, 1,000 ml. starch nutrient agar, and Humic Acid agar composed of (g/l) humic acid, 1; Na₂HPO₄,0.5 ; KCl ,1.71 ; MgSO₄.7H₂O, 0.05; FeSO₄.7H₂O, 0.01; CaCO₃, 0.02; yeast extract,1 and ; agar, 18. Na₂CO₃ was sterilized (at 121 °C, 15 psi) separately as a 25% solution and added to the rest of the medium after cooling. To inhibit Gram-negative bacterial contamination 1 ml nalidixic acid was added to the medium from a stock solution prepared by dissolving 0.01g of nalidixic acid in 10 ml of 100% methanol by swirling and mixing in a 37 °C water bath [19].

2.3.3. Morphological and biochemical characterization of selected isolates

Identification of Actinomycetes to the genus level was done by a polyphasic approach, including cultural, morphological, physiological, and biochemical characteristics as described in the International Streptomyces Project Guidelines. The isolation media for the five sites were chosen based on the culturability of general actinobacterial and the geochemical properties of the soda lakes and forest soils. The four media Starch Casein Agar, Starch Casein Broth, Actinomycetes Isolation Agar, and Glycerol Asparagine Agar media were used in combination with temperature (28 °C- 55 °C), salt (1 – 10 % w/v) and pH (7.0 -13) ranges to simulate the sampling environment. Glycerol Asparagine Agar media was designed for the isolation of actinomycetes from Chitu and Arenguade soda lakes, whilst the rests are general actinobacterial media.

3.3.5. Bacterial and Fungal reference strain

Staphylococcus aureus ATCC25923, *S. aureus* (clinical isolate) *E. coli* ATCC 25922, *Shigella boydii* (clinical isolate), and *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Aspergillus niger*, *Candida albicans* ATCC 62376 and *C. neoformans* (clinical isolates) were used as bacterial and fungal reference (test) strains. The test bacterial and fungal strains were obtained from Ethiopian Health and Nutrition Research Institute (EHNRI) and Biomedical Laboratory, Faculty of Life Sciences, AAU.

3.4. Media for test microorganisms

3.4.1. Nutrient agar and Nutrient broth

For the antimicrobial susceptibility test and screening, those test bacteria were seeded on nutrient agar media. Twenty-eight grams of nutrient agar (NA) (based on manufacturer's recipe) was dissolved in 1L of distilled water and sterilized at 121 °C, 15 psi. Then about 25 ml of sterile NA media was poured on sterile Petri-dishes and allowed to cool overnight before use. Similarly, 13 g Nutrient Broth dissolved in 1L distilled water and sterilized. About 5 ml of sterile nutrient broth was added on sterile screw-capped test tubes then inoculated with test microorganisms and incubated overnight at 37 °C. The broth culture was adjusted to 0.5 McFarland standards before it was used [19].

3.4.2. Muller Hinton agar

For the antibacterial metabolite susceptibility test, thirty-eight grams of Muller Hinton agar based on the manufacturer's recipe were dissolved in 1L of distilled water and sterilized at 121 °C, 15 psi. About 25 ml was poured into sterile Petri dishes to grow the test microorganisms (bacteria).

3.5. Standardization of inoculum

Standardization of the inoculum density of isolates for susceptibility test was done by the method described in [20]. To determine the active phase of test organisms, each isolate was grown in 100 mL of nutrient broth for bacteria and potato dextrose agar for fungi in 250 mL Erlenmeyer flask on a rotary shaker at 120 r/min and 37°C. Samples were taken every 2 h and the density of the turbidity standard was determined using a spectrophotometer (JENWAY, London) at 660 nm. The optical density values were extra plotted against time to determine the different phases of the growth curve. Samples from the exponential phase were taken to adjust the inoculum density with 0.5 McFarland turbidity standard prepared by adding about 0.05 ml of 1 % BaCl₂ and 9.95 ml of 1% H₂SO₄ were mixed to give 0.5 McFarland standards by which approximately 1.5 x 10⁸ bacterial cell suspensions are expected to be present per ml of culture as cited in [8].

3.6. Primary screening of actinomycetes for antimicrobial activity

Antimicrobial activity was first screened using the conventional agar cylinders method on Potato Dextrose Agar (PDA) and Muller Hint Agar (MHA) plates for fungi and bacteria respectively [21-22]. Mycelium plugs (7 mm diameter) of active isolates incubated at a different time (3, 7, 10, and 14 days) were inoculated onto PDA and MHA plates previously inoculated with target pathogens. Secondly, a double layer method was used for confirmation. The active strain was inoculated as a spot in the center of ISP2 plates at 30 °C for 7 days. After incubation, the plates were then covered by 10 ml of PDA and MHA previously inoculated with target fungi and bacteria respectively [21]. The inhibition zones around each spot were measured (mm) after 24 h at 37 °C for bacteria, 48 h and 7 days at 30 °C for yeast and molds respectively. Two replicates were prepared for each test and plates with indicator strain were used as control. The isolates were then selected based on a wide spectrum activity against tested microorganisms for further studies [20].

3.7. Secondary screening

The bioactivity testing of the ethanol crude extracts obtained from the selected active isolates was done for antifungal and antibacterial susceptibility based on primary screening results. Minimum inhibitory concentrations of the crude extract was determined for fungus using the protocol described in NCCLS [23] and for bacteria as described in CLSI [20]. All the secondary bioactivity test with crude extracts was conducted in triplicates and inhibition zone diameters were measured.

4. Data management and analysis

The collected and recorded data were analyzed using Microsoft Excell. The data on temperature and pH tolerance was analyzed by comparing the mean growth in the log colony forming-unit through analysis of variances (ANOVAs). All statistical results with $P<0.05$ were considered to be statistically significant.

Results And Discussions

3.1. Metabolite production and optimization of selected isolates

The preliminary antimicrobial activities of the isolates against selected pathogens by the cross streak method showed that they possess antimicrobial properties, as observed from their inhibitory activities against the selected bacterial and fungal strain. From ethanol extracts obtained from Solid State Fermentation (SSF) (Primary screening), a total of 36 isolates of actinomycetes were subjected to primary screening and tested for activity against bacterial and fungal reference strain. Accordingly, the extracts

from 10 isolates that were most active in the primary screen were 6 (16.66%) from LCHAACC13 (Lake Chitu Alkaliphilic Actinomycete Culture Collection) that were inhibited the growth of Gram-positive bacteria and fungal strain; 4 (11.11%) from LAAACC15 (Lake Arengude Alkaliphilic Actinomycete Culture Collection) that were inhibited Gram-negative bacteria and fungal. From the 10 selected isolates, the best ones in terms of activity spectra were actinomycetes spp. isolated from LCHAACC13. Extracts from 10 isolates that were most active in this primary screen, namely that designated Lc-13, Lc-15, Lc-17, La-13, La-15, La-19, Lc-21, La-17, La-19, and Lc-23, were selected for further characterization. Isolates those were coded with Lc-13-23 and La-13, 15, 17, and 19 were obtained from lake Chitu (Lc) and Arengude (La) respectively (Supplementary materials, Table 3).

3.1.1. Dates of incubation period on the production of antibacterial and antifungal metabolite by isolates of two soda lakes

From evaluated the antibacterial and antifungal potential of two soda lakes water-derived actinomycetes; no antibacterial metabolite production was observed up to 4 and 7 days of incubation for LCHAACC13 (Lake Chitu Alkaliphilic Actinomycete Culture Collection) and LAAACC15 (Lake Arengude/Hora Hadho/Alkaliphilic Actinomycete Culture Collection) respectively as shown in figure 1 and 2 below. However, antibacterial and antifungal metabolite production was detected from isolates of LCHAACC13 with maximum zone of inhibition within 5 - 6 days of incubation against *Candida albicans* ATCC 62376, *Aspergillus niger*, *C. neoformans*, *Fusarium oxysporum f. sp. melonis* (CECT 20474), and *Shigella boydii* (clinical) (Fig.1). The most resistant test isolates as increasing the time of incubation were *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical isolate). All test microorganisms, except *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical isolate) were sensitive to the culture filtrate starting from 8 - 12 days of incubation and in terms of multiple inhibition and diameter of zone of inhibition, the highest level of antibacterial and antifungal production was obtained at 10th, 11th, and 12th day of incubation by LCHAACC13 and LAAACC15 respectively (Fig. 1 and 2). Some decrement in activity starting from 13 days of incubation and least activity were observed from 16 days and above.

On the other hand, no strong antibacterial and antifungal metabolite production and activities were observed from LAACC15 (Lake Arengude/Hora Hadho/Alkaliphilic Culture Collection) up to 7 days of incubation (Fig.2). However, antimicrobial activity of the culture filtrate of LAACC15 was detected on all test microorganisms except *E. coli* ATCC 25922, *S. aureus* ATCC25923, and *S. aureus* (clinical isolate) starting from the 8th, 10th, 11th, and 12th days of incubation with maximum and decrement activity at 14th and 16 days onwards respectively (Fig.2). The best antibacterial activity was found against *Shigella boydii* (12 mm) clinical isolate, followed by *A. niger* (10 mm), *C. neoformans* (clinical isolate), *F. oxysporum f. sp. melonis* (CECT 20474), and *C. albicans* ATCC 62376 (9 mm) (Fig.2).

3.3.2. Carbon sources on growth and production of antibacterial and antifungal metabolite by two soda lakes

Among the carbon sources for the production of antibacterial metabolite by LCHAACC13, high antibacterial and antifungal metabolite production was obtained media containing starch, followed by glycerol, D-xylose, D-glucose, D-fructose, sucrose, and maltose respectively only carbon sources in the submerged fermentation (Fig.3). Interestingly, both strains of soda lakes were unable to use lactose as a sole carbon source. In terms of the effectiveness of the antibacterial activity, the antibacterial metabolite produced from different carbon sources were more effective on *S. aureus* (clinical isolate), *A. niger*, *C. neoformans*, *S. aureus* ATCC25923, *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Shigella boydii* (clinical isolate) (12 mm) respectively. However, the most resistant test isolate with least activity was detected against *Candida albicans* ATCC 62376 and *E. coli* ATCC 25922.

For LAAACC15, high antibacterial and antifungal metabolite production were recorded by the presence of Glycerol, followed by starch, D-xylose respectively (Fig 4). Among tested bacterial reference strains, *S. aureus* ATCC25923, *A. niger*, *C. neoformans* and *S. aureus* (clinical isolate), *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Shigella boydii* (clinical isolate) (12 mm) were more sensitive. Similarly, like the isolate of lake Chitu, for lake Arenguade the most resistant test isolate with the least activity was detected against *Candida albicans* ATCC 62376 and *E. coli* ATCC 25922.

3.3.3. Nitrogen concentration on growth and production of antibacterial and antifungal metabolite by two soda lakes

Primary screening of antibacterial metabolite of LCHACC13 from different Nitrogen sources showed that maximum zone of inhibition against *Shigella boydii* (clinical), *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Aspergillus niger*, *C. neoformans* (clinical isolates), and *S. aureus* (clinical) in the presence of tryptone, NaNO_3 , Casein, and KNO_3 were used as nitrogen sources respectively, while yeast extract and $(\text{NH}_4)_2\text{SO}_4$ were least effective in terms of multiple inhibition of the test microorganisms (Fig.5). The most resistant test isolates were *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *Candida albicans* ATCC 62376.

The culture filtrate of LAAACC15 grown in different nitrogen sources, however showed very strong antibacterial and antifungal activity against *Shigella boydii* (clinical), *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Aspergillus niger*, *Candida albicans* ATCC 62376 and *C. neoformans* (clinical isolates) when peptone and KNO_3 were used as nitrogen sources and very weak antibacterial and antifungal activity when $(\text{NH}_4)_2\text{SO}_4$ and NH_4NO_3 as nitrogen sources (Fig.6). The most resistant test isolates were *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical).

3.3.4. Phosphate concentration on growth and production of antibacterial and antifungal metabolite by two soda lakes

From 12 days culture filtrate of LCHAACC13 grown at different phosphate concentration showed very strong antibacterial and antifungal activity was observed against *Candida albicans* ATCC 62376, *Aspergillus niger*, *Fusarium oxysporum f. sp. melonis* (CECT 20474), *Staphylococcus aureus* ATCC25923, *C. neoformans* (clinical isolates), and *Shigella boydii* (clinical), at the phosphate concentration range between 0.1 - 0.7% and maximum activity at 0.2%, 0.5%, and 0.6% , whereas when the phosphate concentration was increased to 1% and above no antibacterial and antifungal metabolite production and activities were reduced (Fig. 7). The most resistant test isolates at all phosphate concentration were *E. coli* ATCC 25922, and *S. aureus* (clinical).

Antibacterial and antifungal metabolite productions and activities tested from 14 days of culture filtrate of LAACC15 showed very strong antibacterial and antifungal activity was observed against *Shigella boydii* (clinical), *Aspergillus niger*, *Fusarium oxysporum f. sp. melonis* (CECT 20474), *S. aureus* (clinical), *C. neoformans* (clinical isolates), and *Shigella boydii* (clinical), at the phosphate concentration range 0.1 - 1% and maximum activity at 0.5% and 0.6% , whereas when the phosphate concentration was increased to 3% and above antibacterial and antifungal metabolite production and activities were reduced (Fig. 8). The most resistant test isolates at all phosphate concentration were *E. coli* ATCC 25922, *Staphylococcus aureus* ATCC25923, and *Candida albicans* ATCC 62376.

3.3.5. Salt concentration on growth and production of antibacterial and antifungal metabolite by two soda lakes

Extracted CF of LCHAACC13 showed very strong antibacterial and antifungal activity was observed against *C. neoformans* (clinical isolates), *Aspergillus niger*, *S. aureus* (clinical), *Candida albicans* ATCC 62376, and *Fusarium oxysporum f. sp. melonis* (CECT 20474), at the salt concentration range 0.1 - 5% and maximum activity at 0.5%. On the other hand, weak antibacterial and antifungal activity was detected from the crude filtrate of isolates grown on 3%, 5, and 7% NaCl concentration and no antibacterial metabolite from the isolates grown without NaCl (Fig. 9). The most resistant test isolates at all salt concentration were *E. coli* ATCC 25922, *Staphylococcus aureus* ATCC25923, and *Shigella boydii* (clinical).

On the other hand, extracted crude filtrate of LAACC15 showed very strong antibacterial and antifungal activity against *C. neoformans* (clinical isolates), *Aspergillus niger*, *S. aureus* (clinical), *Candida albicans* ATCC 62376, and *Fusarium oxysporum f. sp. melonis* (CECT 20474), at the salt concentration range 0.1 - 3% and maximum activity at 1%. Unlike the crude filtrate of lake Chitu, no antibacterial and antifungal activity was detected from the crude filtrate of the isolates grown on 3%, 5, and 7% NaCl concentration and no antibacterial metabolite from isolates grown without NaCl (Fig.10). The most resistant test isolates at all salt concentration were *E. coli* ATCC 25922, *Staphylococcus aureus* ATCC25923, and *Shigella boydii* (clinical).

3.3.6 Effect of pH on growth and antibiotic production by LCHAACC 13 and LAAACC 15

The final pH measured for both isolates grown on different cultivation media indicated that the pH remained alkaline regardless of the medium constituents and for both isolates antibacterial and antifungal metabolite production was more favored when the pH of the cultivation media adjusted between 9 - 11, with maximum activity at 10 against *C. neoformans* (clinical isolates), *Aspergillus niger*, *S. aureus* (clinical), *Candida albicans* ATCC 62376, and *Fusarium oxysporum f. sp. melonis* (CECT 20474). The most resistant test isolates at all salt concentration was *E. coli* ATCC 25922, *Staphylococcus aureus* ATCC25923, and *Shigella boydii* (clinical). As shown in figure 11 for both isolates production of antibacterial metabolite was favored when the culture condition was alkaline and both culture filtrate of LCHAACC13 and LAACC15 was stable at alkaline pH but weak stability at pH 12 and unstable at pH 13 values.

3.3.7. Effect of cultural temperature and growth period

It was observed that the crude filtrate of LCHAACC13 and LAAACC13 were stable from 40 °C - 60 °C with maximum activity at 40 °C. against *E. coli* and *S. aureus* ATCC25923.

Discussions

From the finding, we observed that growth, antibacterial, and antifungal metabolite production and activities by isolates were positively affected by seven independent variables, namely incubation period, carbon (starch), nitrogen (NaNO_3 , Casein, and KNO_3) sources, phosphate concentrations, pH, salt concentration, and temperature stability. We evaluated *in vitro* antibacterial and antifungal potential of two soda lakes water-derived actinomycetes and we observed that no strong antibacterial and antifungal metabolite production up to 4 and 7 days of incubation for lake Chitu alkaliphilic actinomycete culture collection and lake Arenguade/Hora Hadho/ alkaliphilic actinomycete culture collection) respectively. However, from isolates of LCHAACC13, we observed that with a maximum zone of inhibition within 5 - 6 days of incubation against *Candida albicans* ATCC 62376, *Aspergillus niger*, *C. neoformans*, *Fusarium oxysporum f. sp. melonis* (CECT 20474), and *Shigella boydii* (clinical) (Fig.1). The most resistant test isolates as increasing the time of incubation were *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical isolate). All test microorganisms, except *Staphylococcus aureus* ATCC25923, *E. coli* ATCC 25922, and *S. aureus* (clinical isolate) were sensitive to the culture filtrate starting from 8 - 12 days of incubation and in terms of multiple inhibition and diameter of zone of inhibition, the highest level of antibacterial and antifungal production was obtained at 10th, 11th, and 12th day of incubation by LCHAACC13 and LAAACC15 respectively.

Some decrement in activity starting from 13 days of incubation and least activity were observed from 16 days and above. Our result is in agreement with [24], they evaluated the antimicrobial activity of the strain

ActiF450 against selected *S. aureus* and observed that antimicrobial activity started after 2 days of incubation and reached a maximum after 7 days. They observed a potent and broad-spectrum activity against a range of human fungal pathogens including moulds and yeasts, such as *Arthroderma vanbreuseghemii*, *A. fumigatus*, *A. niger*, *C. albicans*, *C. glabarta*, *C. krusei*, *C. parapsilosis*, *F. oxysporum*, *F. solani*, *Microsporum canis*, *Rhodotorula mucilaginous* and *Scodapulariopsis candida*. In addition, high antibacterial activity was recorded against pathogenic *staphylococci*. [25] screened 180 actinobacteria for their novel biocontrol agents against plant pathogens for anti-fungal activity both *in vitro* and *in vivo* and found that three isolates exhibited potent anti-fungal activity against *Botrytis cinerea* and *F. oxysporum* f. sp. *cucumerinum* evaluated antifungal activity of *Streptomyces rimosus* MY02 [26] and observed the diameter of inhibition zone of the culture supernatant from *S. rimosus* MY02 against *Fusarium oxysporum* f sp. *cucumarinum* was 33.19 mm.

The results showed that among the carbon sources antibacterial and antifungal activity was higher when the isolate was grown in a medium having starch casein agar for lake Chitu isolates enhanced the production of secondary metabolites and the zone of inhibition against reference bacterial and fungal strain which is in agreement with [27-28] that starch casein medium found to be good base for antibacterial and antifungal metabolite production and D-xylose for lake Arenguade in agreement with [29] dextrose prove to be the best carbon source for antibiotic production by *S. kanamyceticus* M27 with a maximum zone of inhibition (22 mm) and 19 mm respectively. Our result is also supported by [30-33] that they concluded that the rate at which carbon source is metabolized can influence the formation of biomass or production of primary or secondary metabolites and fast growth due to high concentrations of rapidly metabolized sugars is often associated with low productivity of secondary metabolites.

Other researchers namely, [8] soybean meal proved to be a suitable carbon source for antibiotic production by *S. tanashiensis* A₂D; [34] no secondary metabolite production with supplemented medium with galactose, lactose, raffinose, and maltose as a sole carbon source; [35] readily metabolized carbon sources, such as glucose, can suppress antibiotic production by preventing the synthesis of a key enzyme in the biosynthesis pathway referred to as "catabolic repression"; [29, 35, 36] *S. griseoarneus*, *Streptomyces violatus*, and *S. sannanensis* strain RJT-1, glucose found to be suitable carbon source for the antibiotic production; while dextrose at a concentration of 2 g/100 ml gave maximal antibiotic titers and a higher dose decreased the yield by *S. kanamyceticus*[29; 34] high concentration of glucose as a repressor of secondary metabolism and maximum cell growth rates can inhibit antimicrobial agent production[33]. An example of catabolic repression of secondary metabolism in actinomycetes is that of actinomycin synthesis by *S. antibioticus* after more glucose is added to the media [37] and in *S. aureofaciens*, the production of tetracycline was critically influenced when 10% soluble starch was enriched with sweet potato residues [38].

Among the nitrogen sources used for the isolate of LCHAACC13 (Fig. 6), yeast extract was suitable in higher production of antibacterial metabolites that were active against *E. coli* and *S. aureus* followed by NaNO₃, casein, and KNO₃ that were effective against most of the test bacteria (Table.10). The culture

filtrate of LAAACC15 grown in different nitrogen sources was showed that very weak antibacterial activity against *Shigella boydii* as compared to the antibacterial activity obtained from the usual cultivation media starch casein agar (SCA) which contains a combination of Casein and KNO_3 as nitrogen sources (Fig. 6). This might because yeast extract is rich in vitamins, minerals, amino acids, digested nucleic acids, and other growth factors which could induce antibacterial metabolite synthesis. The results are comparable with [36] as they observed yeast extract gave the highest antimicrobial metabolite production followed by NaNO_3 , peptone, and KNO_3 by new *Streptomyces* species; [29] in which they tested different nitrogen source contained the basal mineral salts plus 2% dextrose and amino acids and inorganic nitrogen compounds employed at a concentration equivalent to 84 mg of nitrogen per 100 ml. They obtained the highest antibiotic in a synthetic medium containing (NH_4) H_2PO_4 as the nitrogen sources and the optimal concentration for antibiotic production at a concentration of 0.68%.

Additionally,[18], in the production of polyene antibiotics, soybean meal has been considered as a good nitrogen source because of the balance of nutrients, low phosphorous content, and slow hydrolysis and concluded that the presence of nitrogen regulation revealed on the enzymatic level in the fermentation of cephalexin and patulin [39] observed that higher growth and antibacterial activity in ammonium nitrate as nitrogen source followed by sodium nitrate and potassium nitrate exhibited maximum growth and antibacterial activity [40] investigated the effects of nitrogen sources on streptolydigin production and distribution of secondary metabolites from *Streptomyces lydicus*. They found that when soybean meal was used as the source of nitrogen, three analogs of streptolydigin were detected, and among the nitrogen sources glutamic acid was most favorable for the formation of streptolydigin.

Antibacterial and antifungal metabolite production were tested from 12 days culture filtrate of LCHAACC13 grown at different phosphate concentration and the result showed that detectable activity was observed at the phosphate concentration range between 0.1 - 0.7% and maximum activity at 0.2%, whereas when the phosphate concentration was increased to 1% and above no antibacterial metabolite production (Fig. 8 and 9). It has been reported that phosphate concentrations above 0.005% suppress production of natamycin by *Streptomyces natalensis* [41; 42] that KH_2PO_4 is not favorable for the production of antibiotic by *S. violatus*, however, K_2HPO_4 at a concentration of 1g/l yields an inhibition zone of 22 mm, equivalent to an antibiotic concentration of 128 $\mu\text{g}/\text{ml}$. Elucidation of the precise interaction between phosphate metabolism and antibiotic biosynthesis has been complicated by the fact that phosphate is involved in a host of related and non-related metabolic pathways [43].

In addition, [44] also found that when adding a large amount of inorganic phosphate, consumption of carbon and nitrogen sources and respiration are accelerated resulting in good growth, but the production of antibiotics is usually reduced. However, it is presumed that when the concentration of inorganic phosphate in the culture is high, the intracellular concentration of ATP is increased and the primary metabolism is accelerated. When the amount of inorganic phosphate is lowered, the ATP concentration decreases; this decrease depresses metabolic conversions which are required for the production of antibiotics [43]. In general, three separate regulatory roles, viz. independently inhibition of enzyme action, regulation of protein expression, and limit on the cell growth have been postulated for phosphate [43].

For both isolates, antibacterial and antifungal metabolite production was more favored when the pH of the cultivation media adjusted between 9 - 11, with maximum activity at 11 for lake Chitu and 10 Arenguade respectively. The results are comparable with alkaliphilic actinomycetes *Nocardiopsis alkaliphila* sp. nov. [45, 46] grow optimally at pH 9.5 - 10. *N. metallicus* [47] exhibited growth in the pH range 7-10.5, whereas in *Bogoriella caseilytica* [48], a new alkaliphilic actinomycete, optimum growth occurred at pH values 9-10. The results are comparable with some *Streptomyces* species recorded to produce antibiotics against bacteria, fungi, and yeast at alkaline pH [8]. The results are in contrast to the result reported using *Streptomyces* sp. for antimicrobial production [49]. Our finding supports the fact that generally, alkaline environment is more suitable for the growth of *Streptomyces* and thus production of antimicrobial compound [50; 51] reported that when the pH was maintained at 5, production of geldanamycin was increased from 414 to 768 mg/L. [51] studied the effects of the temperature, pH, incubation period, for the production of antimicrobial metabolite production and they found temperature of 35 °C and pH 8 were the best for growth and antimicrobial agent production and 14 to 15 days of incubation was found to be the best for maximum growth and antimicrobial activity, respectively.

An Increase in the concentration of sodium chloride in the medium influenced the growth and antimicrobial compound production. Extracted CF of LCHAACC13 showed that maximum antibacterial metabolite production at 1% salt concentration against *Salmonella typhii* and *Streptococcus Pneumonia*, whereas, *Escherichia coli* and *Pseudomonas aeruginosa* as shown in Figure 10 below. On the other hand, weak antibacterial and antifungal activity was detected from the crude filtrate of both isolates grown on 5%, 7%, and 9% NaCl concentration and no antibacterial metabolite from both isolates grown without NaCl. [18] reported the antimicrobial potential of actinomycetes to isolate that grew in the presence of 20% (w/v) NaCl, while antibiotic production being maximum with 5% (w/v) NaCl in the production medium. [52] reported that good growth as well as an antimicrobial activity when salt concentrations of 12 g/L and a decline in growth and antimicrobial activity after 20 g/L. In *Streptomyces fradiae* NCIM 2418 [53], neomycin production was enhanced by sodium nitrate and ammonium chloride, copper sulphate, and zinc sulphate positively influenced on neomycin production. In *Streptomyces rimosus* [54], supplementation of 1% (w/w) CaCO₃, enhance the production of tracycline in solid state fermentation when corncob was used as the solid medium.

Usually, cultivation for antibiotic production is performed under one constant temperature from the beginning to the end, but the temperature adequate for growth is not always the same as that for production. It is known that thermophilic actinomycetes such as thermo actinomycetes produce new antibiotics at a temperature higher than 40 °C, but *Streptomyces* usually produces antibiotics at temperatures near 27 °C. Generally, the range of a temperature supporting good growth is as wide as 25 °C, but the temperature range adequate for good production of secondary metabolites is narrow and 5 ~ 25 °C [35]. *S. violates* showed a narrow range of incubation temperature for relatively good growth and antibiotic production. The increase of the incubation temperature from 20 °C to 30 °C, increased the growth of the cells and the production of the antibiotic by 3.45-fold and 4.39-fold, respectively. Maximum antibiotic production was obtained at 30 °C. A higher incubation temperature (35 °C) harmed growth and

antibiotic production. The size of inoculum affected the ability of *S. violaceus* to produce the antibiotic in the tested cultures. An increase of the inoculum size from 0.5 ml to 2 ml /50 ml medium enhanced the production of the antibiotic by approximately 4.6 - fold. Thus temperature must be considered separately growth and for production. It would be of interest in antibiotic screening to use temperature shifts [41].

Conclusion

The present study provides evidence that Ethiopian Soda lakes are a rich and valuable resource of potentially active actinomycetes. Concisely, this study highlighted for the first time the antimicrobial potentials of soda lakes associated actinobacteria with the ability to suppress major human fungal and bacterial pathogens *in vitro*. Further studies are needed to enhance the isolation and selection of actinomycetes from unexplored ecosystems for antibiotic discovery. Hopefully, these new agents will meet the challenges as we attempt to manage serious underlying infectious diseases. In addition, knowledge of the actinobacteria gene clusters may provide important answers toward understanding the metabolites biosynthetic pathway.

Declarations

Disclosure of interest

The authors declare no conflicts of interest regarding the publication of this study.

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Tables

Table 1. Preliminary antimicrobial screening of 10 active isolates against different microbial pathogens.

Pathogens	Active isolates									
	Lc-13	Lc-15	Lc-17	La-13	La-15	Lc-19	Lc-21	La-17	La-19	Lc-23
<i>S. aureus</i> ATCC25923	++	++	++	++	++	++	++	++	++	++
<i>S. aureus</i> (clinical)	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>E. coli</i> ATCC 25922	++	++	++	++	++	++	++	++	++	++
<i>Sh. boydii</i> (clinical)	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>F. oxysporum</i> (CECT 20474)	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>Aspergillus niger</i>	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>C. albicans</i> ATCC 62376	++	++	++	++	++	++	++	++	++	++
<i>C. neoformans</i> (clinical)	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++

+++: Good activity; ++: Moderate activity; +: Weak activity; -: No activity.

Table.2. Secondary antimicrobial screening of 10 active isolates against different microbial pathogens.

Pathogens		Active isolates (zone of inhibition in mm)										
		C	Lc-13	Lc-15	Lc-17	La-13	La-15	Lc-19	Lc-21	La-17	La-19	Lc-23
<i>S. aureus</i> ATCC25923	14	10	++	++	++	++	++	++	++	++	++	++
<i>S. aureus</i> (clinical)	14	12	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>E. coli</i> ATCC 25922		++	++	++	++	++	++	++	++	++	++	++
<i>Sh. boydii</i> (clinical)		+++	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>F. oxysporum</i> (CECT 20474)		+++	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>Aspergillus niger</i>	14	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++
<i>C. albicans</i> ATCC 62376	13	++	++	++	++	++	++	++	++	++	++	++
<i>C. neoformans</i> (clinical)		+++	+++	+++	+++	+++	+++	+++	+++	+++	+++	+++

C-Control (Streptomycin for bacteria; ketoconazole for fungi)

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Figures

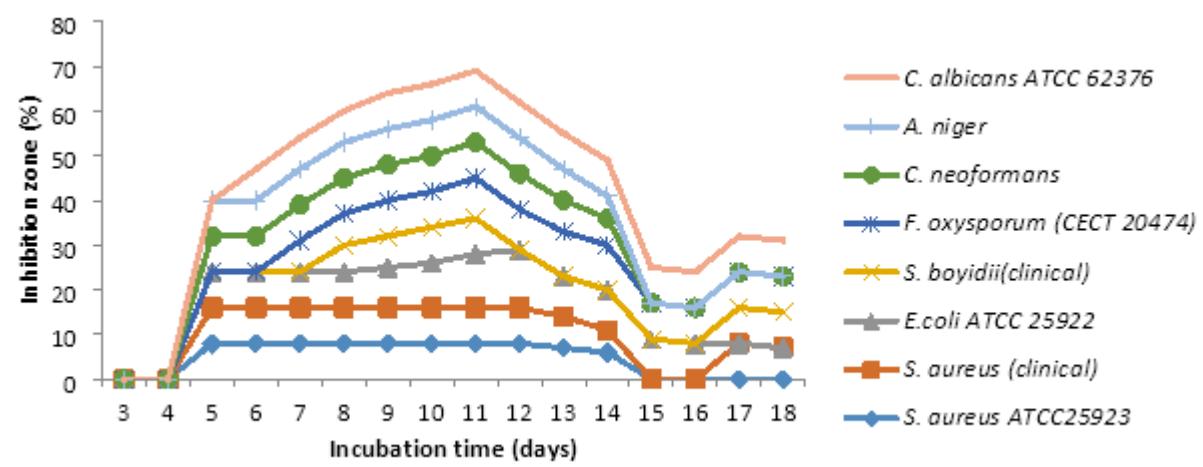


Figure 1

Zone of inhibition (measurement includes the diameter of the disc 6mm) of an antibacterial antifungal metabolite of Lc-13-23 taken at different incubation periods (day).

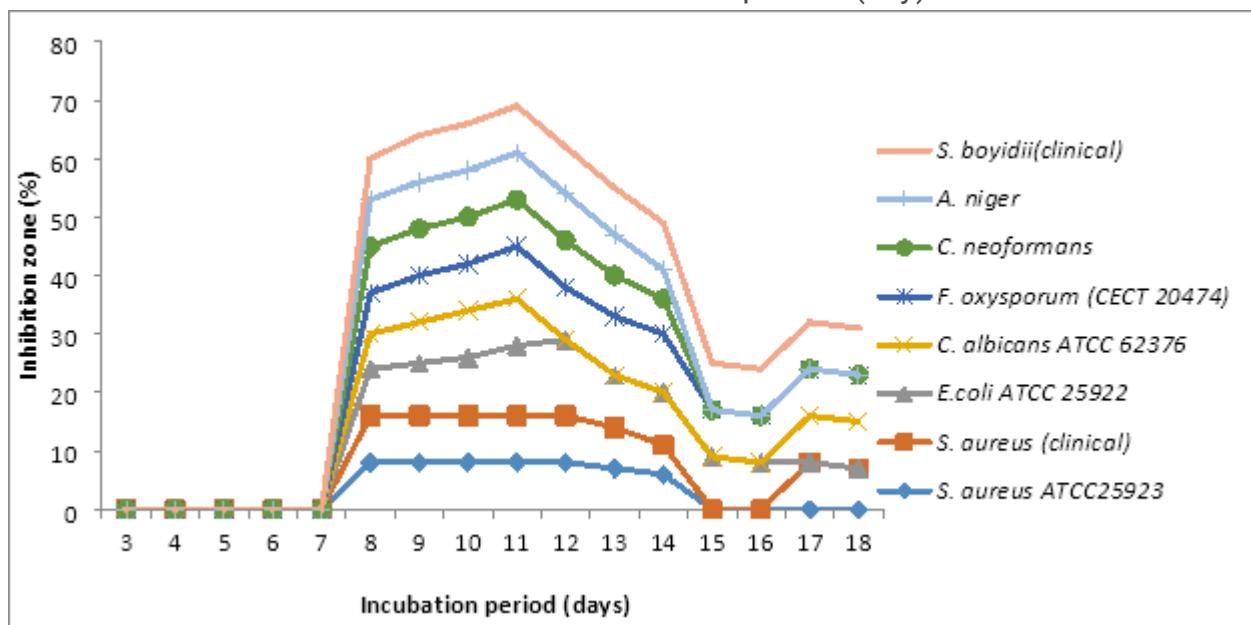


Figure 2

Diameter of zone of inhibition (mm) of an antibacterial and antifungal metabolite of LAAACC15 taken at different incubation periods (day).

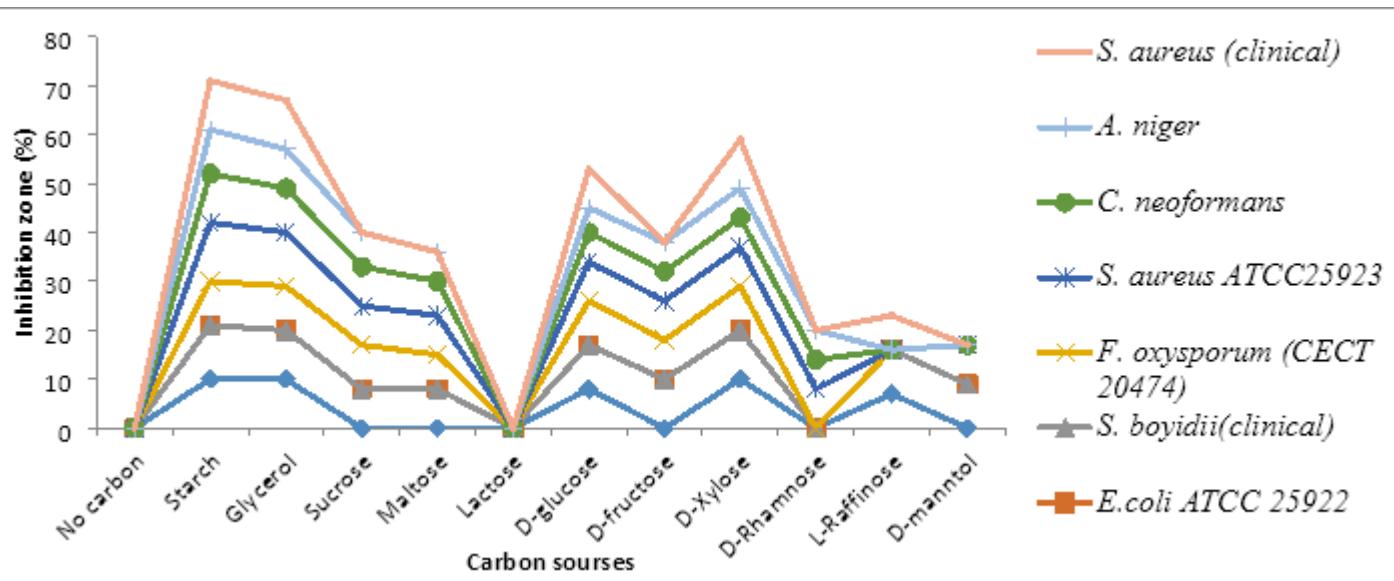


Figure 3

Diameter of the disc (6mm) of zone of inhibition of LCHAACC13 grown on different carbon sources.

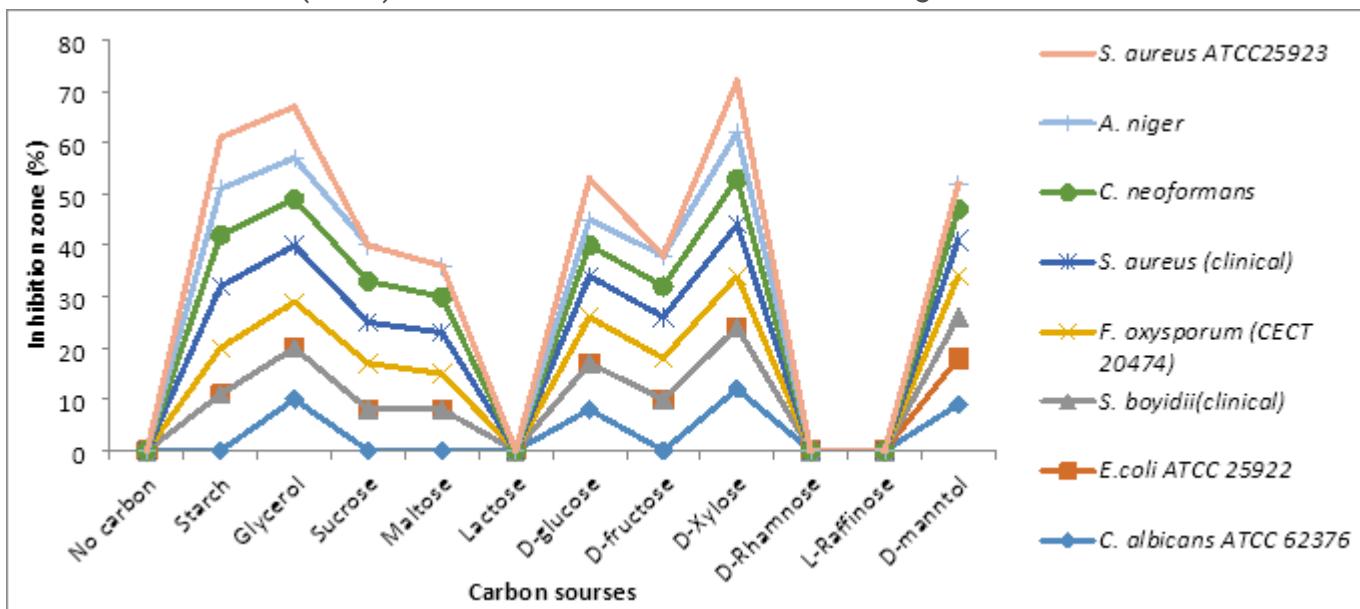


Figure 4

Diameter of zone of inhibition of LAAACC15 grown on different carbon sources.

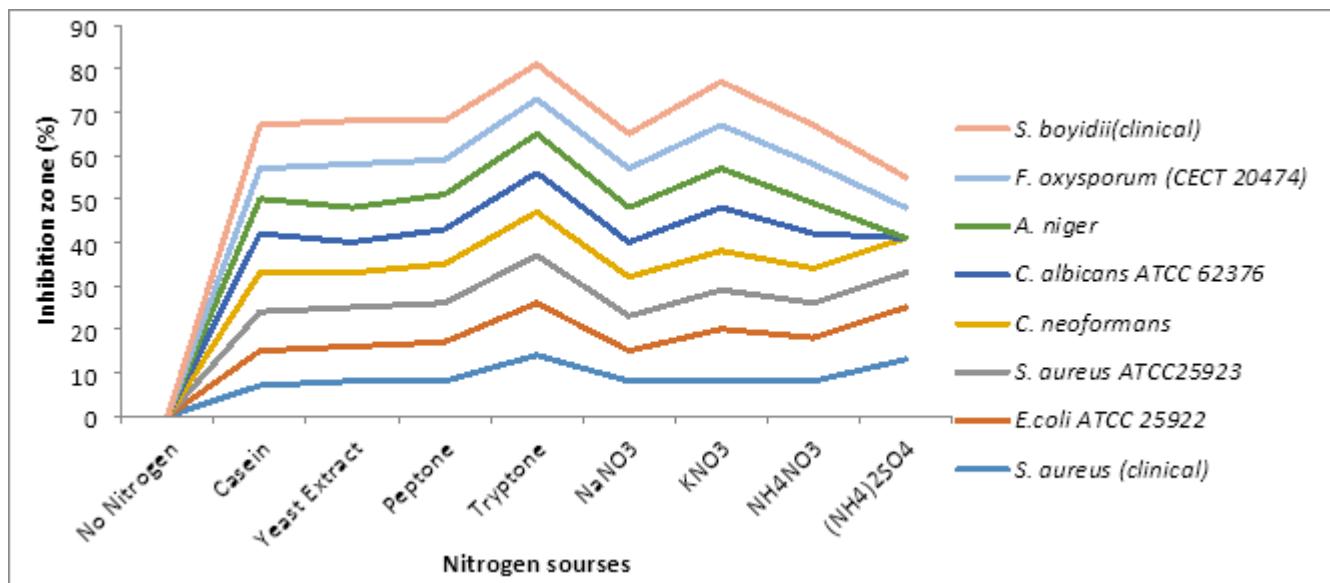


Figure 5

Diameter of zone of inhibition in mm of LCHAACC13 grown on different nitrogen sources.

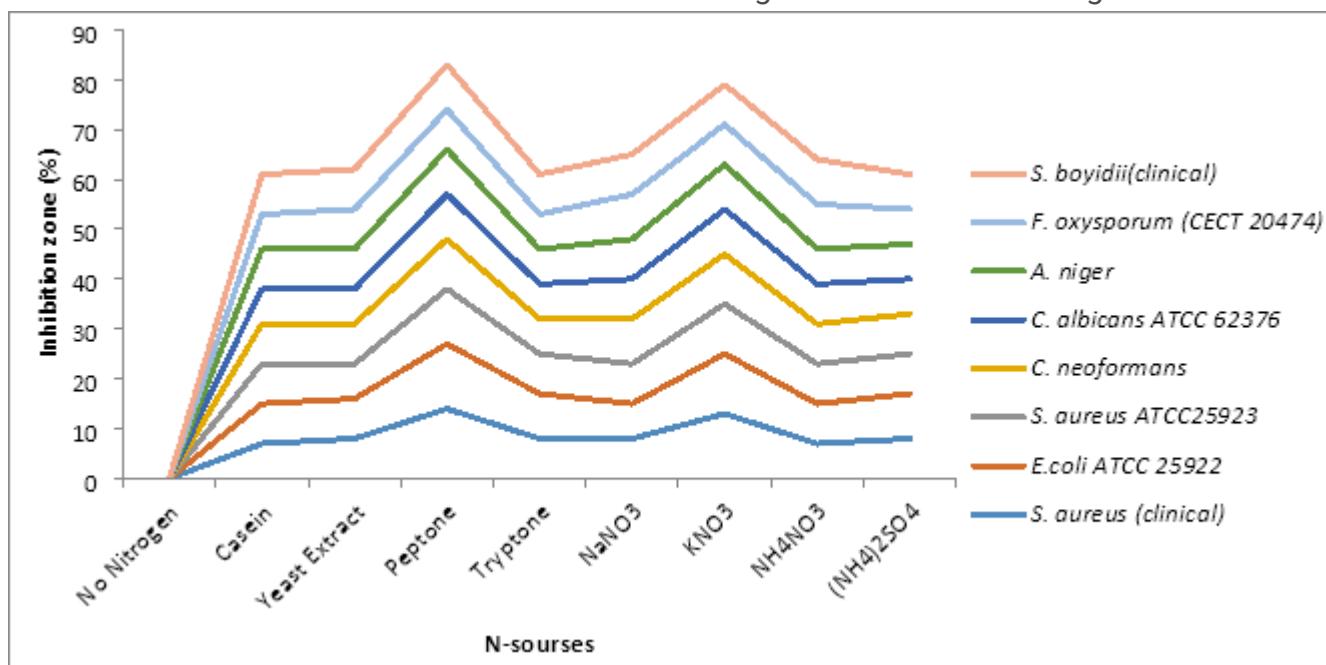


Figure 6

Diameter of zone of inhibition zone in mm of LAAACC15 grown on different nitrogen sources against

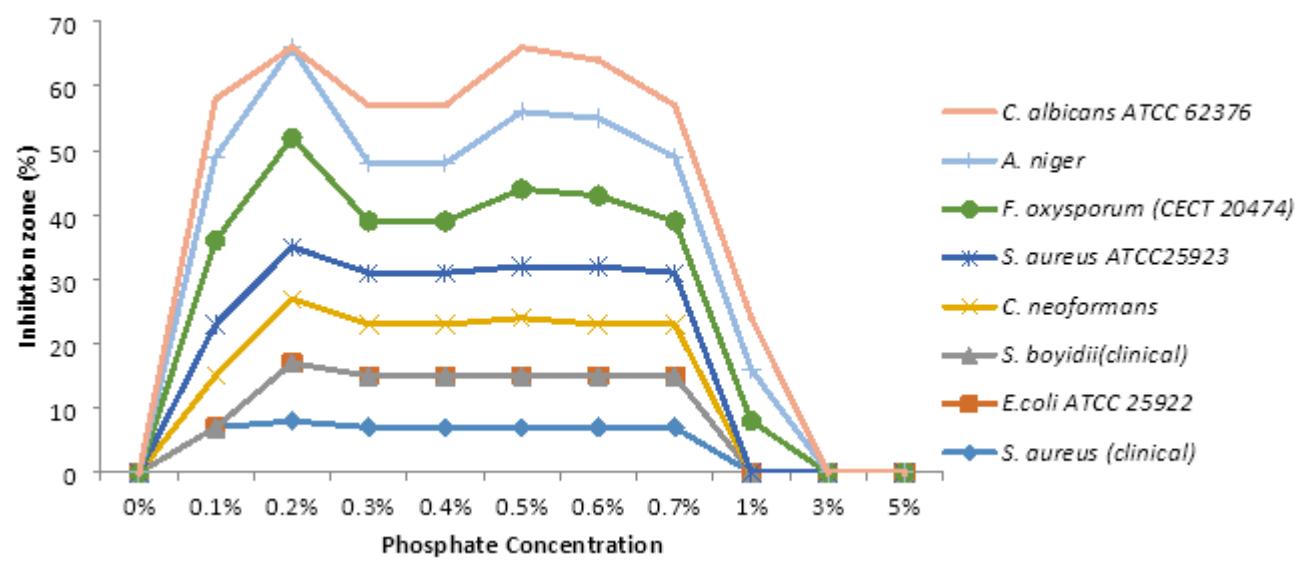


Figure 7

Extracted crude filtrate of LCHAACC13 on 12th day on different K₂HPO₄ concentrations and antibacterial activity.

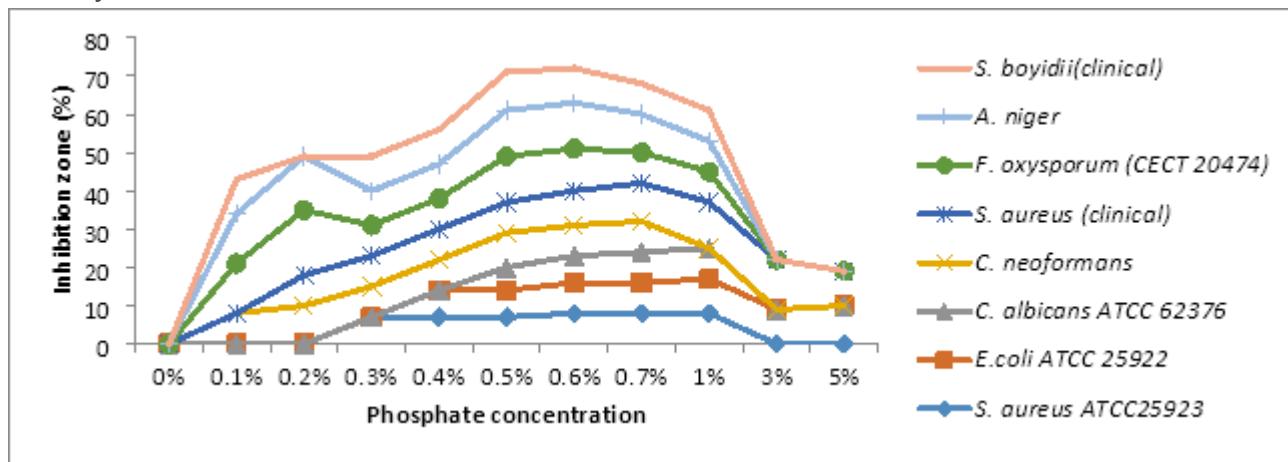


Figure 8

Extracted crude filtrate of LAACC15 on 14th day on different phosphate concentrations and diameter of zone of inhibition in 6 mm disc.

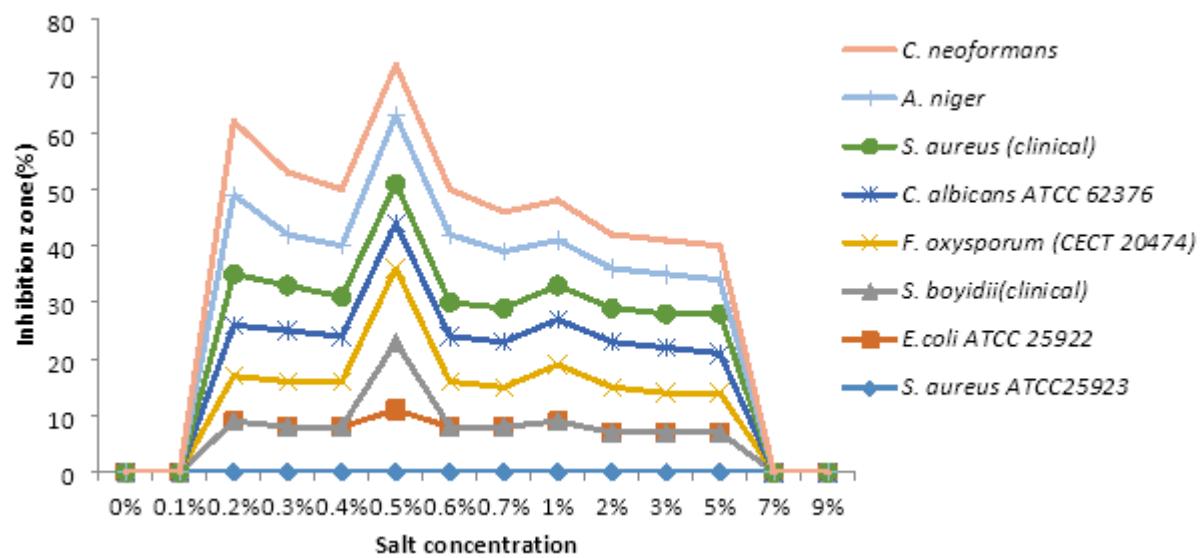


Figure 9

Test microorganisms and diameter of zone of inhibition in mm of LCHACC13 grown on different NaCl concentrations.

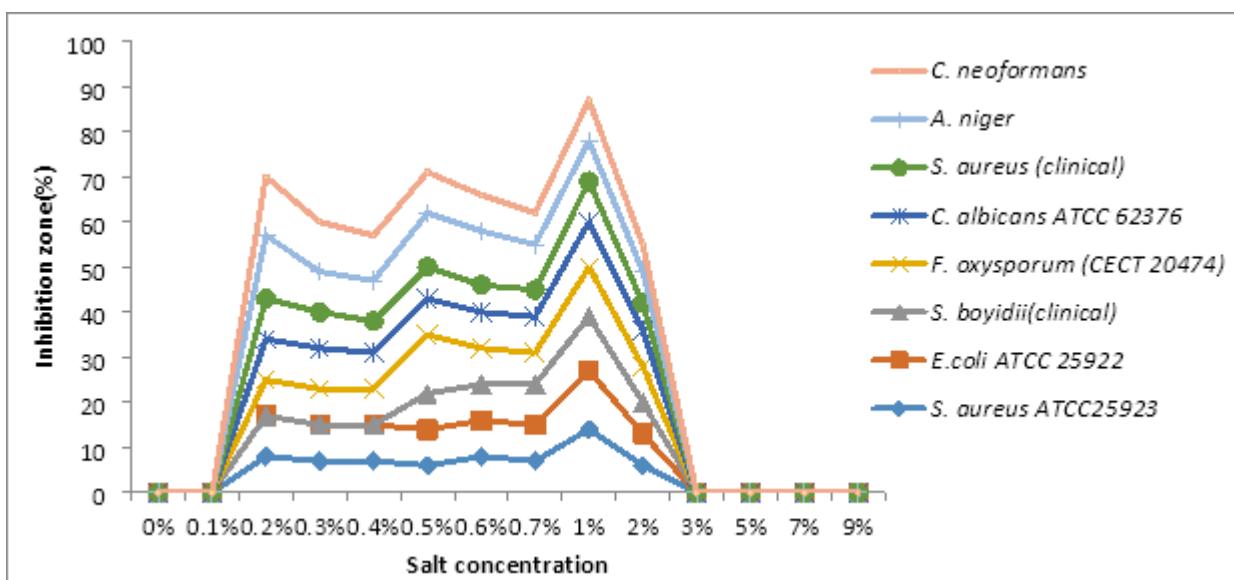


Figure 10

Test microorganisms and diameter of zone of inhibition in mm of LAACC15 grown on different NaCl concentrations.

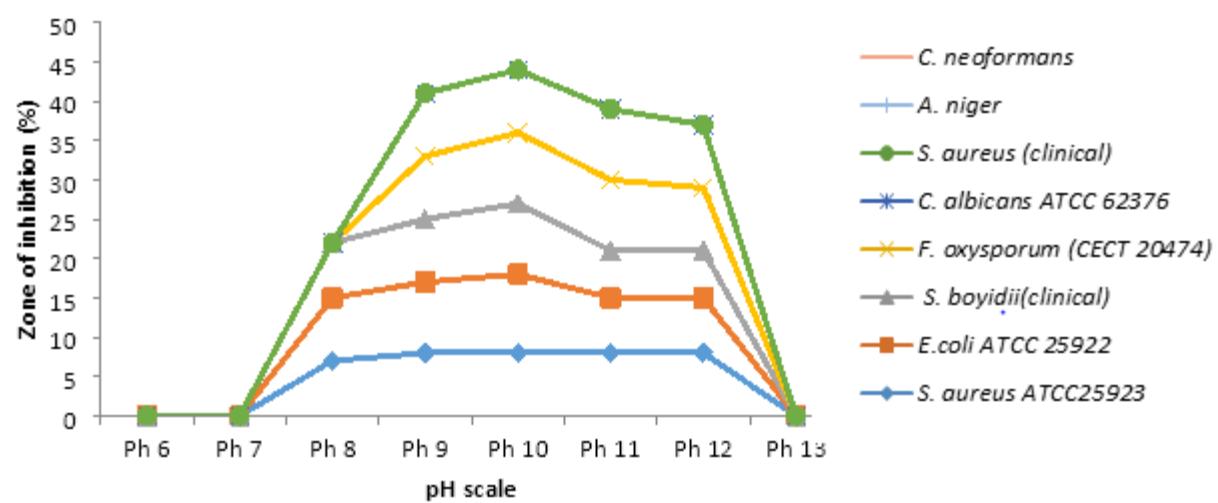


Figure 11

Diameter of zone of inhibition of LCHAACC 13 grown on different pH

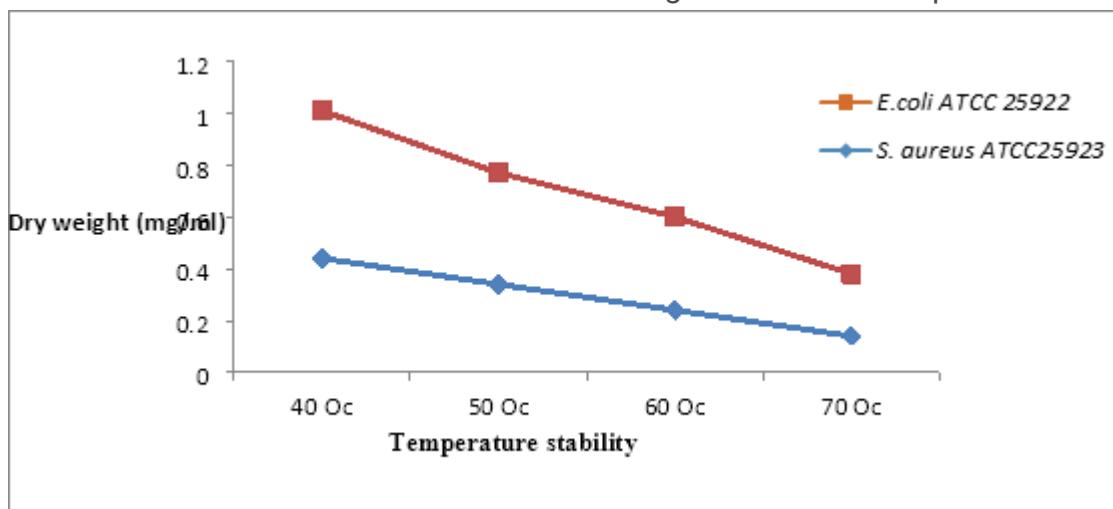


Figure 12

Temperature stability of isolates of two soda lakes. Both isolates of LCHAACC13 and LAAACC15 had the stability of 40 oC- 60 oC, but the range of a temperature supporting good growth was as wide as 20 oC- 30 oC.

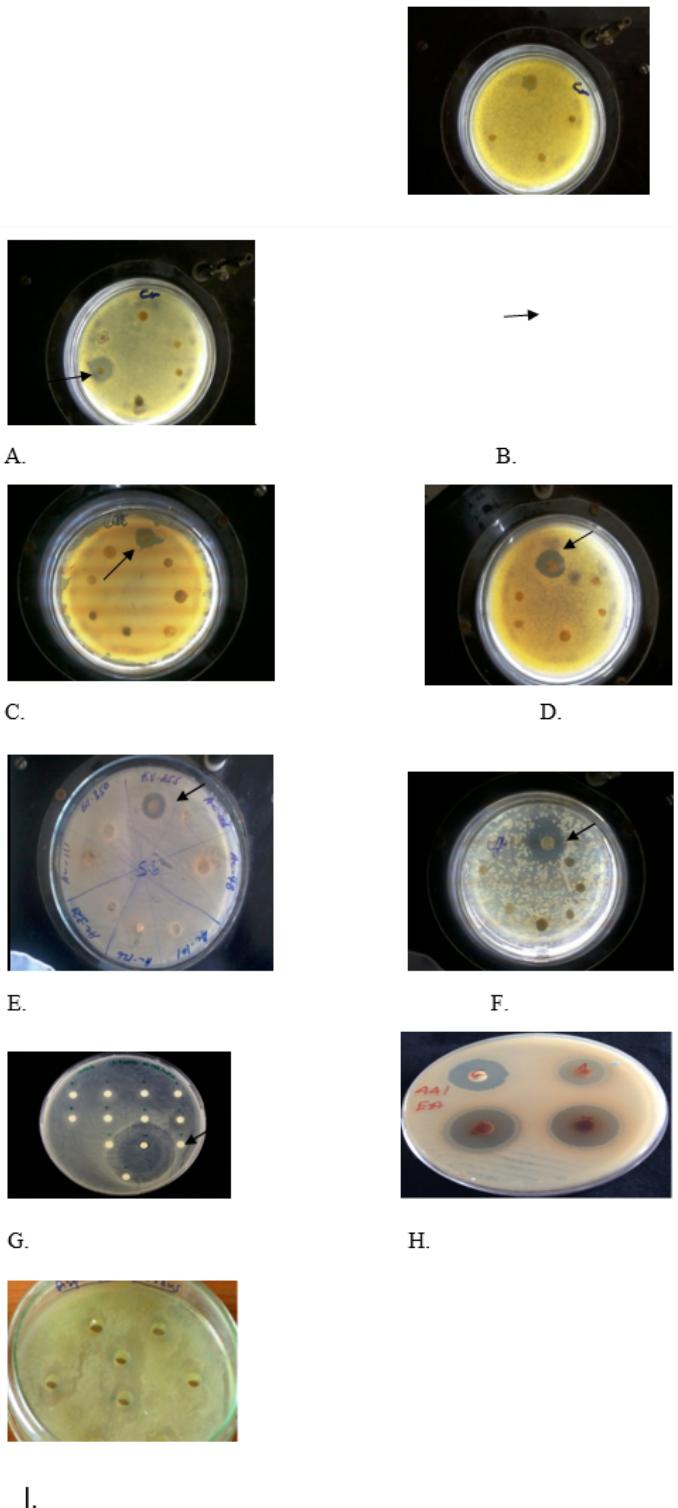


Figure 13

Inhibition of some representative test organisms by the extracts from selected *Streptomyces* spp.
Inhibition of some test organisms by the extracts of 8 active isolates from selected *Streptomyces* species.
A. Lc-13 against *C. neoformans* B. Lc-15 against *C. neoformans* C. Lc-17 against *C. albicans* D. La-13
against *C. neoformans* E. Lc-19 against *S. boydii* F. Lc-21 against *S. aureus* (clinical) G. La-15against *S.*

aureus H. La-15 against *F. oxysporum* (CECT 20474). I. La-17 against *Aspergillus niger*. N.B: Lc- stands for active isolates of Lake Chitu (Lc); La- active isolates of Lake Arenguade (La)

Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [Preliminary antimicrobial screening of 10 active isolates against different microbial pathogens.docx](#)
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- [Pure culture of fungal genera for primary screening.jpg](#)
- [Monop2.jpg](#)
- [pink.jpg](#)
- [Incubation in incubator.jpg](#)