High quality draft genome sequence of *Staphylococcus cohnii* subsp. *cohnii* strain hu-01

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**Keywords:** *Staphylococcus cohnii* subsp. *cohnii*, genome, Hiseq2000

*Staphylococcus cohnii* subsp. *cohnii* belongs to the family *Staphylococcaceae* in the order Bacillales, class Bacilli and phylum Firmicutes. The increasing relevance of *S. cohnii* to human health prompted us to determine the genomic sequence of *Staphylococcus cohnii* subsp. *cohnii* strain hu-01, a multidrug-resistant isolate from a hospital in China. Here we describe the features of *S. cohnii* subsp. *cohnii* strain hu-01, together with the genome sequence and its annotation. This is the first genome sequence of the species *Staphylococcus cohnii*.

**Introduction**

*Staphylococcus cohnii* belongs to the Coagulase-Negative Staphylococci group. It was described by Schleifer and Kloos (1975) and was named for Ferdinand Cohn, a German botanist and bacteriologist [1]. Recently, more cases of *Staphylococcus cohnii* infection have been reported in the literature. This organism may be responsible for brain abscess, pneumonia, acute cholecystitis, endocarditis, bacteremia, urinary tract infection and septic arthritis [2]. *S. cohnii* is comprised of two subspecies that are defined on the basis of their phenotypic characteristics: *Staphylococcus cohnii* subsp. *cohnii* and *Staphylococcus cohnii* subsp. *urealyticus* [3]. *S. cohnii* subsp. *cohnii* is a Gram-positive coccus, coagulase negative and catalase positive, that behaves like a commensal mucocutaneous bacterium [4]. It has more frequently been isolated in hospital than in non-hospital environments [2]. Here we report this draft genome of *S. cohnii* subsp. *cohnii* strain hu-01, the first genome of this species to be sequenced.

**Classification and features**

Strain hu-01 was isolated from a hospital environment in Zhejiang province, China, in October 2012. It is a Gram-positive, coccus-shaped bacterium that can grow on 5% sheep blood enriched Columbia agar (BioMérieux, Marcyl’Etoile, France) at 37°C. Growth occurs under either aerobic or anaerobic conditions. The optimum temperature for growth is 37 ºC, with a temperature range of 15-45 ºC (Table 1). Cell morphology, motility and sporulation were examined by using transmission electron (H-600, Hitachi) microscopy. Cells of strain hu-01 are coccoidal, 0.6 to 1.2 μm in diameter, occurring predominantly singly or in pairs (Figure 1 and Figure 2).
Figure 1. Gram staining of *S. cohnii subsp. cohnii* strain hu-01

Figure 2. Transmission electron micrograph of cells of strain hu-01. Bar: 0.5 µm
Table 1. Classification and general features of *S. cohnii subsp. cohnii* strain hu-01 according to the MIGS recommendations [9].

| MIGS ID   | Property                  | Term                                                                 | Evidence code<sup>a</sup> |
|-----------|---------------------------|----------------------------------------------------------------------|---------------------------|
|           | Domain                    | *Bacteria*                                                           | TAS [20]                  |
|           | Phylum                    | *Firmicutes*                                                         | TAS [21-23]               |
|           | Class                     | *Bacilli*                                                           | TAS [24,25]               |
|           | Current classification    | Order *Bacillales*                                                   | TAS [26,27]               |
|           |                           | Family *Staphylococcaceae*                                           | TAS [24,28]               |
|           |                           | Genus *Staphylococcus*                                              | TAS [26,29-31]            |
|           |                           | Species *Staphylococcus cohnii subsp. cohnii*                       | TAS [1,3]                 |
|           | Gram stain                | Positive                                                            | IDA                       |
|           | Cell shape                | coccus                                                              | IDA                       |
|           | Motility                  | Nonmotile                                                           | IDA                       |
|           | Sporulation               | Nonsporulating                                                      | IDA                       |
|           | Temperature range         | 15-45°C                                                             | IDA                       |
|           | Optimum temperature       | 37°C                                                                | IDA                       |
| MIGS-6.3  | Salinity                  | Tolerates 10% NaCl                                                   | IDA                       |
| MIGS-22   | Oxygen                    | Facultatively anaerobic D-mannitol, fructose, trehalose              | IDA                       |
|           | Carbon source             |                                                                      | IDA                       |
|           | Energy source             | fructose, trehalose                                                 | IDA                       |
| MIGS-6    | Habitat                   | Hospital environment                                               | IDA                       |
| MIGS-15   | Biotic relationship       | Free living                                                         | IDA                       |
| MIGS-14   | Pathogenicity             | Opportunistic pathogen                                              | IDA                       |
|           | Isolation                 | Hospital                                                             | IDA                       |
| MIGS-4    | Geographic location       | Hangzhou, China                                                     | IDA                       |
| MIGS-5    | Sample collection time    |                                                                      | IDA                       |
| MIGS-4.1  | Latitude                  | October, 2012                                                       | IDA                       |
| MIGS-4.2  | Longitude                 | 30°16′N                                                             | IDA                       |
| MIGS-4.3  | Depth                     | unknown                                                             | IDA                       |
| MIGS-4.4  | Altitude                  | 50 (meters)                                                         | IDA                       |

<sup>a</sup>Evidence codes-IDA: Inferred from Direct Assay; TAS: Traceable Author Statement (i.e., a direct report exists in the literature); NAS: Non-traceable Author Statement (i.e., not directly observed for the living, isolated sample, but based on a generally accepted property for the species, or anecdotal evidence). These evidence codes are from the Gene Ontology project [32]. If the evidence code is IDA, then the property should have been directly observed, for the purpose of this specific publication, for a live isolate by one of the authors, or an expert or reputable institution mentioned in the acknowledgements.
Staphylococcus cohnii subsp. cohnii

Figure 3. Phylogenetic tree depicting the relationship between S. cohnii subsp. cohnii strain hu-01 and other members of the genus Staphylococcus. The strains and their corresponding Genbank accession numbers are shown following the organism name and indicated in parentheses. The phylogenetic tree uses 16S rRNA gene sequences aligned by the CLUSTALW [7], and phylogenetic inferences were made using Neighbor-joining method based on Kimura 2-parameter model within the MEGA5 software [8] and rooted with Bacillus subtilis subsp. subtilis. Bootstrap consensus trees were inferred from 100 replicates, only bootstrap values > 50% were indicated.

Comparative 16S rRNA gene sequence analysis by BLASTN [5,6] using the NCBI-NR/NT database revealed 94-99% sequence similarity to members of genus Staphylococcus. Neighbor-Joining phylogenetic analysis based on Kimura 2-parameter model indicated the strain hu-01 is most closely related the strain Staphylococcus urealyticus (AB009936.1) (Figure 3).

Biochemical features were tested by using two automated systems, the Vitek2 Compact (bioMérieux, Marcy l’Etoile, France) and Phoenix 100 ID/AST system (Becton Dickinson Company [BD], Sparks, Maryland, USA). Positive reactions were obtained for D-fructose, trehalose, D-gluconic acid and D-mannitol. Negative reactions were observed for glucose, D-trehalose, D-sucrose, maltose, urea, cellobiose, glucoside, D-tagatose and maltotriose. This strain was susceptible to gentamicin, ciprofloxacin, levofloxacin, moxifloxacin, quinupristin, linezolid, vancomycin, tetracycline, tigecycline, nitrofurantoin, rifampicin, trimethoprim and resistant to cefoxitin, benzylpenicillin, oxacillin, erythromycin, clindamycin.
Genome sequencing information

Genome project history

*S. conhii subsp. cohnii* strain hu-01 was selected for sequencing because of its increasing relevance to human health. The strain was isolated from a hospital environment in China. This whole genome shotgun project of *S. conhii subsp. cohnii* strain hu-01 was deposited at DDBJ/EMBL/GenBank under the accession AYOS00000000. Table 2 presents the project information and its association with MIGS version 2.0 compliance [9].

Table 2. Project information

| MIGS ID | Property                  | Term                      |
|---------|---------------------------|---------------------------|
| MIGS-31 | Finishing quality         | High-quality draft        |
| MIGS-28 | Libraries used            | One pair-end 500 bp library|
| MIGS-29 | Sequencing platforms      | Illumina HiSeq 2000       |
| MIGS-31.2| Fold coverage            | 150×(based on 500 bp library) |
| MIGS-30 | Assemblers                | Velvet 1.2.07             |
| MIGS-32 | Gene calling method       | Glimmer 3.0               |
|         | Genbank ID                | AYOS00000000             |
|         | Genbank Date of Release   | Jan 06, 2014             |
|         | GOLD ID                   | Gi0062613                |
| MIGS-13 | Project relevance         | Biotechnology, Pathway, Pathogenic |

Growth conditions and DNA isolation

*S. conhii subsp. cohnii* strain hu-01 was grown aerobically on Columbia blood agar base, at 37°C for 24h. Genomic DNA was extracted using the DNeasy blood and tissue kit (Qiagen, Germany), according to the manufacturer’s recommended protocol. The quantity of DNA was measured by the NanoDrop Spectrophotometer and Cubit. Then 10μg of DNA was sent to the State Key Laboratory for Diagnosis and Treatment of Infectious Disease at Zhejiang University for sequencing on a Hiseq2000 (Illumina, CA) sequencer.

Genome sequencing and assembly

One DNA library was generated (500 bp insert size, with the Illumina adapter at both ends, detected by Agilent DNA analyzer 2100), then sequencing was performed by using an Illumina HiSeq 2000 genomic sequencer, with a 2×100 pair end sequencing strategy. A total of 1,103 Mbp of sequence data was produced which was assessed for quality by the following criteria: 1) Reads linked to adapters at both end were considered as sequencing artifacts then removed. 2) Bases with a quality index lower than Q20 at both ends were trimmed. 3) Reads with ambiguous bases (N) were removed. 4) Single qualified reads were discarded (In this situation, one read is qualified but its mate is not). A total of 867.94 M clean filtered reads were assembled into scaffolds using the Velvet version 1.2.07 with parameters "-scaffolds no" [10], then we used a PAGIT flow [11] to prolong the initial contigs and correct sequencing errors. to arrive at a set of improved scaffolds.

Genome annotation

Predict genes were identified using Glimmer version 3.0 [12], tRNAscan-SE version 1.21 [13] was used to find tRNA genes, whereas ribosomal RNAs were found by using RNAmer version 1.2 [14]. To annotate predicted genes, we used HMMER version 3.0 [15], with parameters 'hmmscan -E 0.01 -domainE 0.01' to align genes against Pfam version 27.0 [16] (only pfam-A was used) to find genes with conserved domains. The KAAS server [17] was used to assign translated amino acids (with genetic code table 11) into KEGG Orthology with SBH (single-directional best hit) method. Translated genes were aligned with the COG database using NCBI blastp (hits should have scores no less than 60, e-value is no more than 1e-6). To find genes with hypothetical or putative function, we aligned genes against NCBI nucleotide database (nt database was downloaded at Sep 20, 2013) by using NCBI blastn, only if hits have an identity of no less than 0.95, coverage no less than 0.9, and the reference gene had an annotation of putative or hypothetical. To define genes with signal peptide, we use signalp version 4.1 [18] to identify genes with signal peptide with default parameters except "-t gram+ ". TMHMM2.0 [19] was used to identify genes with transmembrane helices.
**Staphylococcus cohnii subsp. cohnii**

**Genome properties**

The draft genome sequence of *S. cohnii subsp. cohnii* strain hu-01 revealed a genome size of 5,761,489 bp and a G+C content of 34.85% (521 scaffolds with N50 is 39,926 bp). These scaffolds contain 5,820 coding sequences (CDSs), 61 tRNAs (excluding 6 Pseudo tRNAs) and incomplete rRNA operons (10 small subunit rRNA and 3 large subunit rRNAs). A total of 1,840 protein-coding genes were assigned as putative function or hypothetical proteins. 3,734 genes were categorized into COGs functional groups. The properties and the statistics of the genome are summarized in Table 3 and Table 4.

| Attribute | Value      | % of total |
|-----------|------------|------------|
| Genome size (bp) | 5,761,489 |            |
| DNA coding region (bp) | 4,751,472 | 82.469     |
| DNA G+C content (bp) | 1,697,984 | 29.471     |
| Total genes | 5,833      |            |
| RNA genes | 13         | 0.221      |
| Protein-coding genes | 5,820      | 99.777     |
| Genes with function prediction | 1,840      | 31.544     |
| Genes assigned to COGs | 3,734      | 64.015     |
| Genes assigned to Pfam domains | 4,943      | 84.741     |
| Genes with signal peptides | 431        | 7.388      |
| Genes with transmembrane helices | 1,629      | 27.927     |

The total is based on either size of the genome in base pairs or total number of genes in the annotated genome.

**Table 3. Genome statistics of *S. cohnii subsp. cohnii* strain hu-01**

| Code | Value | % of total |
|------|-------|------------|
| J    | 230   | 3.95       |
| K    | 452   | 7.77       |
| L    | 184   | 3.16       |
| B    | 3     | 0.05       |
| D    | 72    | 1.24       |
| V    | 187   | 3.21       |
| T    | 238   | 4.09       |
| M    | 254   | 4.36       |
| N    | 70    | 1.20       |
| Z    | 1     | 0.02       |
| W    | 1     | 0.02       |
| U    | 57    | 0.98       |
| O    | 147   | 2.53       |
| C    | 292   | 5.02       |
| G    | 384   | 6.60       |
| E    | 640   | 11.0       |
| F    | 140   | 2.41       |
| H    | 234   | 4.02       |
| I    | 165   | 2.84       |
| P    | 389   | 6.68       |
| Q    | 197   | 3.38       |
| R    | 841   | 14.45      |
| S    | 403   | 6.92       |
| --  | 483   | 8.30       |
| --  | 1603  | 27.54      |

For some genes, qualified alignments can occur with several genes belonging to different COG categories. In such cases only the best match to a single COG category is considered. b) The total is based on the total number of protein coding genes (5,820) in the annotated genome. c) These genes have alignments with reference genes archived in COG, but these reference genes do not have COG categories. d) Genes without a qualified hit to a reference genes.
Conclusion

*Staphylococcus cohnii* ssp. *cohnii* are part of the normal flora of human skin and mucous membranes which, in particular conditions, may become opportunistic pathogens [4]. The genome sequence of *Staphylococcus cohnii subsp. cohnii* strain hu-01 will provide the basis to elucidate the molecular principles of host colonization and insight into the genetic background of this organism’s pathogenesis.

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Acknowledgements

We thank Qiang Ye and Li Liang, Chinese Center of Medical Culture Collections /National Institutes for Food and Drug Control for providing the Biochemical features. This study was supported by the National Basic Research Program of China (973 program) (No. 2013CB531401) and the key Program of the National Basic Research Foundation of China (No. 81330011).
Staphylococcus cohnii subsp. cohnii

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