



Evaluation of *In Vitro* Antibacterial Potential of *Bacillus pseudomycooides* Strain SB138

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ABSTRACT

Bacillus species provide a natural source for the discovery of novel antibacterial substances. However, some of the *Bacillus* species still remained to be explored widely for their antagonism against harmful bacteria. One of them is *Bacillus pseudomycooides* which produces rhizoidal colonies on solid medium. The present study was aimed to explore antibacterial properties of *B. pseudomycooides* isolated from the rhizosphere soil of the indigenous *Mangifera indica* (mango) plant. Identification of the isolated strains was determined by sequencing their 16S rRNA genes. The isolates identified as *B. pseudomycooides* were screened for their antibacterial potential. In order to circumvent the trouble created from rhizoidal colony morphology of *B. pseudomycooides*, a colony mutant of a strain showing strong antibacterial activity was obtained by screening/selection method and designated as *B. pseudomycooides* strain SB138m. The sequence of 16S rRNA gene of both wild type and mutant strains of *B. pseudomycooides* was comparable. *In vitro* test for antagonism of *B. pseudomycooides* strain SB138m against indicator cells showed that the strain contained antibacterial potential. The present study shows inhibitory spectrum of *B. pseudomycooides* SB138m with conclusion that *Bacillus* species from a diverse environment and *Proteus mirabilis* from clinical environment were strongly inhibited.

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Authors' Contribution

RP, SB, SAT conceived and designed research. RP and AAS conducted experiments. ANJ contributed to analytical tools. SB and SAT analyzed data. RP, SB, and SAT wrote the manuscript. SB and SAT reviewed and edited the manuscript critically. All authors have read and approved the manuscript.

Key words

Antagonism, *Bacillus*, Antimicrobials, Biopreservatives, Alternative therapeutics

INTRODUCTION

Bacillus species are Gram-positive, endospore forming rod shaped bacteria. They are found in diverse environment including the rhizosphere soil of plants, soil, food, water environment and gastrointestinal systems of birds, insects and different animals (Nicholson, 2002; Liu *et al.*, 2015; Díez-Méndez *et al.*, 2017). They are motile except *B. anthax*, *B. mycooides* and *B. pseudomycooides*.

A wide array of antibacterial substances with unique properties produced from *Bacillus* species have been discovered and characterized (Sumi *et al.*, 2014; Huang *et al.*, 2016; Zhao and Kuipers, 2016). These include both ribosomally synthesized peptides (RPs) and none ribosomally (enzymatically) synthesized peptides (NRPs). Interestingly, *Bacillus* species have an ability of producing

different types of antimicrobial substances simultaneously. For example, a strain of *B. subtilis* can produce both surfactin and fengycin (Sun *et al.*, 2006). Consequently, a recent study has identified 252 putative antimicrobial gene clusters in only 39 genomes of *B. subtilis* (Zhao and Kuipers, 2016). However, only one genome of *B. pseudomycooides* was included highlighting need of exploration of *B. pseudomycooides* strain with antibacterial activities.

Bacillus pseudomycooides are spore forming non-motile bacteria (Nakamura, 1998). They form characteristic rhizoidal (a symmetric hairy shape) colonies on solid media. They are closely related to *Bacillus cereus* group of bacteria. In recent years, *B. pseudomycooides* have been found producing biosurfactants, extracellular polysaccharides and solubilizing potassium uptake in tea plants (Li *et al.*, 2016; Solmaz *et al.*, 2018; Pramanik *et al.*, 2019). Although there is a rich source of literature available that describes the production of antimicrobials from different species of genus *Bacillus*, only a few researchers have focused on the investigation of antimicrobial activity of *B. pseudomycooides*. In this context, a study has reported the production of antibiotics from *B. pseudomycooides* DSM 12442 suggesting that the bacterium has antibacterial potential and can serve as a natural source of antibacterial agent (Basi-Chipalu *et al.*, 2015). In the present study,

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B. pseudomycooides strain SB138 having antibacterial potential has been characterized and its antagonism against a wide range of bacterial isolates were determined.

MATERIALS AND METHODS

Bacterial strains and growth conditions

Bacterial strains used in the present study are listed in Table I. All the cultures were maintained on nutrient agar at 37°C aerobically unless otherwise mentioned. All media were purchased from Oxoid, UK.

Isolation of B. pseudomycooides

Soil samples were collected from the rhizosphere of *M. indica* plants cultivated at Jamshoro, Pakistan as

described previously (Bano *et al.*, 2018). Briefly, soil adhered to the roots of the plant were shaken against the sides of beaker or flask vigorously and immediately covered with cotton plug or aluminum foil, respectively. Then, one gram of soil sample was mixed into 5ml of sterile nutrient broth and heated for 15 min at 80°C and incubated overnight aerobically. Following the incubation (enrichment stage), tenfold serial dilutions were made using nutrient broth. The last 3 dilutions (10^{-2} , 10^{-3} , and 10^{-4}) were pipette out and poured onto nutrient agar and plates were incubated overnight at 37°C aerobically. After incubation, rhizoidal colonies were selected and streaked onto fresh nutrient agar separately to obtain pure culture colonies. The isolated colonies were initially identified by their phenotypic characteristics.

Table I. Spectrum of the inhibitory effects of *B. pseudomycooides* SB138m.

Bacterial isolates	Source/ Reference	Inhibitory effects of <i>B. pseudomycooides</i> SB138m Positive (+) / negative (-)
<i>Staphylococcus aureus</i> ATCC 6538 (indicator strain)	Purchased (Oxoid, UK)	++
<i>Escherichia coli</i> ATCC 25922	Purchased (Oxoid, UK)	-
<i>S. aureus</i>	Pus specimen	-
<i>S. aureus</i>	(Afreen <i>et al.</i> , 2020)	-
<i>S. aureus</i>		++
<i>Pseudomonas aeruginosa</i>		-
<i>S. aureus</i> (Methicillin resistant)	Wound specimen	++
<i>E. coli</i>	Urine specimen	-
<i>P. fluorescens</i>	(Afreen <i>et al.</i> , 2020)	-
<i>Klebscilla pneumoniae</i>	Device associate infections	-
<i>S. epidermidis</i>	(Afreen <i>et al.</i> , 2020)	++
<i>Enterobacter</i> sp.		-
<i>Citrobacter freundii</i>		++
<i>Proteus mirabilis</i> (4S)	Clinical sample*	++++
<i>Bacillus cereus</i> (SB 47)	Soil*	+++
<i>Enterococcus</i> sp. (SA 44)	Clinical sample*	++
<i>Bacillus subtilis</i> (RP T1)	Spoiled tomato pulp*	++++
<i>Shigella dysenteriae</i>	Drinking water	+
<i>Salmonella</i> sp.	Fresh water sample*	+
<i>Listeria monocytogenes</i>	Raw salad*	+
<i>Bacillus subtilis</i> (SB65)	Rhizosphere soil*	++++
<i>Bacillus</i> sp. (SB70)	Rhizosphere soil*	+++
<i>Bacillus</i> sp. (SB87)	Environment*	++++
<i>Bacillus</i> sp. (SB89)	Raw salad*	+++

++ moderate activity; +++ substantial activity; ++++ strong activity. *Laboratory stock.

16S rRNA sequencing of *B. pseudomycooides* isolates

Isolate identities were determined by sequencing 16S rRNA genes from Macrogen (Korea). The sequencing was performed by using Big Dye terminator cycle sequencing kit (Applied BioSystems, USA). Sequencing products were resolved on an AB model 3730XL automated DNA sequencing system (Applied BioSystems, USA). Universal primers 27F (5'AGAGTTTGATCMTGGCTCAG3') and 1492R (5'TACGGYTA CCTTGTTACGACTT3') were used for PCR amplification of the 16S rRNA gene. Universal primers 785F (GGATTAGAT ACCCTGGTA) and 907R (CCGTCAATTCMTTTRAGTTT) were used for sequencing.

Selection of colony mutant of *B. pseudomycooides*

A colony mutant of *B. pseudomycooides* SB138 was obtained on nutrient agar as described previously (Di Franco *et al.*, 2002). The resultant isolate, *B. pseudomycooides* SB138m, was no more rhizoidal, making round compact colonies. Gram's staining was performed according to standardized method (Claus, 1992). Spore staining was performed according to Schaeffer-Fulton method (Schaeffer and Fulton, 1933). The sequencing of 16S rRNA gene of the mutant strain was also performed again for the confirmation of the isolate identity.

Antibacterial assays

In vitro antibacterial activity of *B. pseudomycooides* SB138m was determined by stab-overlay method and/or spot- overlay method as described previously (Bano *et al.*, 2014). Briefly, a single colony of *B. pseudomycooides* SB138m was stabbed into nutrient agar plate (stab-overlay method) or 2µl of fresh overnight culture of the strain was spotted onto nutrient agar plate (spot-overlay method). The plates were incubated at 28±2°C for at least 16h. After incubation, cells were killed by exposure to chloroform vapours for 30 min for releasing antibacterial substance into the medium. After incubation, 10ml of soft top agar (0.7% agar) seeded with 100µl test culture (10⁸ CFU/ml) was poured onto medium followed by incubation at 37°C for at least 16h. Next day plates were observed for antibacterial activity indicated by the presence of any zone of inhibition of growth of test culture around the stabbed/ spotted culture of *B. pseudomycooides* SB138m.

Determination of effects of heat, ultraviolet rays, and proteolytic enzymes

Moreover, effects of heat, UV and proteolytic enzymes were tested as described previously (Parret *et al.*, 2003). Briefly, an agar plate containing *B. pseudomycooides* SB138m incubated for 6.5 h was exposed to UV light (312 nm) for 30s or placed in an oven at 75°C for 15 min

followed by cooling at room temperature for 30 min. The culture plates were then overlaid as described elsewhere. For testing the sensitivity of *B. pseudomycooides* SB138m to proteolytic enzymes, overnight culture was spotted on the surface of agar plate and incubated. The resultant growth was exposed to the chloroform vapors and a spot of pepsin, trypsin or Proteinase K (20mg/ml, Merck) was put near the growth of *B. pseudomycooides* SB138m. After drying of the drop, plates were incubated for at least 1h at 37°C to allow optimal proteolytic activity.

RESULTS

B. pseudomycooides SB138

A strain of *B. pseudomycooides* was isolated from the rhizosphere soil of *M. indica* plant. The strain was initially identified on the basis of its phenotypic characteristics such as rhizoidal colonies, lack of motility and no hemolysis on blood agar. Subsequently, the strain was identified at species level by 16S rRNA gene sequencing and designated as *B. pseudomycooides* SB138 (Fig. 1). The nucleotide sequence of 16S rRNA gene of the novel strain has been deposited to Gene Bank under accession number MH578628. The sequence had 99% similarity with *B. pseudomycooides* DSM 12442 which is a type strain *B. pseudomycooides* (Nakamura, 1998). *B. pseudomycooides* SB138 appeared as ampicillin resistant and ciprofloxacin sensitive.

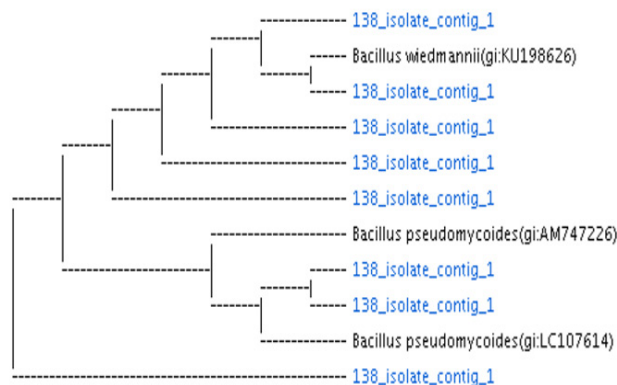


Fig. 1. 16S rRNA gene-based tree showing the phylogenetic relationship of the isolate SB 138 with type strain of *B. pseudomycooides*.

Colony mutant of *B. pseudomycooides* SB138

B. pseudomycooides produces rhizoidal growth on solid medium (Fig. 2a), which brings a challenge for researchers during the performance of antibacterial assays with this bacterium. In order to work on antibacterial aspects of *B. pseudomycooides* conveniently, it was very

necessary to demolish the barrier of rhizoidal growth, therefore a colony mutant of *B. pseudomycooides* SB138 having no more rhizoidal growth morphology was obtained (Fig. 2b). The colony mutant strain was named as *B. pseudomycooides* SB138m. The 16S rRNA gene sequencing of *B. pseudomycooides* SB138m revealed 99% similarity between the sequences of 16S rRNA genes of wild type and the mutant strains (Fig. 3) confirming that the rhizoidal morphology of *B. pseudomycooides* SB138 is lost due to spontaneous mutation as reported previously (Di Franco *et al.*, 2002).

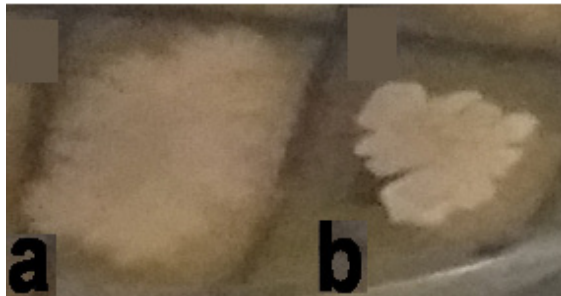


Fig. 2. Colony characteristics of *B. pseudomycooides* SB138 a) the wild type *B. pseudomycooides* SB138, and b) its colony mutant *B. pseudomycooides* SB138m.

```
#
# Aligned_sequences: 2
# 1: Wildtype
# 2: Colonymutant
# Matrix: EDNAFULL
# Gap_penalty: 10.0
# Extend_penalty: 0.5
#
# Length: 1474
# Identity: 1473/1474 (99.9%)
# Similarity: 1473/1474 (99.9%)
# Gaps: 0/1474 (0.0%)
# Score: 7361.0
#
#-----
Wildtype      1 CTCGGATGAACGCTGGCGGCGTGCCCTAATACATGCAAGTCGAGCGAACCG 50
Colonymutant  1 CTGGGATGAACGCTGGCGGCGTGCCCTAATACATGCAAGTCGAGCGAACCG 50
Wildtype     51 ATTAAGAGCTTGCTCTTATGAAGTTAGGGGACGGGTGAGTAAACACGTGGG 100
Colonymutant 51 ATTAAGAGCTTGCTCTTATGAAGTTAGGGGACGGGTGAGTAAACACGTGGG 100
Wildtype    101 TAACCTGCCCATAAAGACTGGGATAACTCCGGGAAACCGGGGCTAATACCG 150
Colonymutant 101 TAACCTGCCCATAAAGACTGGGATAACTCCGGGAAACCGGGGCTAATACCG 150
Wildtype    151 GATAATATTTGCGCCTCATGGCGCAAAATTGAAAGGCGGCTTCGGCTGT 200
Colonymutant 151 GATAATATTTGCGCCTCATGGCGCAAAATTGAAAGGCGGCTTCGGCTGT 200
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Fig. 3. Confirmation of the identification of *B. pseudomycooides* SB138m using 16S rRNA gene sequencing.

Antibacterial activities of B. pseudomycooides strain SB138m

B. pseudomycooides strain SB138 showed antagonism against an indicator strain, *S. aureus* ATCC 6538 (Fig. 4a). The spectrum of the inhibitory effects of the strain was found against a range of the *Bacillus* spp isolated from a diverse environment and *P. mirabilis* isolated from a

clinical sample (Fig. 4b). However, clinical isolates of *E. coli*, *Pseudomonas* spp., *Klebsiella* spp. were not sensitive to the *B. pseudomycooides* strain SB138m (Table I).

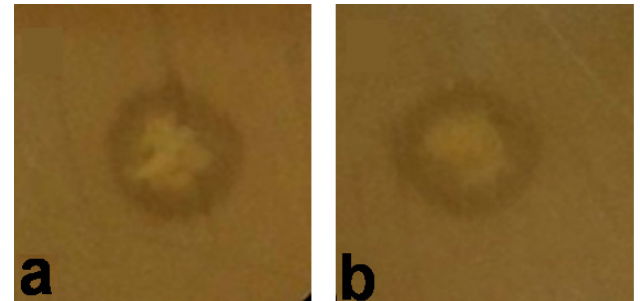


Fig. 4. *In vitro* antibacterial activity of *B. pseudomycooides* strain SB138 against (a) indicator strain, *S. aureus* ATCC 6538 and (b) *P. mirabilis* 4S determined by stab- overlay methods.

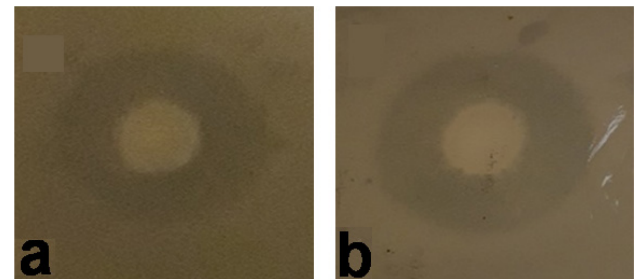


Fig. 5. Effect of (a) heat and (b) UV on antibacterial activity of *B. pseudomycooides* strain SB138m determined by spot-overlay method.

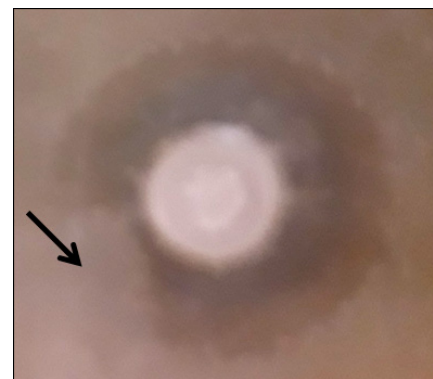


Fig. 6. Determination of the effects of Proteinase K on antibacterial activity of *B. pseudomycooides* strain SB138m.

Stability of antibacterial activity of B. pseudomycooides strain SB138m

The antibacterial activity of *B. pseudomycooides* strain SB138m was found to be stable at high temperature

and UV exposure (Fig. 5a, b). It was also observed that proteinase K enzyme demolished the antibacterial activity of *B. pseudomycooides* strain SB138m completely (Fig. 6). These finding suggested that *B. pseudomycooides* strain SB138m presumably produce bacteriocin like antibacterial substance. Furthermore, the plasmid profiling of *B. pseudomycooides* SB138m suggested that the strain does not contain any plasmid indicating that the gene for bacteriocin production may be genome encoded as supported by a recent study which has found the putative antimicrobial gene clusters on the genome of *B. pseudomycooides* (Zhao and Kuipers, 2016).

DISCUSSION

Discovering natural antimicrobials from *B. pseudomycooides* remained ignored presumably due to its rhizoidal growth features on solid medium. However, a recent study reported that the *B. pseudomycooides* DSM 12442 produces lantibiotic namely Pseudomycoicidin, which was active against a wide range of Gram-positive bacteria only (Basi-Chipalu et al., 2015). Since *Bacillus* species are known to produce antibacterial substances with varied properties concurrently, it is not surprising that genes potentially involved in the synthesis of antimicrobial substances other than Pseudomycoicidin were found on the genome of *B. pseudomycooides* (Basi-Chipalu et al., 2015). Consequently, *B. pseudomycooides* BS6, an isolate of an edible oil contaminated soil was found producing biosurfactants which were lipopeptides in nature (Li et al., 2016).

In the present study, *B. pseudomycooides* strain SB138 was found of holding strong antagonism against *S. aureus* and *Bacillus species*. Notably, inhibition of *B. subtilis* isolated from spoiled pulp of tomato indicated the biopreservative potential of the strain. Furthermore, the strain inhibited the growth of quinolone resistant clinical isolate of *P. mirabilis*. Our findings are supported by another study which identified putative antimicrobial gene clusters for 4 RPs (including a type II lantipeptide) and 3 NRPs on the genome of *B. pseudomycooides* (Zhao and Kuipers, 2016).

The antibacterial activity of *B. pseudomycooides* SB138 was fully retained after 15 min at 75°C and it was completely destroyed by proteinase K treatment, indicating its proteinaceous nature. These finding are in contrast of previous reports which mentioned that the most of antibiotics are sensitive to the activity of proteases (Barbour et al., 2013). However, our findings are in agreement of a recent study reporting that Pseudomycoicidin lantibiotic with four thioether rings and a disulfide bond is resistant to the activity of proteases (Basi-Chipalu et al., 2015).

CONCLUSION

The results of the present study show the antibacterial potential of *B. pseudomycooides* SB138 and have indicated possible use of *B. pseudomycooides* strain SB138m in future strategies for the development of alternative therapeutic agent or bio preservative.

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Ethical statement

This article does not contain any studies with human or animal subjects.

Statement of conflict of interest

The authors have declared no conflict of interest.

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