A report of 29 unrecorded bacterial species belonging to the phylum *Bacteroidetes* in Korea

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Within a comprehensive, widescale investigation of indigenous prokaryotic species in Korea, 29 bacterial strains in the phylum *Bacteroidetes* were isolated from diverse environmental habitats that included soil, plant roots, natural caves, tidal flats, freshwater from lakes, and seawater. Based on their high 16S rRNA gene sequence similarities (>99.1%) and the formation of robust phylogenetic clades with the closest type species, each strain likely belonged to an independent and predefined bacterial species. There are no publications or official reports of the isolation of these 29 species in Korea. Our study provides strong evidence that seven species in three genera in the order *Cytophagales*, 15 species in 13 genera in the order *Flavobacteriales* and seven species in five genera in the order *Sphingobacteriales*, all within the phylum *Bacteriodetes*, are new reports of bacterial species in Korea. Gram reaction, colony and cell morphology, basic biochemical characteristics, isolation source, and strain IDs are described in the species description section.

Keywords: 16S rRNA, bacterial diversity, *Bacteroidetes*, indigenous Korean prokaryotic species, taxonomy, unrecorded species

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Introduction

The phylum *Bacteroidetes*, also known as the *Cytophaga-Flexibacter-Bacteroides* (CFB) group, is a diverse bacterial phylum whose taxonomic designation has changed several times in the past several years. (Woose, 1987; Woose *et al.*, 1990). Species or strains in the *Bacteroidetes* have been isolated from diverse environmental habitats including terrestrial ecosystems and freshwater and marine systems (Bernardet *et al.*, 1996; Bernardet *et al.*, 2002; Kirchman, 2002). Cultured isolates of *Bacteroidetes* are all gram-negative heterotrophs, notable for their ability to efficiently degrade complex biomacromolecules such as protein, chitin, pectin, agar, starch, and cellulose

(Alain et al., 2008).

In 2015, we collected environmental samples from diverse habitats in Korea and isolated many novel and unrecorded bacterial species during a research program supported by NIBR of Korea. This report focused on the description of bacterial species in the phylum *Bacteroidetes* that were not previously reported in Korea. Here, we report 29 unrecorded bacterial species belonging to the *Bacteroidetes*, which consisted of 29 genera within three orders.

MATERIALS AND METHODS

A total of 29 bacterial strains assigned to the phylum

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Bacteroidetes were isolated from various environmental habitats, including soil, plant roots, natural caves, tidal flats, freshwater, and seawater (Table 1). All each environmental samples were processed separately, serially diluted, spread onto diverse culture agar media including R2A agar (BD), marine agar 2216 (MA, BD), Kuster's agar (KUA: glycerol, 10 g; casein, 0.3 g: KNO₃, 2 g; K₂HPO₄, 2 g; soluble starch, 0.5 g; asparagine, 0.1 g; FeSO₄·7H₂O, 0.01 g; CaCO₃, 0.02 g; MgSO₄·7H₂O, 0.05 g; agar, 15 g per 1,000 mL filtered sea water, pH 7.0) and nutrient agar (NA, BD), and incubated at 25-30°C for 2-5 days (Table 1). The designated strain IDs, isolation sources, culture media, and incubation conditions are summarized in Table 1. All strains were isolated as pure cultures and stored as 10-20% glycerol suspension at -80°C and as lyophilized ampoules.

Colony morphology of the strains was observed by eye or a magnifying glass after the cells were cultivated to their stationary phase on their culture agar media. Cellular morphology and cell size were examined using either transmission electron or scanning electron microscopy. Gram staining was performed using a gram-staining kit according to the standard procedures. Biochemical characteristics were tested by using API 20NE galleries (bioMérieux) according to the manufacturer's instructions

Bacterial DNA extraction, PCR amplification, and 16S rRNA gene sequencing were performed using standard procedures as described elsewhere (Lee *et al.*, 2014). The 16S rRNA gene sequences of the strains assigned to the *Bacteriodetes* were compared with those of the type strains of validated bacterial species using the EzTaxone server (Kim *et al.*, 2012). For phylogenetic analyses, multiple alignments of the 16S rRNA gene sequences of the isolates and those of the valid type strains were carried out using the Clustal_X program (Thompson *et al.*, 1997). Evolutionary distances were calculated using the Kimura two-parameter model (Kimura, 1983) and the phylogenetic trees were constructed using a neighborjoining algorithm (Saitou and Nei, 1987) with bootstrap values based on 1,000 replications (Felsenstein, 2002).

RESULTS AND DISCUSSION

The 29 strains were distributed into 3 orders in the phylum *Bacteroidetes*: 15 strains in the order *Flavobacteriales*, seven strains in the *Cytophagales*, and seven strains in the *Sphingobacteriales* (Table 1). These strains were gram-staining-negative, chemoheterotrophic, and rodshaped bacteria (Fig. 1). Colony size, morphology, and physiological characteristics are presented in the species description section.

Seven strains were assigned to the family Cyclobacte-

riaceae, Dyadobacter and Cytophagaceae in the order Cytophagales (Fig. 2). Among these strains, three that were assigned to the family Cyclobacteriaceae belonged to the genus Algoriphagus, one strain that was assigned to the family *Dyadobacter* belonged to the genus Dyadobacter, and three strains that were assigned to the family Cytophagaceae belonged to the genus Arcicella. Strains that were assigned to the order Flavobacteriales (Fig. 3) belonged to one family and 13 genera: Flavobacterium (3 species), Arenibacter (1 species), Muricauda (1 species), Zobellia (1 species), Maribacter (1 species), Formosa (1 species), Mariniflexile (1 species), Cellulophaga (1 species), Mesonia (1 species), Arenitalea (1 species), Sediminicola (1 species), Zeaxanthinibacter (1 species), and Cytophaga (1 species). Seven strains were assigned to the family Sphingobacteriaceae and Chitinophagaceae in the order Sphingobacteriales (Fig. 4). Among these, three strains that were assigned to the family Sphingobacteriaceae belonged to the genus Sphingobacterium (1 species) and *Pedobacter* (2 species) and four strains that were assigned to the family Chitinophagaceae belonged to the genus Chitinophaga (2 species), Niabella (1 species) and Asinibacterium (1 species). Here, we report 29 unrecorded bacterial species belonging to 6 families in 3 orders in the Bacteriodetes, which were isolated in Korea.

Description of Algoriphagus namhaensis HN30-2

Cells are gram-staining-negative, non-flagellated, non-pigmented, and rod-shaped. Colonies are circular, convex, smooth, and reddish orange colored after 3 days of incubation at 25°C on MA. Positive for β -galactosidase activity and esculin hydrolysis. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase, gelatinase, and urease activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, D-maltose, N-acetyl-glucosamine, capric acid, adipic acid, phenylacetic acid, malic acid, trisodium citrate, and potassium gluconate. Strain HN30-2 (=NIBRBAC000497930) was isolated from a seawater sample, Haenam, Korea.

Description of Algoriphagus olei RDH3

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are smooth, shiny, convex, and pinkish red-colored after 2 days of incubation at 30°C on R2A. Positive for esculin hydrolysis and gelatinase and β -galactosidase activities. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase and urease activities. D-Glucose, D-mannose, N-acetyl-glucosamine, and D-maltose are utilized. Does not utilize L-arabinose, D-mannitol, adipic acid, malic acid, phenylacetic acid, trisodium citrate,

 Table 1. Summary of strains isolated belonging to the Bacteriodetes and their taxonomic affiliations.

Order	Family	Genus	Strain ID	NIBR ID	Most closely related species	Similarity (%)	Isolation source	Medium	Incubation conditions
	Cyclobacteriaceae	Algoriphagus Algoriphagus Algoriphagus	G4 RDH3 HN30-2	NIBRBAC000498067 NIBRBAC000498077 NIBRBAC000497930	Algoriphagus winogradskyi Algoriphagus olei Algoriphagus namhaensis	99.5 99.6 99.4	Tidal flat Fresh water Seawater	R2A R2A MA	30°C, 2d 30°C, 2d 25°C, 3d
Cy tophagales	Dyadobacter_f	Dyadobacter	M2015-1	NIBRBAC000498053	Dyadobacter psychrophilus	100	Soil	R2A	25°C, 2d
	Cytophagaceae	Arcicella Arcicella Arcicella	HMF3809 JD36 YBF1	NIBRBAC000497898 NIBRBAC000497944 NIBRBAC000498047	Arcicella aquatica Arcicella rosea Arcicella rigui	99.7 99.8 99.4	Fresh water Fresh water Freshwater	R2A R2A R2A	25°C, 2d 25°C, 3d 25°C, 2d
		Flavobacterium Flavobacterium Flavobacterium Arenibacter	RDH2 HMF4059 2015-2 MW24	NIBRBAC000498076 NIBRBAC000497910 NIBRBAC000498054 NIBRBAC000497863	Flavobacterium cheniae Flavobacterium chilense Flavobacterium columnare Arenibacter echinorum	99.9 99.6 99.0 99.4	Lake Fresh water Fresh water Tidal flat	R2A R2A R2A MA	30°C, 2d 30°C, 3d 25°C, 2d 30°C, 2d
Flavobacteriales	Flavobacteriaceae	Muricauda Zobellia Maribacter Formosa Mariniflexile	MW105 HMF3932 HN1 HN41 HN49	NIBRBACC00497905 NIBRBACC00497905 NIBRBACC00497929 NIBRBACC000497931	Muricauda antarctica Zobellia galactanivorans Maribacter caenipelagi Formosa spongicola Mariniflexile aquimaris	99.7 99.9 99.9 99.7	Tidal flat Tidal flat Tidal flat Tidal flat Tidal flat	MA MA MA	30°C, 2d 30°C, 3d 25°C, 3d 25°C, 3d 25°C, 3d
		Cellulophaga Mexonia Arenitalea Sediminicola Zeaxanthinibacter Cytophaga	KYW1172 KYW1047 LPB0108 LPB0111 LPB0115 DF-2	NIBRBAC000497941 NIBRBAC000497945 NIBRBAC000497984 NIBRBAC000497986 NIBRBAC000497987	Cellulophaga baltica Mesonia algae Arenitalea lutea Sediminicola luteus Zeaxanthinibacter enoshimensis Cytophaga massiliensis	99.6 99.6 100 99.4 100	Seawater Seawater Seawater Seawater Fresh water	MA MA MA MA R2A	25°C, 4d 25°C, 3d 26°C, 2d 26°C, 2d 26°C, 2d 25°C, 2d
	Sphingobacteriaceae	Sphingobacterium Pedobacter Pedobacter	HMF3876 HMF3898 NU3	NIBRBAC000497901 NIBRBAC000497903 NIBRBAC000498049	Sphingobacterium multivorum Pedobacter piscium Pedobacter glucosidilyticus	99.4 99.8 98.8	Soil Soil Fresh water	R2A R2A R2A	30°C, 3d 30°C, 3d 25°C, 2d
Sphingobacteriales	Chitinophagaceae	Chitinophaga Chitinophaga Niabella Asinibacterium	NR 2-02 C2-40 HMF2508 M2015-5	NIBRBAC000498082 NIBRBAC000498025 NIBRBAC000497896 NIBRBAC000498048	Chitinophaga oryziterrae Chitinophaga arvensicola Niabella thaonhiensis Asinibacterium lactis	99.1 99.6 99.9 100	Plant root Natural cave Soil Soil	R2A TSA R2A R2A	30°C, 2d 30°C, 5d 30°C, 3d 25°C, 2d

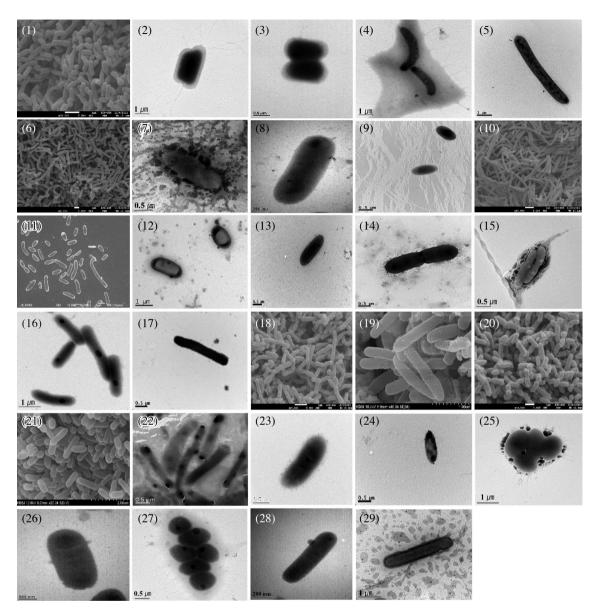


Fig. 1. Transmission electron micrographs or scanning electron micrographs of cells of the strains isolated in this study. The cells were cultured at their optimal growth conditions. Strains: 1, HN30-2; 2, RDH3; 3, G4; 4, HMF3809; 5, YBF1; 6, JD36; 7, MW24; 8, LPB0108; 9, M2015-5; 10, KYW1172; 11, C2-40; 12, NR 2-02; 13, DF-2; 14, M2015-1; 15, RDH2; 16, HMF4059; 17, 2015-2; 18, HN41; 19, HN1; 20, HN49; 21, KYW1047; 22, MW105; 23, HMF2508; 24, NU3; 25, HMF3898; 26, LPB0111; 27, HMF3876; 28, LPB0115; 29, HMF3932.

capric acid, and potassium gluconate. Strain RDH3 (= NIBRBAC000498077) was isolated from a lake water sample, Chung-Ang University, Korea.

Description of Algoriphagus winogradskyi G4

Cells are Gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, shiny, entire, and bright-pink-colored after 2 days of incubation at 30°C on R2A. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase,

gelatinase, and urease activities. D-Glucose, L-arabinose, D-mannitol, D-maltose, adipic acid, malic acid, D-mannose, potassium gluconate, *N*-acetyl-glucosamine, and trisodium citrate are utilized. Does not utilize capric acid and phenylacetic acid. Strain G4 (= NIBRBAC0004980 67) was isolated from a tidal flat sample, Ganghwa-gun, Korea.

Description of Arcicella aquatic HMF3809

Cells are gram-staining-negative, non-flagellated, non-pigmented, and curved rod-shaped. Colonies are circu-

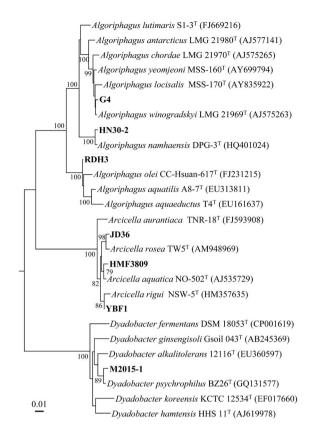


Fig. 2. Neighbor-joining phylogenetic tree based on 16S rRNA gene sequences, showing the phylogenetic relationships between the strains isolated in this study and their relatives of the order *Cytophagales* in the phylum *Bacteroidetes*. *Escherichia coli* ATCC 11775^T (X80725) was used as an outgroup (not shown). Bootstrap values (>70%) are shown above nodes. Scale bar: 0.01 changes per nucleotide.

lar, convex, entire, and pink-colored after 3 days of incubation at 30°C on R2A agar. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. Does not utilize D-glucose, D-mannose, D-mannitol, L-arabinose, malic acid, trisodium citrate, N-acetyl-glucosamine, capric acid, potassium gluconate, adipic acid, D-maltose, and phenylacetic acid. Strain HMF3809 (= NIBRBAC000497898) was isolated from a freshwater sample, Yongin, Korea.

Description of Arcicella rigui YBF1

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, mucous, and pink-colored after 2 days of incubation at 25°C on R2A agar. Positive for β -galactosidase activity. Negative for nitrate reduction, indole production, glucose fermentation, esculin hydrolysis, and arginine dihydrolase, urease, and gelatinase activities. Does not utilize D-glucose, L-arab-

inose, D-mannose, D-mannitol, D-maltose, *N*-acetyl-glucosamine, capric acid, adipic acid, trisodium citrate, phenylacetic acid, potassium gluconate, and malic acid. Strain YBF1 (= NIBRBAC000498047) was isolated a fresh water sample, Yeonggwang-gun, Korea.

Description of Arcicella rosea JD36

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are smooth, circular, translucent, and pale pink-colored after 3 days of incubation at 25°C on R2A agar. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase and urease activities. Positive for esculin hydrolysis and gelatinase and β -galactosidase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-maltose, D-mannitol, *N*-acetyl-glucosamine, adipic acid, malic acid, potassium gluconate, phenylacetic acid, trisodium citrate, and capric acid. Strain JD36 (= NIBRBAC0004 97944) was isolated from a fresh water sample, Changwon, Korea.

Description of Arenibacter echinorum MW24

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, raised, and entire pale yellow-colored after 2 days of incubation at 30°C on MA. Positive for nitrate reduction, esculin hydrolysis, and β -galactosidase activity. Negative for indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. D-Glucose, D-mannose, N-acetyl-glucosamine, and D-maltose are utilized. Does not utilize L-arabinose, D-mannitol, potassium gluconate, capric acid, adipic acid, malic acid, trisodium citrate, and phenylacetic acid. Strain MW24 (= NIBRBAC0004 97863) was isolated from a tidal flat sample, Taean, Korea.

Description of Arenitalea lutea LPB0108

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular and yellow-colored after 2 days of incubation on MA at 26°C. Positive for esculin hydrolysis. Negative for nitrate production, glucose fermentation, indole production, and urease, arginine dihydrolase, β -galactosidase, and gelatinase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, N-acetyl-glucosamine, D-maltose, malic acid, potassium gluconate, capric acid, adipic acid, trisodium citrate, and phenylacetic acid. Strain LPB0108 (= NIBRBAC000497984) was isolated from a seawater sample, Busan, Korea.

Description of Asinibacterium lactis M2015-5

Cells are gram-staining-negative, non-flagellated, and

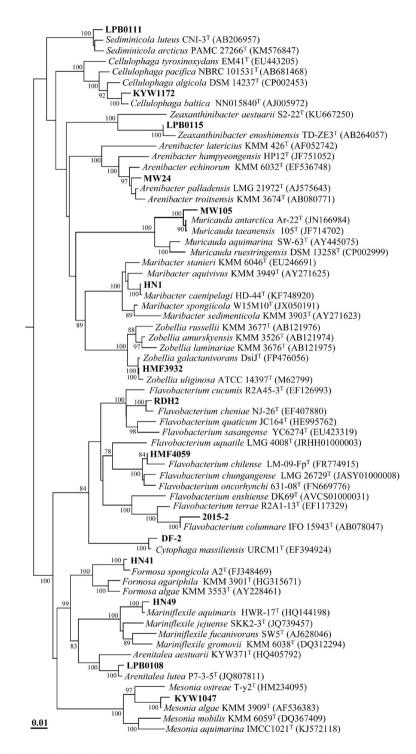


Fig. 3. Neighbor-joining phylogenetic tree based on 16S rRNA gene sequences, showing the phylogenetic relationships between the strains isolated in this study and their relatives of the order *Flavobacteriales* in the phylum *Bacteroidetes*. *Escherichia coli* ATCC 11775^T (X80725) was used as an outgroup (not shown). Bootstrap values (>70%) are shown above nodes. Scale bar: 0.01 changes per nucleotide.

rod-shaped. Colonies are circular, convex, and yellow-colored after 2 days of incubation at 25°C on R2A. Positive for esculin hydrolysis. Negative for nitrate reduction, indole production, glucose fermentation, and ure-

ase, arginine dihydrolase, gelatinase, and β -galactosidase activities. L-Arabinose is utilized. Dose not assimilate D-glucose, D-mannose, D-mannitol, N-acetyl-glucosamine, D-maltose, potassium gluconate, trisodium ci-

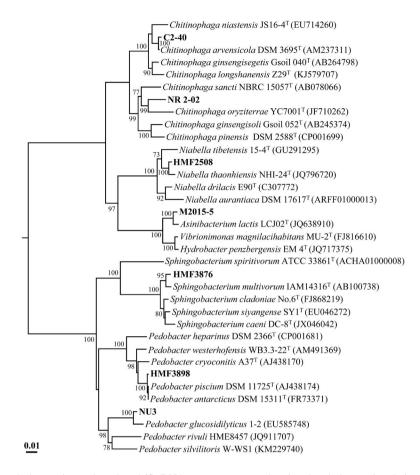


Fig. 4. Neighbor-joining phylogenetic tree based on 16S rRNA gene sequences, showing the phylogenetic relationships between the strains isolated in this study and their relatives of the order *Sphingobacteriales* in the phylum *Bacteroidetes*. *Escherichia coli* ATCC 11775 T (X80725) was used as an outgroup (not shown). Bootstrap values (>70%) are shown above nodes. Scale bar: 0.01 changes per nucleotide.

trate, capric acid, phenylacetic acid, and adipic acid. Strain M2015-5 (= NIBRBAC000498048) was isolated from a soil sample, Wanju, Korea.

Description of Cellulophaga baltica KYW1172

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are irregular, opaque, and yellow-colored after 4 days of incubation at 25°C on MA. Positive for nitrate reduction, esculin hydrolysis, and β -galactosidase activity. Negative for indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, potassium gluconate, capric acid, trisodium citrate, phenylacetic acid, adipic acid, *N*-acetyl-glucosamine, D-maltose, and malic acid. Strain KYW1172 (= NIBRBAC000497941) was isolated from a seawater sample, Gwangyang, Korea.

Description of Chitinophaga arvensicola C2-40

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, entire, and

bright yellow-colored after 5 days of incubation on TSA at 30°C. Positive for esculin hydrolysis and β -galactosidase activities. Negative for nitrate reduction, indole production, glucose fermentation, arginine dihydrolase, urease, and gelatinase activities. D-Glucose, L-arabinose, D-mannose, N-acetyl-glucosamine and D-maltose is utilized. Does not utilize, malic acid, adipic acid, D-mannitol, potassium gluconate, phenylacetic acid, trisodium citrate, and capric acid. Strain C2-40 (= NIBRBAC 000498025) was isolated from a natural cave sample, Jeju island, Korea.

Description of Chitinophaga oryziterrae NR 2-02

Cells are gram-staining-negative, non-flagellated, and short rod-shaped. Colonies are circular, raised, entire and yellow-colored after 3 days of incubation on R2A agar at 30°C. Positive for nitrate reduction, esculin hydrolysis, and β -galactosidase activity. Negative for indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. D-Glucose, L-arabinose, D-mannose, N-acetyl-glucosamine, and

D-maltose are utilized. Does not utilize D-mannitol, potassium gluconate, malic acid, phenylacetic acid, trisodium citrate, capric acid, and adipic acid. Strain NR 2-02 (=NIBRBAC000498082) was isolated from a plant root sample, Jeongeuup, Korea.

Description of Cytophaga massiliensis DF-2

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies irregular, convex, and yellow-colored after 2 days of incubation at 25°C on R2A. Positive for esculin hydrolysis. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase, β -galactosidase, urease, and gelatinase activities. D-Glucose, L-arabinose, D-mannose, and D-maltose are utilized. Does not utilize D-mannitol, N-acetyl-glucosamine, potassium gluconate, adipic acid, malic acid, phenylacetic acid, trisodium citrate, and capric acid. Strain DF-2 (= NIBRBAC000498044) was isolated from a fresh water sample, Jeonju, Korea.

Description of Dyadobacter psychrophilus M2015-1

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, and yellow-colored after 2 days of incubation on R2A agar at 25°C. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, glucose fermentation, indole production, and urease, arginine dihydrolase, and gelatinase activities. D-Glucose, D-mannose, and adipic acid are utilized. Does not utilize L-arabinose, D-mannitol, *N*-acetyl-glucosamine, D-maltose, malic acid, potassium gluconate, phenylacetic acid, trisodium citrate, and capric acid. Strain M2015-1 (= NI BRBAC000498053) was isolated from a soil sample, Wanju, Korea.

Description of Flavobacterium cheniae RDH2

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, smooth, entire, and yellow-colored after 2 days of incubation at 30°C on R2A. Positive for gelatinase activity. D-Glucose, L-arabinose, D-maltose, D-mannose, D-mannitol, trisodium citrate, potassium gluconate, adipic acid, malic acid, and N-acetyl-glucosamine are utilized. Negative for nitrate reduction, indole production, glucose fermentation, esculin hydrolysis, and arginine dihydrolase, urease, and β -galactosidase activities. Does not utilize phenylacetic acid and capric acid. Strain RDH2 (= NIBRBAC00049 8076) was isolated from a lake water sample, Chung-Ang University, Korea.

Description of Flavobacterium chilense HMF4059

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, entire, and

yellow-colored after 3 days of incubation at 30°C on R2A agar. Positive for esculin hydrolysis and gelatinase and β -galactosidase activities. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase and urease activities. Does not utilize D-mannitol, potassium gluconate, capric acid, trisodium citrate, phenylacetic acid, adipic acid, and malic acid. D-Glucose, L-arabinose, D-mannose, *N*-acetyl-glucosamine, and D-maltose are utilized. Strain HMF4059 (=NIBRBAC000497910) was isolated from a fresh water sample, Jeju island, Korea.

Description of Flavobacterium columnare 2015-2

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, and yellow-colored after 2 days of incubation at 25°C on R2A agar. Positive for nitrate reduction and gelatinase activity. Negative for indole production, glucose fermentation, esculin hydrolysis, and arginine dihydrolase, β -galactosidase, and urease activities. Does not utilize D-glucose, D-mannose, D-mannitol, N-acetyl-glucosamine, D-maltose, potassium gluconate, malic acid, trisodium citrate, phenylacetic acid, L-arabinose, capric acid, and adipic acid. Strain 2015-2 (= NIBRBAC000498054) was isolated from a fresh water sample, Jeonju, Korea.

Description of Formosa spongicola HN41

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, and yellow-colored after 3 days on MA at 25°C. Positive for gelatinase activity. Negative for nitrate reduction, indole production, glucose fermentation, esculin hydrolysis, and arginine dihydrolase, urease, and β -galactosidase activities. Does not utilize D-glucose, L-arabinose, D-maltose, D-mannitol, N-acetyl-glucosamine, potassium gluconate, malic acid, trisodium citrate, phenylacetic acid, capric acid, and adipic acid. Strain HN41 (=NIBRBAC000497931) was isolated from a tidal flat sample, Haenam, Korea.

Description of Maribacter caenipelagi HN1

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are irregular, smooth, and light yellow-colored after 3 days of incubation at 25°C on MA. Positive for β -galactosidase activity and esculin hydrolysis. Negative for nitrate reduction, glucose fermentation, indole production, and arginine dihydrolase, urease, and gelatinase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, N-acetylglucosamine, D-maltose, potassium gluconate, adipic acid, malic acid, trisodium citrate, phenylacetic acid, and capric acid. Strain HN1 (= NIBRBAC000497929) was isolated from a tidal flat sample, Haenam, Korea.

Description of Mariniflexile aquimaris HN49

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, smooth, and yellow-colored after 3 days of incubation at 25°C on MA. Positive for nitrate reduction, esculin hydrolysis, and β -galactosidase activity. Negative for indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. D-Glucose, D-mannose, D-mannitol, and D-maltose are utilized. Does not utilize L-arabinose, N-acetyl-glucosamine, potassium gluconate, malic acid, capric acid, trisodium citrate, adipic acid, and phenylacetic acid. Strain HN49 (=NIBRBAC 000497932) was isolated from a seawater sample, Haenam, Korea.

Description of Mesonia algae KYW1047

Cells are Gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, opaque, entire, and yellow-colored after 3 days on MA at 25°C. Positive for β -galactosidase activity. Negative for nitrate reduction, esculin hydrolysis, indole production, glucose fermentation, and gelatinase, arginine dihydrolase, and urease activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, D-maltose, N-acetyl-glucosamine, potassium gluconate, trisodium citrate, malic acid, phenylacetic acid, capric acid, and adipic acid. Strain KYW1047 (= NIBRBAC000497945) was isolated from a sea water sample, Gwangyang, Korea.

Description of Muricauda antarctica MW105

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, raised, entire and pale yellow colored after 2 days on MA at 30°C. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, glucose fermentation, indole production, and arginine dihydrolase, urease, and gelatinase activities. D-Mannose and N-acetyl-glucosamine are utilized. Does not utilize D-glucose, L-arabinose, D-mannitol, D-maltose, potassium gluconate, adipic acid, malic acid, trisodium citrate, phenylacetic acid, and capric acid. Strain MW105 (= NIBRBAC000497883) was isolated from a tidal flat sample, Taean, Korea.

Description of Niabella thaonhiensis HMF2508

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, entire, and yellow-colored after 3 days of incubation at 30°C on R2A agar. Positive esculin hydrolysis and arginine dihydrolase activity. Negative for nitrate reduction, indole production, glucose fermentation, and urease, gelatinase, and β -galactosidase activities. Does not utilize potassium gluconate, malic acid, trisodium citrate, D-mannitol,

capric acid, adipic acid, and phenylacetic acid, but utilizes D-glucose, L-arabinose, D-mannose, *N*-acetyl-glucosamine, and D-maltose. Strain HMF2508 (= NIBR BAC000497896) was isolated from a soil sample, Yongin, Korea.

Description of Pedobacter glucosidilyticus NU3

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, and pink-colored after 2 days of incubation at 25°C on R2A agar. Positive for esculin hydrolysis. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase, β -galactosidase, urease and gelatinase activities. D-Glucose, L-arabinose, D-mannitol, N-acetyl-glucosamine, and D-maltose are utilized. Does not utilize D-mannose, capric acid, adipic acid, trisodium citrate, phenylacetic acid, potassium gluconate, and malic acid. Strain NU3 (=NIBRBAC000498049) was isolated from a fresh water sample, Jeonju, Korea.

Description of Pedobacter piscium HMF3898

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular, convex, entire, and cream-colored after 3 days of incubation at 30°C on R2A. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase, urease, and gelatinase activities. D-Glucose, L-arabinose, D-mannose, D-mannitol, and D-maltose are utilized. Does not utilize *N*-acetyl-glucosamine, potassium gluconate, adipic acid, trisodium citrate, malic acid, phenylacetic acid, and capric acid. Strain HMF3898 (= NI BRBAC000497903) was isolated from a soil sample, Yongin, Korea.

Description of Sediminicola luteus LPB0111

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular and dark yellow after 2 days of incubation at 26°C on MA. Positive for esculin hydrolysis and β -galactosidase activity. Negative for nitrate reduction, glucose fermentation, indole production, and arginine dihydrolase, urease, and gelatinase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, malic acid, trisodium citrate, potassium gluconate, adipic acid, D-maltose, N-acetyl-glucosamine, capric acid, and phenylacetic acid. Strain LPB0111 (= NIBRBAC000497986) was isolated from a sea water sample, Busan, Korea.

Description of *Sphingobacterium multivorum* HMF3876

Cells are gram-staining- negative, non-flagellated, and

rod-shaped. Colonies are circular, convex, entire, and cream-colored after 3 days of incubation at 30°C on R2A agar. Positive for esculin hydrolysis and arginine dihydrolase, urease, and β -galactosidase activities. Negative for nitrate reduction, indole production, glucose fermentation, and gelatinase activity. D-Glucose, L-arabinose, D-mannose, D-maltose, and N-acetyl-glucosamine are utilized. Does not utilize D-mannitol, potassium gluconate, adipic acid, trisodium citrate, malic acid, phenylacetic acid, and capric acid. Strain HMF3876 (= NIBR BAC000497901) was isolated from a soil sample, Yongin, Korea.

Description of Zeaxanthinibacter enoshimensis LPB0115

Cells are gram-staining-negative, non-flagellated, and rod-shaped. Colonies are circular and yellow-colored after 2 days of incubation on MA at 26°C. Positive for esculin hydrolysis. Negative for glucose fermentation, nitrate reduction, indole production, and arginine dihydrolase, urease, gelatinase, and β -galactosidase activities. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, malic acid, capric acid, trisodium citrate, phenylacetic acid, N-acetyl-glucosamine, D-maltose, potassium gluconate, and adipic acid. Strain LPB0115 (= NIBRBAC000497987) was isolated from a sea water sample, Busan, Korea.

Description of Zobellia galactanivorans HMF3932

Cells are gram-staining-negative, flagellated, and rod-shaped. Colonies are circular and yellow-colored after 3 days of incubation at 30°C on MA. Positive for esculin hydrolysis and urease, gelatinase, and β -galactosidase activities. Negative for nitrate reduction, indole production, glucose fermentation, and arginine dihydrolase activity. Does not utilize D-glucose, L-arabinose, D-mannose, D-mannitol, N-acetyl-glucosamine, D-maltose, potassium gluconate, adipic acid, trisodium citrate, capric acid, malic acid, and phenylacetic acid. Strain HMF3932 (=NIBRBAC000497905) was isolated from a seawater sample, Pohang, Korea.

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REFERENCES

- Alain, K., L. Intertaglia, P. Catala and P. Lebaron. 2008. *Eudoraea adriatica* gen. nov., sp. nov., a novel marine bacterium of the family *Flavobacteriaceae*. International Journal of Systematic and Evolutionary Microbiology 58:2275-2281.
- Bernardet, J.F., P. Segers, M. Vancanneyt, F. Berthe, K. Kersters and P. Vandamme. 1996. Cutting a Gordian knot: emended classification and description of the genus *Flavobacterium*, emended description of the family *Flavobacteriaceae*, and proposal of *Flavobacterium hydatis* nom. nov. (basonym, *Cytophaga aquatilis* Strohl and Tait 1978). International Journal of Systematic and Evolutionary Microbiology 46:128-148.
- Bernardet, J.F., Y. Nakagawa and B. Holmes. 2002. Proposed minimal standards for describing new taxa of the family *Flavobacteriaceae* and emended description of the family. International Journal of Systematic and Evolutionary Microbiology 52:1049-1170.
- Felsenstein, J. 2002. PHYLIP (phylogeny inference package), version 3.6a, Seattle: Department of genetics, University of Washington, Seattle, WA, USA.
- Kim, O.S., Y.J. Cho, K. Lee, S.H. Yoon, M. Kim, H. Na, S.C. Park, Y.S. Jeon, J.H. Lee, H. Yi, S. Won and J. Chun. 2012. Introducing EzTaxon-e: a prokaryotic 16S rRNA gene sequence database with phylotypes that represent uncultured species. International Journal of Systematic and Evolutionary Microbiology 62:716-721.
- Kimura, M. 1983. The Neutral Theory of Molecular Evolution. Cambridge: Cambridge University Press, Cambridge, New York.
- Kirchman, D.L. 2002. The ecology of Cytophaga-Flavobacteria in aquatic environments. FEMS Microbiol Ecology 39:91-100.
- Saitou, N. and M. Nei. 1987. The neighbor-joining method: a new method for reconstructing phylogenetic trees. Molecular Biology and Evolution 4:406-425.
- Thompson, J.D., T.J. Gibson, F. Plewniak, F. Jeanmougin and D.G. Higgins. 1997. The Clustal_X windows interface: flexible strategies for multiple sequence alignment aided by quality analysis tools. Nucleic Acids Research 25:4876-4882.
- Woose, C.R. 1987. Bacterial evolution. Microbiological Reviews 51:221-271.
- Woose, C.R., D. Yang, L. Mandelco and K.O. Stetter. 1990.
 The flexibacter-flavobacter connection. Systematic and Applied Microbiology 13:161-165.

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