A report of 34 unrecorded bacterial species in Korea, belonging to the Actinobacteria

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As a subset study to discover indigenous prokaryotic species in Korea in 2014, a total of 34 bacterial strains assigned to the phylum Actinobacteria were isolated from various environmental samples collected from activate sludge, biotite, freshwater, gut of marine organisms, mud flat, sediment, soil, spent mushroom compost and sea water. On the basis of high 16S rRNA gene sequence similarity and a tight phylogenetic association with the closest species, it was revealed that each strain was assigned to independent and previously described bacterial species, with the exception of one isolate. There is no official report that these 34 species included in the phylum Actinobacteria have been described in Korea: 6 species of 5 genera in the order Corynebacteriales, 1 species of 1 genus in the order Frankiales, 2 species of 2 genera in the Micromonosporales, 14 species of 10 genera in Micrococcales, 2 species of 2 genera in the Propionibacteriales, 1 species of 1 genus in the Pseudonocardiales, 4 species of 2 genera in the Streptomycetales, 2 species of 2 genera in the Streptosporangiiales and 1 species of 1 genus in the Solirubrobacterales. Gram reaction, cell and colony morphology, pigmentation, physiological characteristics, isolation sources and strain IDs are described in the section of species description.

Keywords: 16S rRNA gene, Actinobacteria, indigenous prokaryotic species, unrecorded species

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INTRODUCTION

The recently proposed phylum Actinobacteria (Ludwig et al., 2012) is composed of aerobic, Gram-staining-positive bacteria that have high-G+C contents in their genomic DNAs. They exhibit a very wide range of morphology and form cocci, rods or fragmented or branched hyphae depending on species. Some of them produce resting structure such as spores and sporangium on permanent, well-developed mycelium (Goodfellow and Cross, 1984; Ensign, 1992).

The phylum Actinobacteria is well supported by analyses of the 16S and 23S rRNA, presence of conserved insertions and deletions in certain proteins, and characteristic gene rearrangements (Goodfellow and Fiedler, 2010) and encompasses six classes Actinobacteria, Acidimicrobiia, Coriobacteria, Nitriliruptoria, Rubrobacteria and Thermoleophilia (Ludwig et al., 2012).

The class Actinobacteria is composed of 15 orders Actinomycetales, Actinopolysporales, Bifidobacteriales, Catenulisporales, Corynebacteriales, Frankiales, Glycomycetales, Jiannellales, Kineosporiales, Micromonosporales, Micrococcales, Propionibacteriales, Pseudonocardiales, Streptomycetales and Streptosporangiiales. Members of this class have been received considerable
concerns as the source of antibiotics since the discovery of actinomycin from actinomycetes (Waksman and Woodruff, 1940) and widely distributed in natural environment such as soil, fresh or sea water, manure and compost where contribute significantly to turnover of complex biopolymers as the decomposer (Williams et al., 1984).

The class Thermoleophilia contains two orders Solirubrobacterales and Thermoleophilales; the former consists of the families Solirubrobacteraceae, Conexibacteraceae and Patulibacteraceae, while the latter comprises the family Thermoleophilaceae (Reddy and Garcia-Pichel, 2009; Ludwig et al., 2012).

We collected diverse environmental samples for bacterial isolation and recovered a great number of indigenous bacterial species in Korea by the research program supported by NIBR (National Institute of Biological Resources) in 2014. The aim of present study is to deal with the classification and identification of bacterial strains assigned to the phylum Actinobacteria which have not been previously reported in Korea and to describe 34 unrecorded bacterial species belonging to 9 orders of the two classes Actinobacteria and Thermoleophilia.

**MATERIALS AND METHODS**

A total of 34 bacterial strains which were assigned to the phylum Actinobacteria were isolated from various environmental samples collected from activate sludge, biotite, freshwater, gut of marine organisms, mud flat, sediment, soil, spent mushroom compost and sea water (Table 1). Treatment of environmental samples and bacterial isolation was done independently in several laboratories. The pure cultures of isolated bacteria were grown on diverse culture media including R2A agar (TSA; BD), nutrient agar (NA; BD), ISP (International Streptomyces Project) 2 and 4 media (Shirling & Gottlieb, 1966) at 15-30°C for 2-7 day, depending on the strains. The designated strain IDs, isolation sources, culture media, and incubation conditions are summarized in Table 1. The pure cultures were maintained as 10-20% glycerol suspensions at -80°C and lyophilized ampoules.

Cell morphology was observed by either transmission or scanning electron microscopy. Gram staining was performed using a Gram-staining kit according to the instructions of the manufacturer. Colony morphology and pigmentation were observed on agar plates with cells grown to stationary phase. Utilization of carbon sources and some biochemical properties were examined by using API 20NE galleries (bioMérieux) according to the manufacturer’s instructions.

Bacterial DNA extraction, PCR amplification and sequencing of the 16S rRNA gene were performed using the standard procedures described elsewhere. The 16S rRNA gene sequences of the strains assigned to the phylum Actinobacteria were compared with the corresponding sequences collected from and the EzTaxon-e server (Kim et al., 2012) and public database. Multiple alignments of the sequences were performed using the Clustal_X program (Thompson et al., 1997) and optimized manually according to the secondary structure of Escherichia coli. Evolutionary distances were calculated using the correction of Jukes & Cantor (1969). Phylogenetic analyses were performed by using the neighbor-joining (Saitou and Nei, 1987), maximum-parsimony (Fitch, 1971) and maximum-likelihood (Felsenstein, 1981) treeing algorithms contained in the PHYLIp package (Felsenstein, 2008). A neighbour-joining tree was drawn with bootstrap values based on 1,000 replications (Felsenstein, 1985).

**RESULTS AND DISCUSSION**

The 34 strains were found to belong to the two classes Actinobacteria (33 strains) and Thermoleophilia (1 strains) in the phylum Actinobacteria. Among them, 33 strains were distributed in 6 orders of the Actinobacteria; 6 strains for the order Corynebacteriales, 1 strain for the Frankiales, 2 strains for the Micromonosporales, 14 strains for the Micrococcales, 2 strain for the Propionibacteriales, 1 strain for the Pseudonocardiales, 5 strains for the Streptomycetales and 2 strains for the Streptosporangiales. One strain which belonged to the class Thermoleophilia was assigned to the Solirubrobacterales (Table 1). These strains were Gram-staining-positive and chemoheterotrophic and were morphologically characterized by the formation of cocci, rods or mycelia (Fig. 1). The strains of the order Corynebacteriales (Fig. 2) were assigned to 5 genera of 3 families: Dietzia (1 species) the family Dietziaceae, Mycobacterium (2 species) of the Mycobacteriaceae, Gordonia (1 species), Nocardia (1 species) and Rhodococcus (1 species) of the Nocardiaceae. In general, members of the genus Nocardia are morphologically characterized by fragmentation of mycelium, in contrast to other genera of the family Nocardiaceae having coccoid- or rod-shaped morphology (Goodfellow and Maldonado, 2012). The Frankiales strain (Fig. 2) was affiliated to the genus Geodermatophilus (1 species) of the family Geodermatophilaceae. The 2 Micromonosporales strains were distributed to two genera Actinocatenisiopa (1 species) and Micromonospora (1 species) of the family Micromonosporaceae, which were morphologically characterized by the formation of mycelia (Fig. 1). The phylogenetic relationships between the Micrococcales strains and their closest relatives are...
Table 1: Summary of the isolated strains belonging to the Actinobacteria and their taxonomic affiliations.

<table>
<thead>
<tr>
<th>Order</th>
<th>Family</th>
<th>Genus</th>
<th>Strain ID</th>
<th>Most closely related species (Name of type strain)</th>
<th>% Similarity</th>
<th>Isolation conditions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Actinobacteria</td>
<td>Corynebacteriales</td>
<td>Dietzia</td>
<td>IMCC 12370</td>
<td>Dietzia schimae</td>
<td>99.8%</td>
<td>Seawater, MA, 15℃, 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Mycobacteriales</td>
<td>WL1</td>
<td>Mycobacterium septicum</td>
<td>99.7%</td>
<td>Freshwater, R2A, 25℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>WS80</td>
<td>Mycobacterium hodleri</td>
<td>99.2%</td>
<td>Freshwater, R2A, 25℃, 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>M13</td>
<td>Nocardia niigatensis</td>
<td>99.7%</td>
<td>Freshwater, R2A, 25℃, 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>NU 4Y-9-1</td>
<td>Nocardia niigatensis</td>
<td>100%</td>
<td>Ginseng soil, TSA, 30℃, 3 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>YB.Ce-3</td>
<td>Rhodococcus phenolicus</td>
<td>98.9%</td>
<td>Activated sludge, R2A, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>BS8</td>
<td>Actinocatenispora sera</td>
<td>99.0%</td>
<td>Spent mushroom compost, R2A, 30℃, 2 days</td>
</tr>
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<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>61DPE39</td>
<td>Micromonospora siamensis</td>
<td>99.1%</td>
<td>Freshwater, R2A, 25℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>NK 6Y-6-4</td>
<td>Arthrobacter casei</td>
<td>100%</td>
<td>Activated sludge, R2A, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>14-10</td>
<td>Arthrobacter pascens</td>
<td>99.3%</td>
<td>Ginseng soil, R2A, 30℃, 2 days</td>
</tr>
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<td></td>
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<td>Nocardiaceae</td>
<td>K18</td>
<td>Sinomonas flava</td>
<td>99.9%</td>
<td>Ginseng soil, R2A, 30℃, 2 days</td>
</tr>
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<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>KS904</td>
<td>Kribbella antibiotica</td>
<td>99.4%</td>
<td>Ginseng soil, R2A, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>RK 4Y-2-4</td>
<td>Kribbella antibiotica</td>
<td>99.5%</td>
<td>Ginseng soil, ISp4, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>RMD 3Y-3-1</td>
<td>Kutzneria buriramensis</td>
<td>99.5%</td>
<td>Ginseng soil, ISp4, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>Nu 4Y-9-4</td>
<td>Kitasatospora paranensis</td>
<td>99.2%</td>
<td>Freshwater, ISp4, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>BBT1</td>
<td>Streptomyces drozdowiczii</td>
<td>99.0%</td>
<td>Powder of biotite, R2A, 30℃, 2 days</td>
</tr>
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<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>BS22</td>
<td>Streptomyces thermocoprophilus</td>
<td>98.9%</td>
<td>Spent mushroom compost, R2A, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>I1-6</td>
<td>Streptomyces olivochromogenes</td>
<td>99.0%</td>
<td>Soil, ISp2, 30℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nocardiaceae</td>
<td>T1-6</td>
<td>Streptomyces olivochromogenes</td>
<td>99.1%</td>
<td>Soil, ISp2, 30℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>WM35</td>
<td>Streptosporangium amethystogenes</td>
<td>99.3%</td>
<td>Freshwater, R2A, 25℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>7C-18</td>
<td>Actinomadura bangladeshensis</td>
<td>99.3%</td>
<td>Activated sludge, R2A, 30℃, 2 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>3</td>
<td>Thermoleophila composita</td>
<td>99.1%</td>
<td>Spent mushroom compost, R2A, 30℃, 4 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>4Y-1-6</td>
<td>Conexibacter arvalis</td>
<td>99.3%</td>
<td>Spent mushroom compost, R2A, 30℃, 4 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>9-3</td>
<td>Streptomyces thermocyanogenes</td>
<td>99.1%</td>
<td>Freshwater, ISp4, 30℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>4Y-2-4</td>
<td>Streptomyces thermocyanogenes</td>
<td>99.2%</td>
<td>Freshwater, ISp4, 30℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>BBT1</td>
<td>Streptomyces thermocyanogenes</td>
<td>99.3%</td>
<td>Freshwater, ISp4, 30℃, 7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>BS10</td>
<td>Streptomyces thermocyanogenes</td>
<td>99.8%</td>
<td>Spent mushroom compost, R2A, 30℃, 4 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Streptosporangiaceae</td>
<td>BS15</td>
<td>Streptomyces thermocyanogenes</td>
<td>99.8%</td>
<td>Spent mushroom compost, R2A, 30℃, 4 days</td>
</tr>
</tbody>
</table>
Fig. 1. Transmission electron micrographs or scanning electron micrographs of cells of the strains isolated in this study. Strains: 1, IMCC 12370; 2, WL1; 3, WS80; 4, MN13; 5, NU 4Y-9-1; 6, YB.Ce-3; 7, WM99; 8, BS8; 9, 61DPR39; 10, Ho-02; 11, RMD 3Y-15-4; 12, HMF2762; 13, NK 4Y-9-3; 14, WW28; 15, EgT0207; 16, KHS04; 17, NK 6Y-6-4; 18, NS 4Y-8-4; 19, 145-10; 20, AX5; 21, NGS 3Y-15-2; 22, R1-6; 23, N1-9; 24, RK 4Y-2-4; 25, RS 4Y-2-4; 26, RMD 3Y-3-1; 27, NU 4Y-9-4; 28, BBT1; 29, BS22; 30, 11-6; 31, T1-6; 32, 7C-18; 33, WM35; 34, BS10.
given in Fig. 3. The 14 strains belonged to the 9 genera of 7 families: the genus *Brevibacterium* (2 species) of the family *Brevibacteriaceae*, *Cellulomonas* (3 species) of the *Cellulomadaceae*, *Brachybacterium* (1 species) of the *Dermabacteraceae*, *Ornithimicrobium* (1 species) of the *Intrasporangiaceae*, *Arthrobacter* (2 species), *Kocuria* (1 species), *Micrococcus* (1 species) and *Sinomonas* (1 species) of the *Micrococcaceae*, *Diaminobutyricibacter*
(1 species) of the Microbacteriaceae and Luteimicrobium (1 species) of the Promicromonosporaceae. Among them, the strain belonging to the family Intrasporangiaceae showed low 16S rRNA gene sequence similarity (96.20%) to the closest species Ornithinimicrobium murale (Table 1), suggesting that this isolate represents a novel taxon at the taxonomic ranks of species and genus.

Fig. 4 shows phylogenetic affiliations of the 10 strains of the orders Propionibacteriales, Pseudonocardiales, Streptomycetales and Streptosporangiiales. The 2 strains belonged to the orders Propionibacteriales; each strain was assigned to the genera Kribbella and Nocardioides of the family Nocardioidaceae, respectively. The order Pseudonocardiales contained only 1 strain assigned to the
The order Streptomyces consisted of 5 isolates that were affiliated to the genera Kitasatospora (1 species) and Streptomyces (3 species) of the family Streptomycetaceae, some of which formed branched mycelia (Fig. 1). Members of the family Streptomycetaceae are morphologically characterized by the formation of well-developed, branched hypha. Most of them further produced the chains of spores with the various surface ornamentations borne on the tips of branched mycelia (Kämpfer, 2012a; 2012b). The 2 strains of the order Streptosporangiales were assigned to the genus Actinomadura of the family

Fig. 4. Neighbor-joining phylogenetic tree, based on 16S rRNA gene sequences, showing the relationships between the strains isolated in this study and their relatives of the orders Propionibacteriales, Pseudonocardiales, Streptomyces and Streptosporangiales in the class Actinobacteria. Bootstrap values (>70%) are shown at the branching points. Asterisks indicate that the corresponding branches were also recovered in both the maximum-likelihood and maximum-parsimony trees. Bar, 0.02 substitutions per nucleotide position.
Thermomonosporaceae and the genus Streptosporangium of the family Stereptosporangiaceae, respectively. These strains were also characterized by the formation of branched mycelium (Fig. 1). Members of the genera Actinomadura and Streptosporangium are further characterized by the formation of short chains of spores and globose sporangia, respectively (Trujillo and Goodfellow, 2012; Quintana and Goodfellow, 2012). Lastly, 1 strain were found to belong to the class Thermoleophilia and further assigned to the genus Conexibacter of the Conexibacteraceae in the order Solirubrobacterales (Fig. 5).

**Description of Dietzia schimae IMCC 12370**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and short rod-shaped. Colonies are circular, convex, smooth and white-colored after 3 days of incubation on MA at 15°C. Positive for nitrate reduction, glucose fermentation, urease and β-galactosidase in API 20NE but negative for indole production, arginine dihydrolase, esculin hydrolysis, gelatinase and β-galactosidase. Does not utilize D-glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate. Strain IMCC 12370 (NIBRBA0000114860) has been isolated from seawater, Sokcho, Korea.

**Description of Mycobacterium septicum WL1**

Cells are Gram-staining-negative, flagellated and rod-shaped. Colonies are circular, raised, entire and white-colored after 2 days on R2A at 25°C. Positive for nitrate reduction, glucose fermentation, urease and β-galactosidase in API 20NE but negative for indole production, arginine dihydrolase, esculin hydrolysis and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, malate, citrate and phenylacetate are utilized. Does not utilize caprate and adipate. Strain WL1 (NIBRBA0000114765) has
been isolated from freshwater, Jeonju, Korea.

**Description of Mycobacterium hodleri WS80**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and rod-shaped. Colonies are circular, convex, smooth and orange-colored after 3 days on R2A at 25°C. Negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis, gelatinase and β-galactosidase in API 20NE. Gluconate is utilized. Does not utilize D-glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, caprate, adipate, malate, citrate and phenylacetate. Strain WS80 (= NIBRBA0000115031) has been isolated from freshwater, Changnyeong, Korea.

**Description of Gordonia neofelifaecis MN13**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular and cream-colored after 3 days on MA at 15°C. Positive for β-galactosidase in API 20NE but negative for indole production, arginine dihydrolase, esculin hydrolysis and gelatinase. D-Glucose, L-arabinose, D-mannitol, N-acetyl-glucosamine, gluconate and adipate are utilized. Does not utilize D-mannose, D-maltose, caprate, malate, citrate and phenylacetate. Strain MN13 (= NIBRBA0000114947) has been isolated from gut of mugil cephalus, Korea.

**Description of Nocardia niigatensis NU 4Y-9-1**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are punctiform dry, flat and orange-colored after 3 days on TSA at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. L-Arabinose, N-acetyl-glucosamine, gluconate and malate are utilized. Does not utilize D-glucose, D-mannitol, D-mannose, D-maltose, caprate, adipate, citrate and phenylacetate. Strain NU 4Y-9-1 (= NIBRBA0000114874) has been isolated from ginseng soil, Anseong, Korea.

**Description of Rhodococcus phenolicus YB.Ce-3**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and coccus or coccoid-rod-shaped. Colonies are circular, raised, entire and pale-yellow-colored after 2 days on R2A at 30°C. Positive for nitrate reduction, arginine dihydrolase, urease, esculin hydrolysis and β-galactosidase in API 20NE but negative for indole production, glucose fermentation and gelatinase. N-Acetyl-glucosamine and adipate are utilized. Does not utilize D-glucose, L-arabinose, D-mannitol, D-mannose, D-maltose, gluconate, caprate malate, citrate and phenylacetate. Strain YB.Ce-3 (= NIBRBA0000114843) has been isolated from activated sludge, Daejeon, Korea.

**Description of Geodermatophilus terrae WM99**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and rod-shaped. Colonies are circular, convex, smooth and pink-colored after 3 days on R2A at 25°C. Positive for esculin hydrolysis and nitrate reduction (weak) but negative for indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannose, D-maltose and malate are utilized. Does not utilize D-mannitol, N-acetyl-glucosamine, gluconate, caprate, adipate, citrate and phenylacetate. Strain WM99 (= NIBRBA0000115029) has been isolated from freshwater, Changnyeong, Korea.

**Description of Actinocatenispora sera BS8**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and branched mycelium-forming. Colonies are lenticular, raised, entire and white-colored after 2 days on R2A at 30°C. Positive for nitrate reduction, arginine dihydrolase, esculin hydrolysis and β-galactosidase in API 20NE but negative for indole production, glucose fermentation and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, adipate, malate and citrate are utilized. Does not utilize caprate and phenylacetate. Strain BS8 (= NIBRBA0000114830) has been isolated from spent mushroom compost, Yesan, Korea.

**Description of Micromonospora siamensis 61DPR39**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are punctiform, digging and orange-colored on R2A at 25°C. Positive for esculin hydrolysis, gelatinase and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase and urease. D-Glucose, D-mannitol, D-mannose, D-maltose and malate are utilized. Does not utilize L-arabinose, N-acetyl-glucosamine, gluconate, caprate, adipate, citrate and phenylacetate. Strain 61DPR39 (= NIBRBA0000114798) has been isolated from freshwater, Daejeon, Korea.

**Description of Brevibacterium casei Ho-02**

Cells are Gram-staining-negative, non-flagellated, non-pigmented and rod-shaped. Colonies are circular, raised, undulate and white-yellow-colored after 2 days on R2A at 30°C. Positive for gelatinase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis and β-galactosidase. D-Glucose, D-mannitol,
N-acetyl-glucosamine, D-maltose, gluconate, adipate, malate, citrate and phenylacetate are utilized. Does not utilize L-arabinose, D-mannose and caprate. Strain Ho-02 (= NIBRBA0000114825) has been isolated from activated sludge, Daejeon, Korea.

**Description of Brevibacterium epidermidis RMD 3Y-15-4**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, entire, convex and pale-yellow-colored after 2 days on R2A at 30°C. Positive for glucose fermentation and gelatinase in API 20NE but negative for nitrate reduction, indole production, arginine dihydrolase, urease, esculin hydrolysis and β-galactosidase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, glucanate, caprate, adipate, malate, citrate, phenylacetate are utilized. Strain RMD 3Y-15-4 (= NIBRBA0000114878) has been isolated from ginseng soil, Anseong, Korea.

**Description of Cellulomonas cellasea HMF2762**

Cells are Gram-staining-positive, flagellated, non-pigmented and rod-shaped. Colonies are circular, convex, entire and yellow-colored after 5 days on R2A at 30°C. Positive for nitrate reduction and esculin hydrolysis in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease, gelatinase and β-galactosidase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate are not utilized. Strain HMF2762 (= NIBRBA 0000115002) has been isolated from sediment, Taebaek, Korea.

**Description of Cellulomonas biazotea NK 4Y-9-3**

Cells are Gram-staining-negative, non-flagellated and rod-shaped. Colonies are circular, entire, raised and yellow-colored after 2 days on R2A at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, N-acetyl-glucosamine, D-maltose and gluconate are utilized. Does not utilize D-mannose, caprate, adipate, malate, citrate and phenylacetate. Strain NK 4Y-9-3 (= NIBRBA 0000114888) has been isolated from ginseng soil, Anseong, Korea.

**Description of Cellulomonas persica WW28**

Cells are Gram-staining-positive, non-pigmented and rod-shaped. Colonies are circular, smooth, convex and light-yellow-colored after 3 days on R2A at 25°C. Positive for nitrate reduction and esculin hydrolysis and β-galactosidase in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose and D-maltose are utilized. Does not utilize N-acetyl-glucosamine, gluconate, caprate, adipate, malate, citrate and phenylacetate. Strain WW28 (= NIBRBA0000115028) has been isolated from freshwater, Changnyeong, Korea.

**Description of Brachybacterium saurashtreense EgT0207**

Cells are Gram-staining-positive, non-flagellated and cocccoid-shaped. Colonies are circular and cream-colored after 3 days on TSA at 25°C. Positive for nitrate reduction, esculin hydrolysis and β-galactosidase in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, D-mannitol and gluconate are utilized. Does not utilize L-arabinose, D-mannose, N-acetyl-glucosamine, D-maltose, caprate, adipate, malate, citrate and phenylacetate. Strain EgT0207 (= NIBRBA0000114942) has been isolated from gut of fulvia mutica, Korea.

**Description of Ornithinimicrobium murale KHS04**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular and cream-colored after 3 days on NA at 25°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, D-mannitol, D-mannose, D-maltose and citrate are utilized. Does not utilize L-arabinose, N-acetyl-glucosamine, gluconate, caprate, adipate, malate and phenylacetate. Strain KHS04 (= NIBRBA0000114950) has been isolated from gut of Todarodes pacificus, Korea.

**Description of Arthrobacter chlorophenolicus NK 6Y-6-4**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, entire, raised and white-colored after 3 days on NA at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, N-acetyl-glucosamine, D-maltose and gluconate are utilized. Does not utilize D-mannose, caprate, adipate, malate, citrate and phenylacetate. Strain NK 6Y-6-4 (= NIBRBA 0000114890) has been isolated from ginseng soil, Anseong, Korea.
**Description of Arthrobacter pascens NS 4Y-8-4**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, entire, convex and white-colored after 2 days on R2A at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, malate and citrate are utilized. Does not utilize caprate, adipate and phenylacetate. Strain NS 4Y-8-4 (= NIBRBA0000114873) has been isolated from ginseng soil, Anseong, Korea.

**Description of Kocuria rosea 145-10**

Cells are Gram-staining-negative, non-flagellated and rod-shaped. Colonies are circular, raised, entire and yellow-colored after 2 days on TSA at 25°C. Positive for nitrate reduction and β-galactosidase in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis and gelatinase. D-Glucose, D-maltose and malate are utilized. Does not utilize L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, gluconate, ca-prate, adipate, citrate and phenylacetate. Strain 145-10 (= NIBRBA0000114763) has been isolated from freshwater, Cheongsong, Korea.

**Description of Micrococcus antarcticus AX5**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, raised, entire and pale-yellow-colored after 2 days on R2A at 30°C. Negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis, gelatinase and β-galactosidase in API 20NE. D-Glucose, N-acetyl-glucosamine, D-maltose and malate are utilized. Does not utilize L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, gluconate, ca-prate, adipate, citrate and phenylacetate. Strain AX5 (= NIBRBA0000114894) has been isolated from mud flat, Incheon, Korea.

**Description of Sinomonas flava NGS 3Y-15-2**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, raised, entire and pale-yellow-colored after 2 days on R2A at 30°C. Positive for nitrate reduction, esculin hydrolysis and β-galactosidase in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, D-mannitol, D-mannose, D-maltose, gluconate, malate, citrate and phenylacetate are utilized. Does not utilize L-arabinose, N-acetyl-glucosamine, caprate and adipate. Strain NGS 3Y-15-2 (= NIBRBA0000114885) has been isolated from Ginseng soil, Anseong, Korea.

**Description of Diaminobutyricibacter tongyongensis R1-6**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and rod-shaped. Colonies are circular, convex, entire and yellow-colored on R2A at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate are not utilized. Strain R1-6 (= NIBRBA0000114812) has been isolated from soil, Daejeon, Korea.

**Description of Luteimicrobium subarcticum N1-9**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and rod-coccus-shaped. Colonies are circular, convex, entire and white-colored on NA at 30°C. Positive for esculin hydrolysis and β-galactosidase in API 20NE, weakly positive for nitrate reduction and glucose fermentation but negative for indole production, arginine dihydrolase, urease and gelatinase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate are not utilized. Strain N1-9 (= NIBRBA0000114811) has been isolated from soil, Daejeon, Korea.

**Description of Kribbella antibiotica RK 4Y-2-4**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are punctiform, dry, flat and white-colored after 2 days on R2A at 30°C. Positive for urease, esculin hydrolysis, gelatinase and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation and arginine dihydrolase. L-Arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine and D-maltose are utilized. Does not utilize D-glucose, gluconate, caprate, adipate, malate, citrate and phenylacetate. Strain RK 4Y-2-4 (= NIBRBA0000114877) has been isolated from ginseng soil, Anseong, Korea.

**Description of Nocardioides albus RS 4Y-2-4**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are circular, convex, raised and white-colored after 2 days on ISP4 at 30°C. Positive for esculin hydrolysis, gelatinase and β-galactosidase in API 20NE but negative for nitrate reduction, indole pro-
duction, glucose fermentation, arginine dihydrolase and urease. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, gluconate, adipate and malate, citrate are utilized. Does not utilize D-maltose, caprate and phenylacetate. Strain BS22 (= NIBRBA0000114835) has been isolated from spent mushroom compost, Yesan, Korea.

**Description of Kutzneria buriramensis RMD 3Y-3-1**

Cells are Gram-staining-positive, non-flagellated and rod-shaped. Colonies are undulate, dry, flat and white-colored after 2 days on ISP4 at 30°C. Positive for β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis and gelatinase. L-Arabinose and malate are utilized. Does not utilize D-glucose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, citrate and phenylacetate. Strain RMD 3Y-3-1 (= NIBRBA0000114879) has been isolated from ginseng soil, Anseong, Korea.

**Description of Kitasatospora paranensis NU 4Y-9-4**

Cells are Gram-staining-positive, non-flagellated and coccus-shaped. Colonies are circular, dry, flat and brown-colored after 3 days on ISP4 at 30°C. Positive for gelatinase and β-galactosidase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease and esculin hydrolysis. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate are utilized. Does not utilize D-glucose and caprate. Strain NU 4Y-9-4 (= NIBRBA0000114875) has been isolated from ginseng soil, Anseong, Korea.

**Description of Streptomyces drozdowiczii BBT1**

Cells are Gram-staining-positive, non-flagellated and non-pigmented and branched mycelium-forming. Colonies are lenticular, raised, entire and white-colored after 2 days on R2A at 30°C. Positive for nitrate reduction, arginine dihydrolase, urease, esculin hydrolysis, gelatinase, β-galactosidase in API 20NE but negative for indole production and glucose fermentation. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, adipate, malate, citrate and phenylacetate are utilized. Does not utilize caprate. Strain BBT1 (= NIBRBA0000114837) has been isolated from powder of biotite, Yesan, Korea.

**Description of Streptomyces thermocophilus BS22**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and branched mycelium-forming. Colonies are lenticular, raised, entire and white-colored after 2 days on R2A at 30°C. Positive for nitrate reduction, arginine dihydrolase, urease, esculin hydrolysis, gelatinase, β-galactosidase in API 20NE but negative for indole production and glucose fermentation. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, adipate, malate, citrate and phenylacetate are utilized. Does not utilize caprate. Strain BS22 (= NIBRBA0000114835) has been isolated from spent mushroom compost, Yesan, Korea.

**Description of Streptomyces olivochromogenes I1-6**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and branched mycelium-forming. Colonies are rhizoid, penetrate and light brownish-colored on ISP2 at 30°C. Positive for esculin hydrolysis in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, gelatinase and β-galactosidase. L-Arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, adipate, malate, citrate and phenylacetate are utilized. Does not utilize D-glucose and caprate. Strain I1-6 (= NIBRBA0000114809) has been isolated from soil, Dajeon, Korea.

**Description of Streptomyces tsukubensis T1-6**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and rod-shaped. Colonies are rhizoid, penetrate and white to creamy-colored on TSA at 30°C. Positive for nitrate reduction arginine dihydrolase and urease in API 20NE but negative for indole production, glucose fermentation, esculin hydrolysis, gelatinase and β-galactosidase. D-Glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate, adipate, malate, citrate and phenylacetate are not utilized. Strain T1-6 (= NIBRBA0000114816) has been isolated from soil, Dajeon, Korea.

**Description of Actinomadura bangladeshensis 7C-18**

Cells are Gram-staining-positive, non-flagellated, non-pigmented and branched mycelium-forming. Colonies are lenticular, raised, entire and white-colored after 2 days on R2A at 30°C. Positive for arginine dihydrolase and urease in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, esculin hydrolysis, gelatinase and β-galactosidase. D-Glucose, D-mannose, D-maltose and gluconate are utilized. Does not utilize L-arabinose, D-mannitol, N-acetyl-glucosamine, caprate, adipate, malate, citrate and phenylacetate. Strain 7C-18 (= NIBRBA0000114839) has been isolated from activated sludge, Dajeon, Korea.
Description of *Streptosporangium amethystogenes* WM35

Cells are Gram-staining-positive, non-flagellated, non-pigmented and branched mycelium-forming. Colonies are round, circular, convex and pale red-colored after 7 days on R2A at 25°C. Positive for nitrate reduction and gelatinase in API 20NE but negative for indole production, glucose fermentation, arginine dihydrolase, urease, esculin hydrolysis and β-galactosidase. D-Maltose and citrate are utilized. Does not utilize D-glucose, L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, gluconate, caprate, adipate, malate and phenylacetate. Strain WM35 (＝NIBRBA0000115020) has been isolated from freshwater, Changnyeong, Korea.

Description of *Conexibacter arvalis* BS10

Cells are Gram-staining-positive, non-pigmented and rod-shaped. Colonies are circular, raised, entire and white-colored after 4 days on R2A at 30°C. Positive for urease and gelatinase in API 20NE but negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, esculin hydrolysis and β-galactosidase. D-Glucose is utilized. Does not utilize L-arabinose, D-mannitol, D-mannose, N-acetyl-glucosamine, D-maltose, gluconate, caprate adipate, malate, citrate and phenylacetate. Strain BS10 (＝NIBRBA0000114831) has been isolated from spent mushroom compost, Yesan, Korea.

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REFERENCES


