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Data in Brief

Whole genome sequencing and annotation of halophilic *Salinicoccus* sp. BAB 3246 isolated from the coastal region of Gujarat



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ABSTRACT

Salinicoccus sp. BAB 3246 is a halophilic bacterium isolated from a marine water sample collected from the coastal region of Gujarat, India, from a surface water stream. Based on 16sRNA sequencing, the organism was identified as *Salinicoccus* sp. BAB 3246 (Genebank ID: KF889285). The present work was performed to determine the whole genome sequence of the organism using Ion Torrent PGM platform followed by assembly using the CLC genomics workbench and genome annotation using RAST, BASys and MaGe. The complete genome sequence was 713,204 bp identified by with second largest size for *Salinicoccus* sp. reported in the NCBI genome database. A total of 652 degradative pathways were identified by KEGG map analysis. Comparative genomic analysis revealed *Salinicoccus* sp. BAB 3246 as most highly related to *Salinicoccus* halodurans H3B36. Data mining identified stress response genes and operator pathway for degradation of various environmental pollutants. Annotation data and analysis indicate potential use in pollution control in industrial influent and saline environment.

Specifications Organism/cell Salinicoccus sp. BAB 3246 line/tissue Sex Not applicable Sequencer or Ion Torrent PGM platform array type Data format Fasta complete genome Experimental Marine water sample factors Experimental Shotgun whole genome sequencing followed by features genome annotation using RAST, BASys and MaGe. Gujarat, India (21.672439 N 72.275925 E) Sample source location BioProject: PRJNA342322 Data submission RAST: genome ID 1437774.4 - Salinicoccus sp.

1. Direct link to deposited data

BAB-3246

BioProject: https://www.ncbi.nlm.nih.gov/bioproject/PRJNA342322.

2. Introduction

The genus *Salinicoccus*, belonging to family *Staphylococcaceae* was first proposed by Ventosa et al., (1990) and is defined as moderately halophilic, aerobic, Gram-positive, non-motile, non-sporulating, and heterotrophic cocci [1]. The genomic DNA G + C content of the species in this genus lies within the range of 46–51 mol%. Most species in genus *Salinicoccus* including *Salinicoccus albus*, *Salinicoccus carnicancri*, *Salinicoccus roseu*, *Salinicoccus halodurans*, *Salinicoccus luteus* have been found in salty environments, such as fermented foods, solar salterns, salt mines, salt lakes, and saline soils [1–7]. Alongside, genus *Salinicoccus* is also reported for production of Amylase, Protease, Gelatinase like enzymes in hyper saline environments [8].

The members of the *Salinicoccus* genus are abundant in the marine environments suggesting that they play important roles in marine ecosystems, such as the degradation of aromatic compounds and the biogeochemical cycles of carbon and sulfur [5]. *S. roseus* has been reported to exhibit high salinity and high lactate resistance [9]. *Salinicocci* have much importance in biotechnology applications such as serinemetabolism strategies to adapt to lactate stress [10]. In order to understand the genetic variability and industrial applications of those genes, genome sequencing and annotation of strain *Salinicoccus* sp. BAB 3246 was executed. The prime interest was to identify presence of

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Table 1

Summary of RAST annotation.

Genome	Salinicoccus sp. BAB 3246			
Size (bp)	7,13,204			
G + C content	49.1			
Number of coding sequences	1691			
Number of features	1762			
Number of subsystems	1009			
Number of RNAs	71			
Number of contigs	1			

distinctive enzymes for potential industrial applications.

3. Experimental design, materials and methods

The halophilic organism was isolated from marine water collected from surface streams of coastal region near Bhavnagar, Gujarat, India (latitude, longitude: 21.67 N, 72.27E). The isolation was performed by providing 15% Sodium Chloride containing Medium. The identification of Salinococcus sp. BAB 3246 was validated by 16 s rRNA sequencing and submitted to Genebank (accession no: KF889285.1). Furthermore, the DNA was extracted using Hi-Media Kit for Genomic DNA isolation Kit. The genome sequencing was performed using Ion Torrent PGM generating 15,26,815 sequencing reads. Initially all reads were subjected to preprocessing and conversion of BAM to fasta file format using Galaxy NGS: BamTools, online server using default parameters provided by the developer [11]. The genome data were assembled using CLC Genomic Workbench 5. The final whole genome assembly size was reported is 7,13,204 bp. The genome annotation was performed using RAST (Rapid Annotation using Subsystem Technology) [12], BASys (a web server for automated bacterial genome annotation) [13] and MaGe



Fig. 1. Subsystem category distribution.

(Microscope Genome Annotation) [14]. The RAST analysis revealed total 1691 coding sequences (Table 1). A total of 1009 subsystems were identified, including Stress Response (42), Sulfur Metabolism (4), Potassium metabolism (4) and Iron metabolism (1). However, the highest numbers of subsystems were observed for Amino Acids and Derivatives (159), Protein Metabolism (153) and Carbohydrate synthesis (150) (Fig. 1). KEGG pathway analysis was performing using seed viewer system of RAST. The KEGG map analysis revealed 652 pathways associated with only degradation of metabolites (Table 2).

The genome annotation using BASys annotate 955 genes amongst total 2330 genes reported in and automated mode. The amino acid composition was also examined using BASys (Fig. 2). The highest amino acid residue content was predicted for Leucine followed by Glycine, Glutamic acid and Alanine. Annotated data were displayed in the form of circular DNA as a genome browser map for easy representation of genome data (Fig. 3). The genome annotation using Microscope Genome Annotation identified 1772 Genomic Objects (without artifacts): CDS, 1326; fCDS, 358; misc_RNA, 16; rRNA, 12; tRNA, 60.

4. Quantitative comparison of coding sequences, rna and subsystem

The comparison of genome size for six different strains available in NCBI genome database revealed that, *S. halodurans* strain had the largest genome size of 2,778,379 bp followed by 873,136 bp, 713,204 bp, 679,606 bp, 461,933 bp and 342,819 bp respectively for *S. carnicancri* Crm, *Salinicoccus* sp. BAB 3246, *S. luteus* DSM 17002, *S. roseus and S. albus* DSM 19776 strain. A maximum of 2839 coding sequences was reported for *S. halodurans* followed by 1691, 863, 668, 449 and 334 respectively for *Salinicoccus* sp. BAB 3246, *S. carnicancri* Crm, *S. luteus* DSM 17002, *S. roseus* and *S. albus* DSM 19776 strain (Table 3).

Table 2

KEGG map analysis for degradation pathway.

No	Name of derivative	KEGG map	Salinicoccus sp. BAB-3246
1	1,1,1-Trichloro-2,2-bis(4-chlorophenyl)ethane (DDT) degradation	Tyrosine metabolism	4
2	1,2-Dichloroethane degradation	1,2-Dichloroethane degradation	1
		Glyoxylate and dicarboxylate metabolism	9
3	1,4-Dichlorobenzene degradation	Benzoate degradation via hydroxylation	2
		Glycolysis/gluconeogenesis	17
		Glyoxylate and dicarboxylate metabolism	9
4	1 and 2 Mathedran the land doors dation	Pyruvate metabolism	14
4	1- and 2-methymaphthalene degradation	1- and 2-Methylnaphthalene degradation	2
5	2 4-Dichlorobenzoate degradation	Benzoate degradation via hydroxylation	1
5	2,4-Dicinorobenzoare degradation	Naphthalene and anthracene degradation	1
6	3-Chloroacrylic acid degradation	3-Chloroacrylic acid degradation	1
0	o emotodelyne dela degradation	Pyruvate metabolism	14
7	Atrazine degradation	Atrazine degradation	1
		Folate biosynthesis	5
8	Benzoate degradation via CoA ligation	Benzoate degradation via CoA ligation	4
		Benzoate degradation via hydroxylation	2
		Butanoate metabolism	9
		Ethylbenzene degradation	1
		Phenylalanine metabolism	1
		Pyruvate metabolism	14
9	Benzoate degradation via hydroxylation	Benzoate degradation via CoA ligation	4
		Benzoate degradation via hydroxylation	2
		Caprolactam degradation	2
		Glycolysis/gluconeogenesis	17
		Naphthalene and anthracene degradation	1
		Phenylalanine metabolism	1
		Pyruvate metabolism	14
		Tryptophan metabolism	9
		Tyrosine metabolism	4
10	Biphenyl degradation	Benzoate degradation via CoA ligation	4
		Benzoate degradation via hydroxylation	2
		Glycolysis/gluconeogenesis	17
	Discharged A. das an destant	Pyruvate metabolism	14
11	Bisphenol A degradation	Benzoate degradation via hydroxylation	2
12	Caprolactam degradation	Convolution degradation via hydroxylation	2
12	Carbazale degradation	Caprolaciani degradation	2
15	Carbazole degradation	Benzoate degradation via budroxulation	4
		Chycolycis /gluconeogenesis	2
		Pyrivate metabolism	1/
		Tryptophan metabolism	9
14	Ethylbenzene degradation	Benzoate degradation via CoA ligation	4
		Ethylbenzene degradation	1
		Glycolysis/gluconeogenesis	17
		Propanoate metabolism	6
		Pyruvate metabolism	14
15	Fluorene degradation	Benzoate degradation via hydroxylation	2
		Glycolysis/gluconeogenesis	17
		Pyruvate metabolism	14
16	Fluorobenzoate degradation	Benzoate degradation via hydroxylation	2
17	Geraniol degradation	Geraniol degradation	3
		Valine, leucine and isoleucine degradation	9
18	Limonene and pinene degradation	Limonene and pinene degradation	3
19	Lysine degradation	Biotin metabolism	1
		Citrate cycle (TCA cycle)	14
		Lysine biosynthesis	5
		Lysine degradation	6
20	Naphthalene and anthracene degradation	Benzoate degradation via hydroxylation	2
		Naphthalene and anthracene degradation	1
		ryruvate metabolism	14
		Turcoino metabolism	9
21	Other always degradation	fyrosine metabolism Glucosphingolipid biosynthesis - gonglio agrice	4
21 22	Styrene degradation	Citrate cycle (TCA cycle)	14
<u></u>	organical degradation	Ethylhenzene degradation	1 1
		Glycolysis/gluconeogenesis	17
		Propanoate metabolism	17 6
		Pyruvate metabolism	14
23	Synthesis and degradation of ketone bodies	Butanoate metabolism	9
	2,	Fatty acid metabolism	5
		Glycolysis/gluconeogenesis	17
		Pyruvate metabolism	14
24	Tetrachloroethene degradation	Glyoxylate and dicarboxylate metabolism	9

(continued on next page)

Table 2 (continued)

No	Name of derivative	KEGG map	Salinicoccus sp. BAB-3246
		Pyruvate metabolism	14
25	Toluene and xylene degradation	Benzoate degradation via CoA ligation	4
		Benzoate degradation via hydroxylation	2
		Glycerolipid metabolism	3
		Glycolysis/gluconeogenesis	17
		Pyruvate metabolism	14
26	Trinitrotoluene degradation	Trinitrotoluene degradation	1
27	Valine, leucine and isoleucine degradation	Biosynthesis of type II polyketide backbone	1
		Citrate cycle (TCA cycle)	14
		Propanoate metabolism	6
		Pyrimidine metabolism	17
		Valine, leucine and isoleucine biosynthesis	12
		Valine, leucine and isoleucine degradation	9
28	Gamma-Hexachlorocyclohexane degradation	Benzoate degradation via hydroxylation	2
		Citrate cycle (TCA cycle)	14
		Glyoxylate and dicarboxylate metabolism	9
		Naphthalene and anthracene degradation	1







Fig. 3. Genome browser map for Salinicoccussp. BAB 3246.

Quantitative comparison of coding sequence, RNA and subsystem.

Genome	Size (bp)	G + C content	Coding sequences	Features	RNAs	Subsystems	BioProject
Salinicoccus sp. BAB_3246	713,204	49.1	1691	1762	71	202	PRJNA342322
Salinicoccus roseus	461,933	49.9	449	459	10	80	PRJNA272357
Salinicoccus carnicancri Crm	873,136	47.6	863	909	46	138	PRJNA175941
Salinicoccus albus DSM 19776	342,819	45.2	334	334	0	77	PRJNA185242
Salinicoccus luteus DSM 17002	679,606	49.7	668	669	1	114	PRJNA235106
Salinicoccus halodurans	2,778,379	44.5	2839	2912	73	388	PRJNA282445

5. Nucleotide sequence accession number

The complete sequence of *Salinicoccus* sp. BAB 3246 genome can be accessed under the NCBI BioProject: PRJNA342322.

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