



Optimization of Degenerate PCR Conditions for Reducing Error Rates in Detection of PKS and NRPS Gene groups in Actinomycetes

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Abstract

Background: The screen of Polyketide Synthase (PKS) and Nonribosomal Peptide Synthetase (NRPS) gene groups is a quick way to discover new therapeutic agents. However, errors in laboratory techniques cause a loss of touch with reality. This study aimed to evaluate the presence of PKS and NRPS gene groups in previously isolated strains by optimizing their specialized amplification by degenerate primers and indicating the evolutionary relationships with reference strains.

Methods: PKS-I, II, and NRPS genes PCR amplification was performed using three degenerate primer sets for 22 actinomycete strains with antibacterial activity. Annealing temperature and the amount of template DNA and primers were optimized. PCR products of PKS-I, II, and NRPS from three strains were sequenced after TA cloning. Besides, strains with high antibacterial activity were identified by biochemical features and partial 16S rDNA sequencing and hypothetically classified by a phylogenetic tree.

Results: High frequencies of PKS-I (86.4%), PKS-II (81.8%), and NRPS (95.4%) genes were found among the strains after optimization. Fourteen strains (64%) contained all of the genes, and 100% of strains had at least two genes. These numbers are pretty distinct in comparison with the previous researches. All of the sequenced strains were members of *Streptomyces* genus.

Conclusion: Our research showed that degenerate PCR strongly depends on the variation of annealing temperature and primer concentration, resulting in an unexpected shift in PCR outputs. The sequencing results confirmed the optimized conditions for specialized PCR of PKS-I, PKS-II, and NRPS gene groups.

Keywords: Antimicrobial peptides, Biological products, PCR, Polyketide synthase, *Streptomyces*

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Introduction

Actinobacteria represent a significant component of the microbial population in most soils. This phylum of bacteria has been extremely useful to the pharmaceutical industry due to its seemingly unlimited capacity to produce secondary metabolites with diverse biological activities and chemical structures. Approximately 50% of Actinobacteria are from the genus *Streptomyces*, and about 75% of commercially valuable antibiotics are derived from this genus¹.

However, novel technologies are demanded for discovering new bioactive secondary metabolites from

these bacteria to meet the persistent and urgent medical need for newer and better drug candidates. Based on the hypothesis that secondary metabolites with similar structures are likely biosynthesized by gene clusters harboring certain homologous genes, such homologous genes could serve as appropriate selectable markers for different natural product biosynthetic gene clusters^{2,3}. A wide range of structurally diverse bioactive compounds, including toxins, siderophores, pigments, antibiotics, antitumor agents, and immunosuppressants, are synthesized on large Nonribosomal Peptide Synthetase

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(NRPS) and Polyketide Synthase (PKS) enzyme complexes⁴⁻⁶. Therefore, screening for genes associated with secondary metabolism gives a better insight into the genetic capabilities of actinomycetes^{2,3}. NRPSs are composed of catalytic units called modules⁶ working in an assembly-line fashion, where each module is responsible for incorporating a single monomer⁷. According to the collinearity rule, the modules' number and order define the number and order of amino acids in NRPs⁶. Enzyme modules contain three core catalytic domains: adenylation (A), thiolation or peptidyl carrier (T or PCP), and condensation (C), respectively responsible for recognition and activation of an amino acid, transfer of the activated substrate to the peptidyl-carrier domain of the same module, and the formation of the C-N bond between the elongated chain and the activated amino acid^{6,8}. Actually, in the biosynthesis of NRPs, the enzymatic machinery itself functions as a template for the synthesized peptide product⁷. PKSs are synthesized through sequential decarboxylative condensation of the acyl CoA precursors by PKS. PKSs are classified into types I, II, and III⁹. Type I PKSs, similar to type I Fatty Acid Synthase (FAS), are large multifunctional enzymes with a modular organization, with each module comprising catalytic domains^{5,9,10}. Typically, each module contains three essential catalytic domains for chain elongation: Ketosynthase (KS), Acyl Transferase (AT), and Acyl Carrier Protein (ACP). Bacterial type I PKSs are usually noniterative¹¹. Type II PKSs, similar to type II FAS, are multienzyme complexes acting in an iterative manner^{9,11}. They consist of a heterodimer formed by a KS α subunit (active site) and a KS β or Chain Length Factor (CLF) subunit, which collaborate with ACP¹². Type III PKSs are homodimeric enzymes. Their single active site in each monomer catalyzes the priming, extension, and cyclization reactions iteratively to form PK products^{9,11}. Despite the structural simplicity, bacterial type III PKSs fall into five groups, many of which have been identified from *Streptomyces*¹¹. In addition to the common form of types I and II, exceptions such as iterative type I PKSs and non-iterative type II PKSs have also been reported by now⁹. Due to the evolution of multidrug-resistant strains, screening of natural resources, especially actinomycetes, represents an important route to discover new therapeutic agents. In addition, the screening of *PKS* and *NRPS* genes is a quick way to show the potential for antibacterial agent production.

The overall aims of this study were (i) to evaluate the presence of *PKS* (types I and II) and *NRPS* gene groups involved in the biosynthesis of secondary metabolites in twenty-two antibacterial agent-producing actinomycete strains isolated in our previous study¹³. For example, A3F and A7R primers target the conserved motifs A3 and A7 in adenylation domain. Therefore, PCR products can have different sizes (between 700-800) from the conserved motifs 3 to 7, (ii)

to study the effect of different factors on degenerate PCR amplification, (iii) to scrutinize the power of the optimized reaction conditions for specialized amplification of *PKS-I*, *PKS-II* and *NRPS* gene groups by sequencing, and (iv) to show the phylogenetic and evolutionary relationships between some of the high potential antibacterial agent-producing strains and reference strains.

Materials and Methods

Actinobacterial strains

Twenty-two antibacterial agent-producing actinomycete strains (F4, F6, F9, F10, Is1, Is4, Is5, Ih1, Ih6, Iz1, Iz2, Iz6, Iz8, Iz9, Iz11, Iz14, Iz16, Iz17, D2, B4, N1, and A2) isolated in our previous study¹³ were selected for further investigation. All strains were screened for the presence of *PKS* and *NRPS* genes. The high potential antibacterial agent-producing strains (F4, F6, F9, F10, Iz6, Iz8, Iz16, and B4) were also selected for molecular identification by 16S rDNA sequencing. The most active isolates against hospital-resistant strains and, in the next step, the strains with the largest halo diameter (Iz8, F9, and F6) were selected for cloning.

DNA extraction

DNA was extracted by CTAB method¹⁴. The only difference for strain F9 was that a lysis treatment was applied at the beginning of protocol¹⁵.

PCR detection of *PKS-I*, *PKS-II*, and *NRPS* genes

Genomic potential for producing bioactive metabolites was evaluated using three sets of specific degenerate primers: (1) K1F (5'-TSAAGTCSAACATCGGB CA-3'), and M6R (5'-CGCAGGTTSCSGTACCAGT A-3') targeting *PKS-I* KS and methyl malonyl transferase domains¹⁶⁻¹⁸; (2) KS α F (5'-TSGCSTGCTTGGA YGCSATC-3'), and KS α R (5'-TGGAANCCGCCGAA BCCTCT-3') targeting *PKS-II* KS α subunit^{17,18}; and (3) A3F (5'-GCSTACSYSATSTACACSTCSGG-3'), and A7R (5'-SASGTCVCCSGTSCGGTAS-3') targeting the conserved motifs A3 and A7 in NRPS adenylation domain¹⁶⁻¹⁸. PCR conditions for the initial amplification of *PKS-I*, *PKS-II*, and *NRPS* were as follows: a 25 μ l sample of the PCR reaction mixture containing 10-300 ng of genomic DNA, 0.4-4 μ M of each primer, 200 μ M dNTPs, 1.5 mM MgCl₂, 2.5 μ l of 10x PCR buffer, 2.5 μ l (1.4 M) of DMSO, and 2.5 U of Taq polymerase. The PCR was performed in an Eppendorf Mastercycler Gradient with the following cycling conditions: (i) 5 min at 95°C; (ii) 35 cycles of 30 s at 95°C, 2 min at 54-62°C (for K1F-M6R and KS α F-KS α R) and 55-60°C (for A3F-A7R), and 4 min at 72°C; and (iii) 10 min at 72°C. Amplification products were analyzed by electrophoresis in 1% (w/v) agarose gels stained with ethidium bromide and bands of 1200-1400 bp, 600-700 bp, and 700-800 bp were classified as products of *PKS-I*, *PKS-II*, and *NRPS* genes, respectively¹⁶⁻¹⁸.

TA cloning

PCR products of *PKS-I*, *PKS-II*, and *NRPS* associated with strains Iz8, F9, and F6, respectively, were selected to study the power of the optimized reaction conditions by sequencing. They were purified by gel purification kit (GeNet Bio, Korea) and cloned using the TA cloning kit (Thermo Fisher Scientific) as per the manufacturer's instructions. *Escherichia coli* (*E. coli*) strain TOP10 was used as a cloning host. Then, colony PCR was used to confirm the length of the inserted product in all clones. The plasmid was extracted from one of the positive clones in each group by plasmid extraction kit (GeNet Bio, Korea) as per the manufacturer's instructions. The purified plasmid DNAs were sequenced using universal primers M13F and M13R by Macrogen Inc. (Seoul, Republic of Korea). Sequences were tested against all DNA sequences available in GenBank database with the BLASTN program.

Identification and phylogenetic analysis of the high potential antibacterial agent-producing strains

Determination of cultural, morphological, and biochemical characteristics: The color of the sporulated aerial and substrate mycelia was examined on Starch Casein Agar (SCA) which had the following composition: 10.0 g/l soluble starch, 0.3 g/l casein, and basal mineral salt medium containing 2.0 g/l K_2HPO_4 , 2.0 g/l KNO_3 , 2.0 g/l NaCl, 0.05 g/l $MgSO_4 \cdot 7H_2O$, 0.02 g/l $CaCO_3$, 0.01 g/l $FeSO_4 \cdot 7H_2O$ (scharlau, Spain), and 15.0 g/l agar (Merck, Germany)^{19,20}. The pH of the medium was adjusted to 7.2 with the addition of HCl.

The utilization of different carbon sources (10 g/l) and degradation of starch (10 g/l)²¹ and cellulose (2 g/l)¹⁹ were studied in the basal mineral salt medium. The carbon sources were filtrated separately and added just before inoculation. Growth was compared with basal medium without carbon source as the negative control. The following carbon sources were tested: glucose, fructose, lactose, mannitol, raffinose, sorbitol, sucrose, and xylose. Proteolytic activity was determined using milk casein (plate) and gelatin (in tubes). Skimmed milk agar plates contained 5.0 g/l peptone, 3.0 g/l yeast extract, 1.0 g/l skimmed milk, and 12.0 g/l agar, pH=7.0²². The gelatin hydrolysis test was performed on nutrient gelatin containing 100-150 g/l gelatin^{23,24}, final pH=7.2. Urease test media contained 1.0 g/l peptone, 1.0 g/l glucose, 5.0 g/l NaCl, 2.0 g/l KH_2PO_4 , 0.012 g/l phenol red, 20.0 g/l urea, and 15.0 g/l agar. The pH of the medium was adjusted to 6.8 with the addition of HCl. The urea was filtered and added to the rest of the medium²⁵.

Molecular identification

The partial length sequence of 16S rDNA gene was amplified from the isolated genomic DNAs using the following two sets of universal bacterial 16S rDNA primers, "27F-YM (5'-AGAGTTTGTATYMTGGCTCAG-3')^{26,27} and 1492R (5'-TACGGYTACCTTGTTACGACTT-3')²⁸" and "16SP0 (5'-GAAGAGTTTGATC

CTGGCTCAG-3') and 16SP6 (5'-CTACGGCTACCTTGTTACGA-3')"²⁹. A 50 μ l sample of the PCR reaction mixture contained 10-100 ng of genomic DNA, 0.4 μ M of each primer, 200 μ M dNTPs, 1.5 mM $MgCl_2$, 5 μ l of 10x PCR buffer, 5 μ l (1.4 M) of DMSO, and 2.5 U of Taq polymerase, all obtained from Cinnagen Inc, Iran. The PCR was performed in an Eppendorf Thermal Cycler using appropriate programs optimized for each primer pair after denaturation of DNA by heating for 4 min at 94°C. The PCR program involved 35 cycles, and each cycle consisted of denaturation at 94°C for 1 min, annealing at 55°C for 1 min, extension at 72°C for 2 min for "27F-YM/1492R"²⁸, and denaturation at 95°C for 30 s, annealing at 55°C for 30 s, extension at 72°C for 2 min for "16SP0/16SP6"²⁹. This was followed by a final elongation step for 5 min at 72°C. Moreover, we tested colony PCR without prior extraction of DNA as the template for rapid and simple detection of DNA sequences according to the method of Ishikawa *et al.*³⁰ The PCR products were separated on a 1% agarose gel containing ethidium bromide in 1xTBE buffer. The purified PCR product was sequenced in both directions using an automated sequencer by Macrogen.

Phylogenetic analysis of 16S rDNA gene sequences

The sequences were edited using Finch TV V.1.4.0., then aligned with sequences from the closest related species of the genus *Streptomyces* retrieved from the GenBank/EzTaxon databases using PhyDE. The alignment length was 1402 characters (including indels). Phylogenetic relationships were studied through Bayesian inference using MrBayes version 3.2.6³¹. JModeltest version 2.3³² was used to find the best model of molecular evolution, and TIM1ef+I+G model was selected according to Akaike Information Criterion (AIC). Two parallel runs of four MCMC (Markov chain Monte Carlo) chains, including three heated chains and one cold chain, were carried out simultaneously for 8 million generations with a sample frequency of 1000. The analyses were stopped when the standard deviation of split frequencies was well below 0.01. The first 25% of generations were set to burn-in. Trees were visualized using TreeGraph2³³.

Results

Study of the presence of *PKS-I*, *PKS-II*, and *NRPS* genes in twenty-two antibacterial agent-producing actinomycete strains

Twenty-two antibacterial agent-producing actinomycete strains (F4, F6, F9, F10, Is1, Is4, Is5, Ih1, Ih6, Iz1, Iz2, Iz6, Iz8, Iz9, Iz11, Iz14, Iz16, Iz17, D2, B4, N1, and A2) isolated in our previous study¹³ (Table 1, Figure 1) were screened for the presence of *PKS-I*, *PKS-II* and *NRPS* gene sequences by specific amplification of chromosomal DNA with primer sets K1F/M6R, KS α F/KS α R, and A3F/A7R.

At first, multiple bands were found after PCR amplification of *PKS-I* genes with the annealing tempera-

Table 1. Antibacterial activity of some actinomycete strains isolated from soil samples of Iran

The name of strain	S	Antibacterial activity ^a against					R	Ca	Cg
		B	E	M	V	A			
F4	++	++	-	++	-	-	++	+	-
F6	-	+++	-	+++	-	-	-	-	-
F9	+++	+++	-	+++	+++	-	-	-	-
F10	+++	++	-	++	++	-	-	-	-
Is1	-	+	-	-	-	-	++	-	-
Is4	+++	+	-	-	-	-	++	-	-
Is5	+++	++	+	-	-	-	-	-	-
Ih1	++	-	-	-	-	-	-	-	-
Ih6	+++	+++	-	-	-	-	-	-	-
Iz1	+++	+	-	-	-	-	-	-	-
Iz2	+++	-	-	-	-	-	-	-	+
Iz6	++	+	-	++	++	-	-	-	-
Iz8	+	++	-	+	+++	+	-	-	-
Iz9	+	++	-	-	-	-	-	-	-
Iz11	-	+++	-	+++	-	-	-	-	-
Iz14	+++	+	-	-	-	-	-	-	-
Iz16	+++	+	-	+++	-	-	++	-	-
Iz17	++	+	-	+++	-	-	+	-	-
D2	+++	+++	-	-	-	-	-	-	-
B4	+	+	-	-	+	-	-	-	-
N1	++	-	-	-	-	+	-	-	-
A2	+	++	-	-	-	-	-	-	-

a: Estimated by measuring the mean diameter of the clear zone of growth inhibition obtained from triplicate. Symbols: S, *Staphylococcus aureus* (ATCC 6538); B) *Bacillus subtilis* (ATCC 12711); E) *Escherichia coli* (ATCC 1399); M) methicillin-resistant *Staphylococcus aureus*; V) vancomycin-resistant enterococci; A) *Aspergillus fumigatus*; R, *Rhizopus* sp.; Ca, *C. albicans*; Cg, *C. glabrata*; -, no activity; +, weak activity; ++, moderate activity and +++, strong activity.

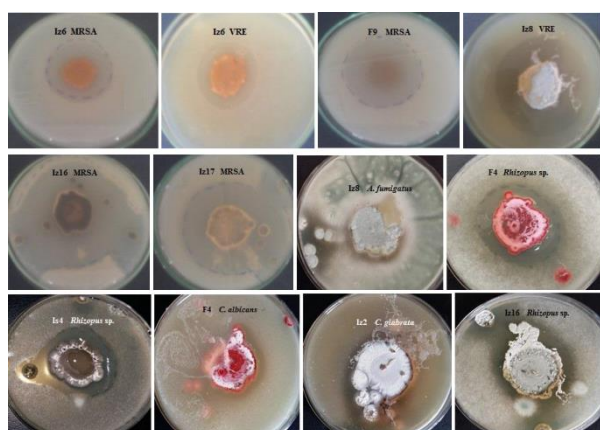


Figure 1. The antimicrobial activity of several actinomycete strains against hospital-acquired bacterial and fungal isolates.

ture estimated in previous researches ¹⁶⁻¹⁸ (Figure 2A only shows the results of some strains). Therefore, we tried to find the optimal annealing temperature for K1F/M6R primer set.

The optimal annealing temperature was variable for *PKS-I* depending on template DNA. It was: 55°C for Iz1, Iz11, Iz17, and Is1; 57°C for Iz2, Iz8, Iz9, Iz14, Is4, and Ih1; 58°C for B4 and D2; 59°C for F4, F9, and F10; 60°C for A2, Ih6, and N1, 61°C for F6 and Iz16; and 62°C for Iz6 and Is5. Concerning A2, all sub-bands were removed with the annealing temperature increased from 57 to 60°C. However, in some cases, we couldn't get a single band at the mentioned temperatures, but as can be seen for Iz6, Iz16, F6, N1, and Ih6, the majority of non-specific bands with variable lengths were eliminated, and the rest were reduced to the minimum sharpness. (Figures 3A and 3B).

The optimal annealing temperature was 58°C for *PKS-II* because there was no shift from negative to positive at lower annealing temperatures, leading to non-specific DNA products (Figure 2B only shows the results of some strains). A comparison between Figures 2B, 3C and 3D shows that PCR amplification of F10, Iz6, and Iz8 at 58°C gives a single band on 600-700 bp while at 55°C gives multiple bands at different lengths. For *NRPS* genes, in addition to the specific band, a

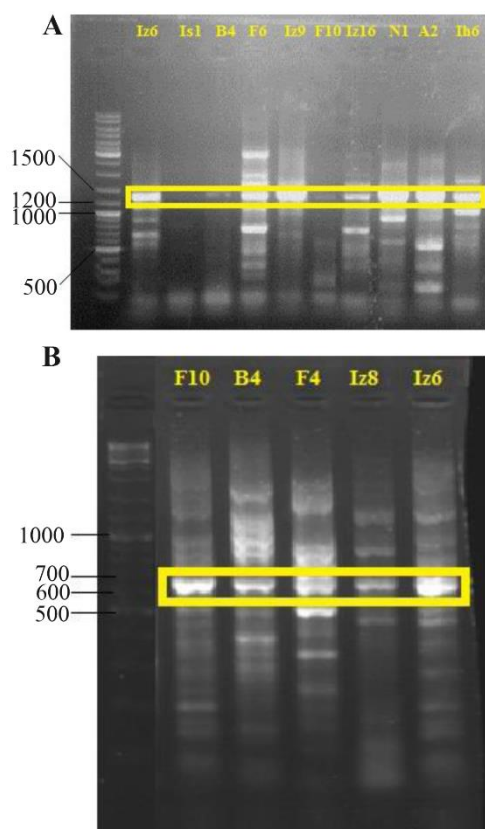


Figure 2. PCR amplification of PKS-I (A) and PKS-II (B) gene sequences with an amplicon size of 1200-1400 and 600-700 bp, respectively, for a number of strains at 57 and 55°C, respectively. The name of strains has mentioned at the top of gel.

non-specific band was consistently observed at about 400-500 bp, and there was no change in the quality of the PCR output in the temperature range of 55-60°C. Some sequences were amplified at 55 up to 60°C, while some were amplified only at 55°C. Therefore, 55°C was the optimal annealing temperature for the degenerate primer set A3F/A7R (Figures 3E and 3F). Another factor that affects the success of PCR is primer concentration. The optimal concentrations to remove the false-negative results were 2, 0.8, and 2 μ M for amplification of *PKS-I*, *PKS-II*, and *NRPS* genes, respectively. Actually, the results shifted from negative to positive or weak to strong by increasing primer concentrations from 0.8 to 2 μ M for *PKS-I* and *NRPS* and from 0.4 to 0.8 μ M for *PKS-II*. It is important to note that the number of non-specific bands sometimes increases with an excessive primer concentration. The minimum required amount of DNA that serves as the initial template or target sequence was 20-30 ng, and a higher amount did not have any marked effect on the outcome of PCR. Among investigated strains, 19 (86.4%), 18 (81.8%), and 21 (95.4%) strains were positive for *PKS-I*, *PKS-II*, and *NRPS* genes, respectively, after optimization (Table 2, Figure 3).

Table 2. The presence of PKS-I, PKS-II and NRPS gene groups in twenty-two antimicrobial agent-producing *Streptomyces* strains

The name of strain	Presence ^a of gene		
	PKS-I	PKS-II	NRPS
F4	+	+	+
F6	+	+	+
F9	-	+	+
F10	+	+	+
Is1	+	-	+
Is4	+	-	+
Is5	+	-	+
Ih1	+	+	+
Ih6	+	+	+
Iz1	+	+	+
Iz2	-	+	+
Iz6	+	+	+
Iz8	+	+	+
Iz9	+	+	-
Iz11	+	+	+
Iz14	-	+	+
Iz16	+	+	+
Iz17	+	+	+
D2	+	-	+
B4	+	+	+
N1	+	+	+
A2	+	+	+

a: +, present; -, absent.

Sequence analysis of *K1F/M6R*, *KSaF/KSaR*, and *A3F/A7R* PCR products

The clones selected from strains Iz8, F9, and F6 had high sequence similarity (85, 99, and 92%, respectively) with *PKS-I* gene on plasmid pSPA1 in *Streptomyces parvulus* (*S. parvulus*) strain 2297, the first *PKSII* gene previously reported in *Streptomyces* sp. strain MM1, and *NRPS* gene cluster in *Streptomyces pactum* (*S. pactum*) strain ACT12. These sequences were deposited in the NCBI GenBank database under the accession numbers KY637058, KY637060, and KY637057. The sequencing results confirmed the optimized conditions for specialized PCR of *PKS-I*, *PKS-II*, and *NRPS* gene groups.

Identification of morphological, biochemical, and molecular characteristics and assessment of phylogenetic relationships of eight actinomycete strains with high antimicrobial activity (F4, F6, F9, F10, Iz6, Iz8, Iz16, and B4)

The ability of strains F4, F6, F9, F10, Iz6, Iz8, Iz16, and B4 to utilize various carbon sources for growth, proteolyze casein and gelatin, and degrade urea, was tested. All of them used glucose and starch, while none was able to use sorbitol and cellulose as the sole carbon source for growth. Every eight strains were urease-positive. "F4 and Iz8" and "F10 and Iz6" were able to degrade the same carbon sources (Table 3).

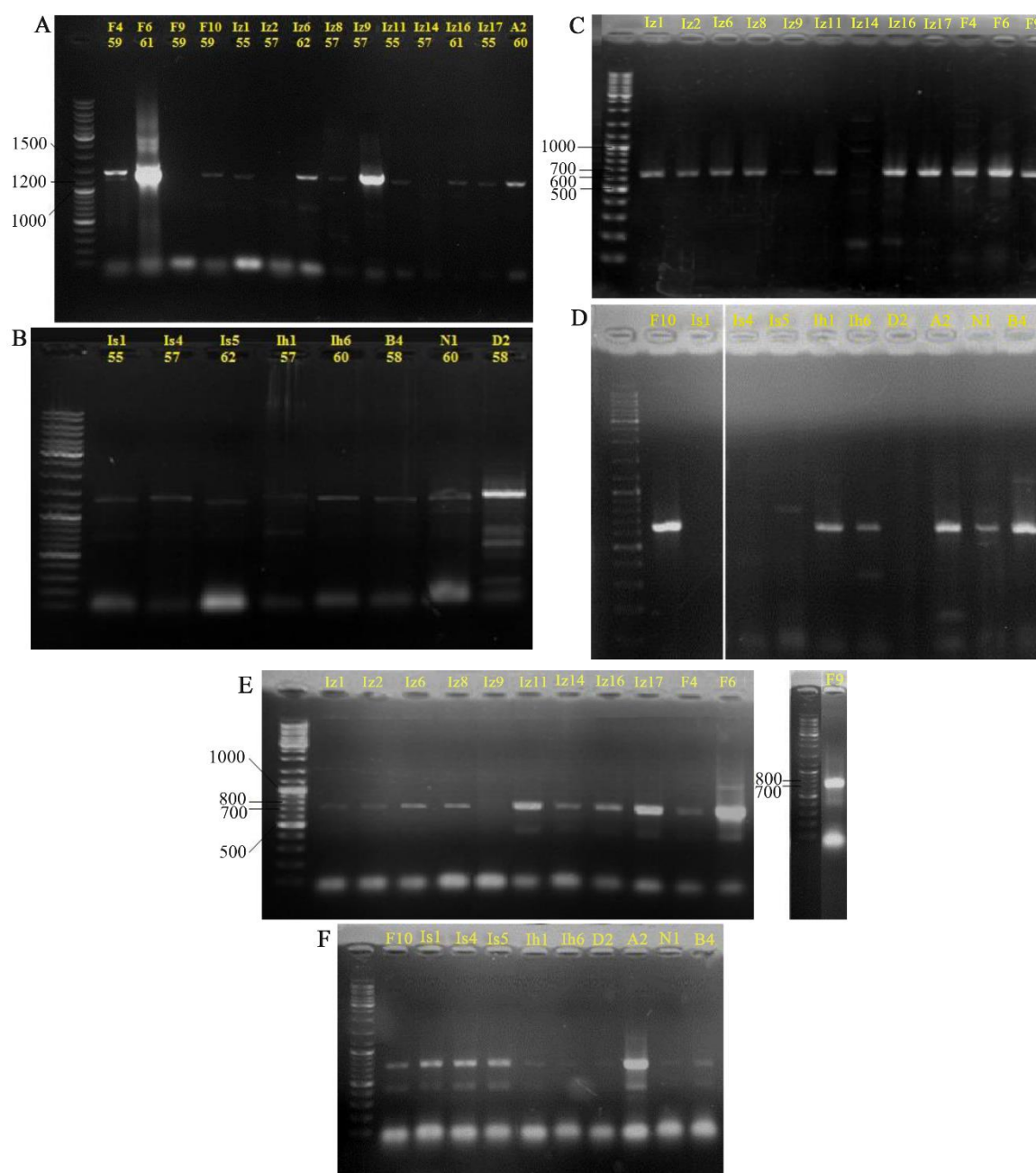


Figure 3. PCR amplification of PKS-I (A and B), PKS-II (C and D) and NRPS (E and F) gene sequences with an amplicon size of 1200-1400, 600-700 and 700-800 bp using primers K1F/M6R, KSaF/KSaR and A3F/A7R, respectively, on agarose gel 1%. The name of strains has mentioned at the top of gel.

As shown in figure 4, amplification of 16S rDNA gene gives ≈ 1500 bp product using each of the two primer sets (six strains B4, F4, F6, F9, Iz6, and Iz16 using 27F-YM/1492R and two strains F10 and Iz8 using SP0/SP6). 16S rDNA gene of strains F10 and Iz8 had a very weak amplification by 27F-YM/1492R primer set, while, no product was observed by colony PCR.

Nearly full-length 16S rDNA gene sequences of all eight strains were compared with 16S rDNA sequences

of type strains retrieved from GenBank/EzTaxon. All of them were assigned to genus *Streptomyces* and were deposited in the NCBI GenBank database under the accession numbers KX229767-73 (strains B4, F4, F6, F10, Iz6, Iz8 and Iz16, respectively) and KX417085 (strain F9) (Table 3).

Moreover, the alignment of 16S rDNA sequences showed a very high degree of similarity and conservation among the isolated strains. It was one of the most important reasons for using MrBayes instead of Mega

Optimization of Degenerate PCR Conditions

Table 3. Morphological, physiological and biochemical characteristics and molecular identification of the high potential antibacterial agent-producing strains

Strain	Closest type strain	Sequence identity (%)	Carbone utilization							Degradation		Protease			Urease	Color	
			Glucose	Fructose	Lactose	Mannitol	Raffinose	Sorbitol	Sucrose	Xylose	Starch	Cellulose	Casein	Gelatin		Aerial mycelia	Substrate mycelia
B4	<i>S. tauricus</i>	99.57	+	-	-	+	+	-	-	-	+	-	+	-	+	red	dark orange-red
F4	<i>S. rectiviolaceus</i>	99.14	+	+	+	+	+	-	+	+	+	-	-	+	+	light red-orange	dark red
F6	<i>S. marokkonensis</i>	99.49	+	+	+	+	+	-	-	+	+	-	+	-	+	gray	blackish gray
F9	<i>S. chryseus</i>	99.88	+	-	-	-	-	-	+	-	+	-	+	-	+	dark brown with white dots	dark brown
F10	<i>S. galilaeus</i>	100	+	+	+	-	+	-	+	-	+	-	-	-	+	orange-light green	light orange
Iz6	<i>S. bobili</i>	99.93	+	+	+	-	+	-	+	-	+	-	-	-	+	orange	orange
Iz8	<i>S. coerulescens</i>	100	+	+	+	+	+	-	+	+	+	-	-	-	+	blue	yellow
Iz16	<i>S. flavoviridis</i>	99.29	+	+	+	+	-	-	+	+	+	-	+	-	+	green with white dots	green

+, positive result; -, negative result

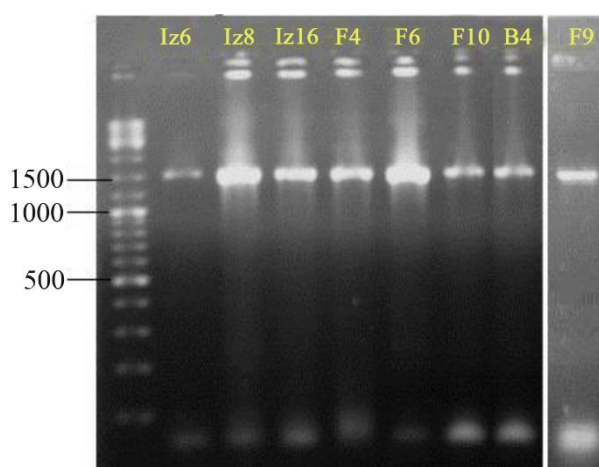


Figure 4. Agarose gel electrophoresis of 16S rDNA gene PCR-amplified region. The ladder includes fragments ranging from 100-10,000 bp. The name of strains has mentioned at the top of gel.

to draw a phylogenetic tree. Phylogenetic analysis was performed by Bayesian inference implemented in Mr-Bayes based on the partially complete 16S rDNA gene sequences of eight selected strains and the type strains of closely related species (Figure 5). They were divided into four separate clades. Three strains of F9, F4, and B4 were placed in distinct clades of I, III, and IV, respectively, and five strains of F6, Iz16, Iz8, F10, and Iz6 were in a joint clade (clade II). Strain F9 and four

strains, including *Streptomyces chryseus* (*S. chryseus*), *Streptomyces helveticus* (*S. helveticus*), *S. albidocromogenes*, and *Streptomyces albosporeus* (*S. albosporeus*), formed a polytomic subclade within clade I, which was moderately supported (posterior probability=0.75). Clade II contained several differential subclades. Strain F6 was one of these subclades. Strains Iz8 and Iz16 are branched together on one sub-branch, while Iz6 and F10 are branched together on another with the posterior probability of 0.77 and 0.83. "Strain F4 and type strain of *Streptomyces rectiviolaceus* (*S. rectiviolaceus*)" and "strain B4 and type strain of *Streptomyces tauricus* (*S. tauricus*)" were in an internal subclade of clades III and IV, respectively, with the posterior probability>0.8.

Discussion

NRPS and PKS pathways can be detected in many actinomycete isolates as the characteristics of secondary metabolite production¹. Prescreening of isolates with PCR primers targeting bioactive compounds-encoding genes (*PKS-I*, *PKS-II*, *NRPS*, etc.) is an effective and practical approach for detecting novel and useful secondary metabolites³⁴. On the other hand, the PCR detection of these genes is vital to discover isolates with high potential for the production of valuable secondary metabolites because some of the secondary metabolite biosynthetic gene clusters appear to be non-active or silent under standard culture conditions, and

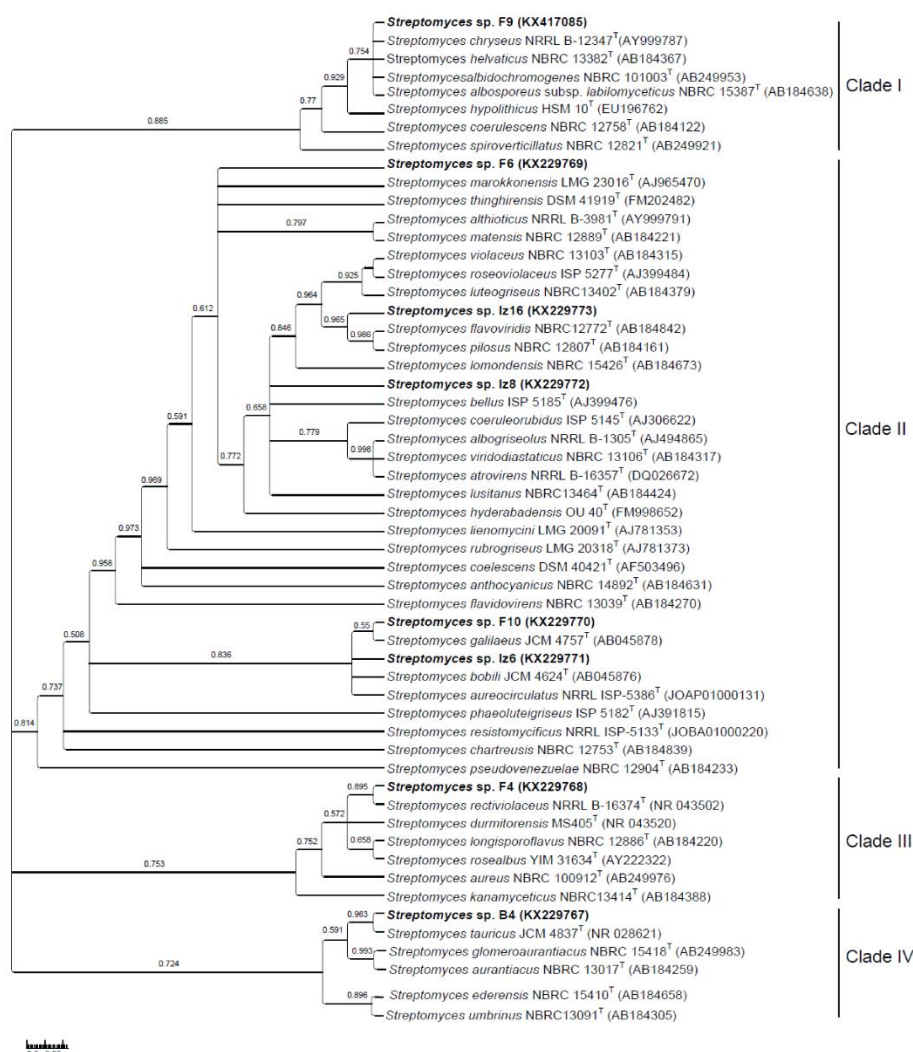


Figure 5. Bayesian inference tree showing the phylogenetic relationship among the eight selected strains and the type strain species of *Streptomyces* with the highest 16S rDNA sequence similarity. Posterior probabilities for the nodes are shown under the branches.

they may require some specific triggers to be activated. From these, it can be concluded that the nonbioactive isolates hold their genetic capability for producing valuable secondary metabolites if they are cultivated under appropriate conditions¹. Therefore, it would be reasonable to utilize the screening of *PKS* and *NRPS* systems in combination with antimicrobial activity assay for biotechnological applications¹⁸. Several research studies have been conducted and reported on the presence of *PKS* and *NRPS* genes as the characteristics of secondary metabolite production in Actinobacteria. It is noted that the low detection of *PKS* (types I and II) and *NRPS* gene fragments in the antibacterial agent-producing strains may reflect the lack of targeted genes or the implication of other genes in the biosynthesis of these bioactive products. Further, this may be caused by the types of selected degenerate primer pairs, which

are unsuitable for amplifying these genes^{2,5,18}. Amos *et al*³⁵ validated two new updated PCR assays using new degenerate primers and concluded that these assays are specific, can detect clusters from a wide range of phyla, and have a good and reasonable hit rate.

To account for the discrepancies and inappropriate PCR conditions, twenty-two strains were screened for the presence of *PKSI*, *PKS-II*, and *NRPS* gene sequences by specific amplification of chromosomal DNA with primer sets K1F/M6R, KSaF/KSaR, and A3F/A7R¹⁶⁻¹⁸. Actually, in the previous researches, PCR was performed at the fixed annealing temperature of 67.5°C³⁶ for A3F/A7R, 56°C³ or 57°C^{1,2,17,18,34} for K1F/M6R and A3F/A7R and 57°C¹, 58°C^{2,17,18,34} or 63°C³ for KSaF/KSaR and KSaF/KSβR. Lee *et al*¹ used the annealing temperature of 62°C for A3F/A7R. In contrast, our analysis showed that PCR amplifica-

tion of the KS domain belonging to *PKS-I* using the degenerate primer set K1F/M6R needs different annealing temperatures depending on the template DNA. This is not inconsistent with the fact that some strains do not have this gene but may raise the possibility of getting the actual positive results. Moreover, some sequences can amplify only at a lower annealing temperature (55°C) by primer set A3F/A7R.

Another factor that affects the success of PCR is primer concentration, as sometimes the low concentration might lead to an undetectable product giving a false negative result. The optimal concentrations were 2 μ M for *PKS-I* and *NRPS* and 0.8 μ M for *PKS-II*, while previous researchers used primers at the final concentration of 0.2-0.8 μ M^{1-3,17,18,35}. The dependence of amplification efficiency on the template amount was low. In this study, 14 strains (64%) contained all of the studied genes, and 100% of strains had at least two of the target genes. These numbers are quite distinct in comparison with the previous researches: 19.5, 42.5, and 5.7¹; 34.1, 63.4, and 61²; 10.9, 15.2, and 26.1³⁴; 11.5, 23, and 76.9¹⁷; 46.6, 60, and 46.6¹⁸; 79.6, 70.4, and 57.4; and 65.6, 45.9, and 34.4³ for *PKS-I*, *PKS-II* and *NRPS* genes, respectively.

The sequencing results indicated the high presence of genus *Streptomyces* compared to other actinomycetes in our soil samples. However, it should be noted that starch casein agar, containing starch as the sole carbon source, was the only medium used to separate these actinomycete strains¹³. It is well accepted that members of the genus *Streptomyces* are prolific producers of novel antibiotics, with approximately 75% of commercially valuable antibiotics being derived from *Streptomyces*¹.

Conclusion

The screening of *PKS* and *NRPS* genes is a quick way to show the potential for antimicrobial agent production. However, errors in laboratory techniques sometimes can make us move away from reality. The optimization of degenerate PCR will help us reduce error rates in the lab. Based on the results obtained in the present study, it can be concluded that degenerate PCR strongly depends on the annealing temperature and primer concentration, as any changes in these parameters can result in an unexpected shift in PCR outputs. Actually in degenerate PCR, we are dealing with different primers, and naturally, they have different annealing temperatures. For example, the primer set KSaF/ KSaR can be combined in 192 modes, and the annealing temperature will be different depending on which F and R primers are matched with DNA. Moreover, in some cases, bacteria in one common clade produce identical antibiotics. The phylogenetic tree helps us to choose different antibiotic-producing strains and avoid the purification of similar antibiotics by classifying the strains.

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Conflict of Interest

The authors declare that they have no conflict of interest.

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