## Draft genome sequences of *Sabulilitoribacter multivorans* KCTC 32326<sup>T</sup> and *Sabulilitoribacter arenilitoris* KCTC 52401<sup>T</sup>

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# Sabulilitoribacter multivorans KCTC 32326<sup>T</sup>와 Sabulilitoribacter arenilitoris KCTC 52401<sup>T</sup>의 유전체 염기서열 분석

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The draft genome sequences of *Sabulilitoribacter multivorans* KCTC 32326<sup>T</sup> and *Sabulilitoribacter arenilitoris* KCTC 52401<sup>T</sup> were determined using Illumuna Hiseq X-ten platform. The assembled genome of *Sabulilitoribacter multivorans* KCTC 32326<sup>T</sup> was composed of 13 contigs with a total length of 3,278,864 bp and G + C content was 32.7%. The assembled genome of *Sabulilitoribacter arenilitoris* KCTC 52401<sup>T</sup> comprises 41 contigs with a total length of 3,491,608 bp and the genomic DNA G+C content was 32.0%. Two genomes showed differences in carotenoid biosynthesis, nitrogen metabolism and carbohydrate-active enzymes. These genome information of type strains will be expected to be helpful for classification of other isolated *Sabulilitoribacter* strains.

**Keywords:** Sabulilitoribacter multivorans KCTC 32326<sup>T</sup>, Sabulilitoribacter arenilitoris KCTC 52401<sup>T</sup>, draft genome sequence

The genus *Sabulilitoribacter*, belong to the family *Flavo-bacteriaceae*, was first proposed by Park *et al.* (2013) and currently comprises 2 recognized species, including *Sabulilitoribacter multivorans* and *S. arenilitoris*. The members of the genus *Sabulilitoribacter* were isolated from sand of seashore, and were characterized by Gram-stain-negative, aerobic, catalase-

and oxidase-positive, non-motile, rod shaped. Herein, we report the draft genome sequences and annotations of *S. multivorans* KCTC 32326<sup>T</sup> and *S. arenilitoris* KCTC 52401<sup>T</sup>.

Sabulilitoribacter multivorans KCTC 32326<sup>T</sup> and S. arenilitoris KCTC 52401<sup>T</sup> were obtained from the Korean Collection for Type Cultures (KCTC) and revival and routinely cultured on marine agar 2216 (Difco) at 30°C for 3 days. Genomic DNA was extracted using MagAttract® HMW DNA kit (Qiagen) according to the manufacturer's instructions. The draft genome sequencing was performed on the Illumina Hiseq X-ten platform with TruSeq Nano DNA (350 bp insert size) library by Macrogen Inc. Raw reads were qualified by FastQC (version 0.11.5) and were assembled by SPAdes (version 3.13.0) (Bankevich et al., 2012) or Platanus-allee (version 2.2.2) (Kajitani et al., 2019). Orthologous average nucleotide identity (OrthoANI) and 16S rRNA gene sequence similarity were calculated using the Orthologous Average Nucleotide Identity Tool (http://www.ezbiocloud.net/tools/orthoani) and the online pairwise sequence alignment tool for the taxonomy (http:// www.ezbiocloud.net/tools/pairAlign), respectively (Yoon et al., 2017). The genome annotation was conducted using NCBI Prokaryotic Genome Annotation Pipeline (PGAP) (Tatusova et al., 2016), and additional function of the predicted genes were conducted by BlastKOALA with KEGG database (Kanehisa

Table 1. Genomic features of Sabulilitoribacter multivorans KCTC 32326<sup>T</sup> and Sabulilitoribacter arenilitoris KCTC 52401<sup>T</sup>

Property	Value	
	S. multivorans KCTC 32326 <sup>T</sup>	S. arenilitoris KCTC 52401 <sup>T</sup>
Genome assembly		
Assemble method	SPAdes (version 3.15.0)	Paltanus-allee (version 2.2.2)
Genome coverage	147.9X	149.3X
Sequencing technology	Illumina Hiseq X-ten	Illumina Hiseq X-ten
Genome features		
Genome size (bp)	3,278,864	3,491,608
Number of scaffolds	11	37
Number of contigs	13	41
G + C content (%)	32.7	32.0
Protein-coding genes (CDSs)	2,886	2,945
rRNA genes (5S, 16S, 23S)	1, 2, 1	1, 1, 1
tRNA genes	38	34
ncRNA genes	4	4
Pseudogenes	8	56
Accession number (GenBank)	JAKKDV000000000	JAKKDU000000000

et al., 2016) and EggNOG 5.0 (Huerta-Cepas et al., 2018). The dbCAN2 meta server was used for carbohydrate-active enzyme (CAZyme) annotation (Zhang et al., 2018). A Bacterial Pan Genome Analysis pipeline (BPGA) was used to find the number of core gene (Chaudhari et al., 2016).

The genome features of *S. multivorans* KCTC 32326<sup>T</sup> and *S.* arenilitoris KCTC 52401T are shown in Table 1. The draft genome of S. multivorans KCTC 32326<sup>T</sup> contained 11 scaffolds with a total length of 3,278,864 bp (N50 value, 766,194 bp). The G+C content was 32.7%, and 2,886 protein-coding genes, 4 rRNA genes, 38 tRNA genes, 4 non-coding RNA genes and 8 pseudo genes were predicted. The draft genome of S. arenilitoris KCTC 52401<sup>T</sup> was composed of 37 scaffolds with a total length of 3,491,608 bp (N50 value, 204,683 bp). The G + C content was 32.0%, and 2,945 protein-coding genes, 3 rRNA genes, 34 tRNA genes, 4 non-coding RNA genes and 56 pseudo genes were predicted. Average nucleotide identity (ANI) value and 16S rRNA gene sequence similarity value between S. multivorans KCTC 32326<sup>T</sup> and S. arenilitoris KCTC 52401<sup>T</sup> were 79.3% and 96.9%, respectively. The number of core genes in both strains was 1,863 and a unique gene of each strain accounted for about one-third.

While both genomes contained carotenoid biosynthesis related genes such as 15-cis-phytoene synthase CrtB, phytoene desaturase CrtI and  $\beta$ -carotene 3-hydroxylase CrtZ, lycopene  $\beta$ -

cyclase CrtY, β-carotene/zeaxanthin 4-ketolase CrtW and zeaxanthin glucosyltransferase CrtX were found only in S. arenilitoris KCTC 52401<sup>T</sup>. In the nitrogen metabolism, both genomes commonly contained nitrous-oxide reductase NosZ, but ferredoxin-nitrate reductase NarB and nitrite reductase (NADH) large subunit NirB were found only in S. multivorans KCTC 32326<sup>T</sup>. As a result of CAZyme annotation, S. multivorans KCTC 32326<sup>T</sup> involved 138 CAZymes (4 auxiliary activities, 3 carbohydrate-binding modules, 11 carbohydrate esterases, 55 glycoside hydrolases, 49 glycosyl transferases and 16 polysaccharide lyases), and S. arenilitoris KCTC 52401<sup>T</sup> involved 149 CAZymes (1 auxiliary activity, 2 carbohydratebinding modules, 8 carbohydrate esterases, 84 glycoside hydrolases, 53 glycosyl transferases and 1 polysaccharide lyases). Both genomes common encoded biopolymeric degradation related genes such as exodeoxyribonuclease III XthA,  $\beta$ glucosidase BglX,  $\alpha$ -glucosidase MalZ,  $\beta$ -galactosidase LacZ, xylose isomerase XylA and xylose kinase XylB. α-Amylase AmyA, pullanase PulA and xylan 1,4-β-xylosidase XynB were only found in S. multivorans KCTC 32326<sup>T</sup>, and chitinase was only found in S. arenilitoris KCTC 52401<sup>T</sup>. These genomic data might be useful for the genetic classification of the genus Sabulilitoribacter, and for study the ecological status and organic material circulation of the marine Flavobacteriaceae.

#### Nucleotide sequence accession numbers

The draft genome sequence of *Sabulilitoribacter multivorans* KCTC 32326<sup>T</sup> and *Sabulilitoribacter arenilitoris* KCTC 52401<sup>T</sup> has been deposited to GenBank under the accession number JAKKDV000000000 and JAKKDU0000000000, respectively. The version described in this paper are JAKKDV010000000 and JAKKDU0100000000.

#### 적 요

Sabulilitoribacter multivorans KCTC 32326<sup>T</sup>와 Sabulilitoribacter arenilitoris KCTC 52401<sup>T</sup>의 유전체 초안을 Illumina Hiseq X-ten platform을 사용하여 결정하였다. Sabulilitoribacter multivorans KCTC 32326<sup>T</sup>의 조립된 유전체는 전체 길이 3,278,864 bp의 13개 contig로 구성되었고 G+C 함량은 32.7% 이었다. Sabulilitoribacter arenilitoris KCTC 52401<sup>T</sup>의 조립된 유전체는 전체 길이 3,491,608 bp의 41개 contig로 구성되었고 G+C 함량은 32.0% 이었다. 이들 유전체는 carotenoid 생합성, 질소 대사 및 탄수화물 활성 효소 들에서 차이를 보였다. 이러한 type 균주의 유전 정보는 다른 분리된 Subulilitoribacter 균의 분류에 도움이 될 것으로 기대된다.

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## Conflict of Interest

The authors have no conflict of interest to report.

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