Genome-wide analysis reveals the emergence of multidrug resistant *Stenotrophomonas acidaminiphila* strain SINDOREI isolated from a patient with sepsis

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*Stenotrophomonas acidaminiphila*, the most recent reported species in genus *Stenotrophomonas*, is a relatively rare bacteria and is an aerobic, glucose non-fermentative, Gram-negative bacterium. However, little information of *S. acidaminiphila* is known to cause human infections. In this research, we firstly reported a multidrug-resistant strain *S. acidaminiphila* SINDOREI isolated from the blood of a patient with sepsis, who was dead of infection eventually. The whole genome of strain SINDOREI was sequenced, and genome comparisons were performed among six closely related *S. acidaminiphila* strains. The core genes (2,506 genes) and strain-specific genes were identified, respectively, to know about the strain-level diversity in six *S. acidaminiphila* stains. The presence of a unique gene (narG) and essential genes involved in biofilm formation in strain SINDOREI are important for the pathogenesis of infections. Strain SINDOREI was resistant to trimethoprim/sulfamethoxazole, ciprofloxacin, ofloxacin, cefepime, ceftazidime, and aztreonam. Several common and specific antibiotic resistance genes were identified in strain SINDOREI. The presence of two sul genes and exclusive determinants GES-1, aadA3, qacL, and cmlA5 is responsible for the resistance to multidrug. The virulence factors and resistance determinants can show the relationship between the phenotype and genotype and afford potential therapeutic strategies for infections.

**KEYWORDS**

*Stenotrophomonas acidaminiphila*, sepsis, genome sequencing, genome comparisons, virulence factors, multidrug resistant, human infection
Introduction

The genus *Stenotrophomonas*, which was first described with the type species *Stenotrophomonas maltophilia* (Palleroni and Bradbury, 1993), currently comprises 16 validly described species (Wang et al., 2018). Members of the genus *Stenotrophomonas* demonstrated great metabolic versatility and intraspecific heterogeneity (Ryan et al., 2009). The most recently reported species, *Stenotrophomonas acidaminiphila*, is an aerobic, glucose non-fermentative, Gram-negative bacterium, which is initially isolated from a petrochemical wastewater treated by an upflow anaerobic sludge blanket (UASB) reactor (Assih et al., 2002), and strains of *S. acidaminiphila* occur ubiquitously in the environment (Virues and Ochoa-Sánchez, 2015; Patil et al., 2016; Huang et al., 2018; Huyan et al., 2020). However, the information on the characteristic of strains is still limited. To date, the reported strains of *S. acidaminiphila* were limited, especially clinical isolates. Additionally, the further genomic analysis was necessary to dig the features of *S. acidaminiphila*.

*Stenotrophomonas* species, especially *S. maltophilia* and *S. acidaminiphila*, can cause human infections (Looney et al., 2009; Huang et al., 2018). Multidrug-resistant (MDR) strains of *Stenotrophomonas* are associated with a high rate of mortality in immunocompromised patients (Paez and Costa, 2008). In the genomes of *Stenotrophomonas*, various genes encoding virulence determinants are involved in the infections (Trifonova and Strateva, 2019). Surveillance of the presence of virulence genes is important to supplement knowledge about the pathogenesis of infections (Windhorst et al., 2002; de Oliveira-Garcia et al., 2003).

Trimethoprim/sulfamethoxazole (TMP/SMX) was considered as the first-line therapy (Abbott and Peleg, 2015; Kumar et al., 2020), but was plagued by increasing resistance worldwide (Al-Jasser, 2006; Tan et al., 2008; Looney et al., 2009). Fluoroquinolones and β-lactam drugs have been used as potential alternative antibiotics to TMP/SMX for *Stenotrophomonas* infections. However, recent studies have revealed a trend in decreasing susceptibility (Wei et al., 2016; Ko et al., 2019). The strains’ intrinsic and acquired mechanisms of antibiotic resistance to most antibiotics limited the antimicrobial options for *Stenotrophomonas* infections.

The molecular mechanisms involved in its extensive antimicrobial resistance include efflux pumps and encoded genes. The *sul1* gene, *sul2* gene and *dfrA* gene are well known to be responsible for resistance to TMP/SMX (Barbolla et al., 2004; Domínguez et al., 2019). Two chromosomal-mediated β-lactamases, namely L1 and L2, with several regulatory genes, such as *ampR*, *ampN* and *ampG*, are associated with β-lactam resistance (Okazaki and Avison, 2008; Huang et al., 2010; Lin et al., 2011). A chromosomally encoded *qnr* gene protects both gyrase and topoisomerase IV from quinolones and confers resistance to fluoroquinolone (Ko et al., 2019). Moreover, efflux pumps are shown to be associated with resistance to multidrug and are classified to five families, namely the resistance-nodulation-cell-division (RND) family, the major facilitator superfamilies (MFS), the small multidrug resistance (SMR) family, the ATP binding cassette (ABC) family, and the multidrug and toxic compound extrusion (MATE) family (Putman et al., 2000).

To our best knowledge, *S. acidaminiphila* was mostly isolated from aquatic environments. It is worth noting that the first clinical isolate *S. acidaminiphila* SUNE0 (Huang et al., 2018) was isolated from the bile of a cholangiocarcinoma patient with obstructive jaundice and cholangitis and was found resistant to sulfamethoxazole and imipenem based on the antimicrobial susceptibility testing. Meanwhile, the comparisons and analysis of whole genomes aid the identification of resistant determinants to develop the antimicrobial strategies.

In this study, the first MDR clinical isolate, *S. acidaminiphila* SINDOREI, was cultured and isolated from the blood of a patient with sepsis in China. The complete genome of SINDOREI was assembled to highlight the virulence factors and resistant genes characterizing the specific isolates. Additionally, we profiled the adaptive changes in *S. acidaminiphila* SINDORI through the characteristic of genome and genomic comparisons of six *S. acidaminiphila* strains.

Materials and methods

Bacterial isolation and culture conditions

Pure strain SINDOREI was cultured from the blood of a patient with sepsis. The 51-year-old man was transferred to Xiangya Hospital of Central South University (Changsha, China) on March 16, 2020, complaining of intermittent fever and full-body pain for more than 1 month without obvious causes. Based on physical examination, laboratory test results and related examinations including bone marrow (BM) aspirate smear, flow cytometry and RT-PCR, etc., a diagnosis of sepsis was made. The antibiotic treatment did not improve his condition. The patient had a continuous high fever (39.5°C) with some new symptoms such as chills, high fever, appetite, fatigue, and shortness of breath and eventually died for multiple organ failure. The detailed medical record of the patient was shown in Supplementary Table S1. The blood samples of this patient were inoculated on nutrient agar with 5% sheep blood and incubated aerobically at 37°C overnight for three times. Only smooth, opaque and yellow colonies showing clear zones were isolated. Then, the isolates were conducted species identification by using MALDI Biotyper (Bruker, Germany) and returned no match in the database. At last, the purified isolate was further performed the whole genome sequencing and was classified as *S. acidaminiphila*.

Antimicrobial susceptibility testing

Antimicrobial susceptibility testing was performed by VITEK2 system (bioMérieux, France) for minimum inhibitory concentration (MIC) according to the manufacturer’s instructions. MIC breakpoint was determined referring to the clinical and laboratory standards institute (CLSI) guidelines (M100, 30th Ed.; Wayne, 2020).
Phylogenetic analysis based on the 16S rRNA sequence

The phylogenetic tree based on the 16S rRNA sequence which recovered from the genome of SINDOREI were constructed using MEGA version 11 software using maximum-likelihood method with 1,000 bootstrap replications (Tamura et al., 2021). The phylogenomic analysis based on whole genomes of the members of genus *Stenotrophomonas* were performed using the genome taxonomic database toolkit (GTDB-Tk; Chaumeil et al., 2019) and iqtree (Nguyen et al., 2015).

Genome sequencing, assembly and annotation

The genomic DNA of *S. acidaminiphila* SINDOREI was extracted using cetyltrimethylammonium bromide (CTAB)-based methods. Then genomic DNA was randomly fragmented and was selected to average size of 200–400 bp. Adaptors were ligated to the ends of fragments by PCR assay. PCR products were processed into the sequencing library. The qualified BGI-seq libraries were sequenced on a BGISEQ-500 platform with read length of PE100. The Nanopore libraries were prepared and sequenced according to the manufacturer’s instructions (Oxford Nanopore, Oxford, UK) and sequenced on MinION with flowcell version of R9.4.1. The short reads and nanopore long reads were assembled using software Unicycler (Wick et al., 2017). The genes were predicted using Glimmer software (Delcher et al., 2007) and were functionally annotated by querying eggNOG, NR, Pfam and Swiss-Prot databases to obtain the corresponding annotations.

SNP/InDel identification

Firstly, the genomes were broken into 500,000 reads with length of 150bp using wgsim (v1.11) with parameters of (-e 0 -r 0 -R 0 -X 0). Then, these reads were mapped to SINDOREI genome using bwa (0.7.17-r1188) with default parameters. Next, the SNPs were identified by bcftools’ and variations with quality <20 were filtered.

Average nucleotide identity values and digital DNA–DNA hybridization values

Average nucleotide identity (ANI) values between any two genomes were calculated using fastANI (Jain et al., 2018). The digital DNA–DNA hybridization (dDDH) values were obtained by means of genome-to-genome distance calculator via GGDC 3.0 using Formula 2.

Genome comparisons and identification of core and strain-specific genes

For genome comparisons, six genomes of the *S. acidaminiphila* strains, including a type strain and other five cultured strains, were downloaded from the NCBI database (Table 1). The clusters of homologous genes among the investigated genome sequences were determined using OrthoMCL (Fischer et al., 2011). The numbers of unique core genes and strain-specific genes of all isolates were mapped. The strain-specific genes that are present in strain SINDOREI were annotated through the clusters of orthologous genes (COG) database using software EggNOG-mapper v2 (Cantalapiedra et al., 2021).

Virulence factor and resistance genes comparisons

To analyze the virulence factors of the members of species *S. acidaminiphila*, virulence factors database (VFDB) was used (Liu et al., 2022). The protein-coding sequences were aligned against the comprehensive antibiotic resistance database (CARD; Alcock et al., 2020) and resistance-related genes were analyzed. Comparisons of specific genes related to resistance genes and efflux pumps between *S. acidaminiphila* strains were proceeded using local BLASTP alignment.

Phylogeny based on the amino acid of sul genes

The phylogenetic tree based on the amino acid of *sul* genes was also constructed by MEGA version 11 software, using maximum-likelihood method with 1,000 bootstrap replications (Tamura et al., 2021). The amino acid sequences of dihydropteroate synthase from *Stenotrophomonas* strain were downloaded from the National Center for Biotechnology Information.

Results

Genomic features of *Stenotrophomonas acidaminiphila* SINDOREI

The complete genome of SINDOREI was assembled into one circular chromosome with a total size of 3,996,619 bp and a GC content of 68.73%. SINDOREI genome contained no plasmid, and the details of the genome were listed in Supplementary Tables S2, S3. The genome of SINDOREI were functionally annotated in databases: eggNOG (Supplementary Table S4), NR

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1. https://github.com/lh3/wgsim
2. https://github.com/lh3/bwa
3. https://github.com/samtools/bcftools
TABLE 1  The genomic information of the isolated strains of *Stenotrophomonas acidaminiphila*.

| Strain            | Genome size (bp) | GC content (%) | Genes (total) | CDSs (total) | Isolation source                      | Country | Accession number |
|-------------------|------------------|----------------|---------------|--------------|---------------------------------------|---------|------------------|
| *S. acidaminiphila* SINDOREI | 3,996,619        | 68.73          | 3,623         | 3,536        | Blood of septic patient               | China   | PRJCA009493      |
| *S. acidaminiphila* SUNEO      | 3,660,864        | 69.75          | 3,300         | 3,226        | Human bile                            | China   | GCA_002951995.1  |
| *S. acidaminiphila* ZAC14D2_NAIMI4_2 | 4,138,397       | 68.48          | 3,752         | 3,677        | Sediments of polluted river           | Mexico  | GCA_001314305.1  |
| *S. acidaminiphila* T25-65     | 3,915,662        | 68.96          | 3,556         | 3,481        | Aerobic biofilm reactors with antibiotics | China   | GCA_014076435.1  |
| *S. acidaminiphila* T0-18      | 3,848,207        | 69.17          | 3,480         | 3,406        | Aerobic biofilm reactors with antibiotics | China   | GCA_014109845.1  |
| *S. acidaminiphila* JCM13310   | 3,942,520        | 68.81          | 3,598         | 3,425        | Sludge from anaerobic chemical wastewater reactor | Mexico  | GCA_001431595.1  |

(Supplementary Table S5), Pfam (Supplementary Table S6) and Swiss-Prot (Supplementary Table S7). A total of 3,623 genes, including 3,536 coding sequences (CDSs), were predicted in the genome, along with 68 transfer RNA (tRNA genes) and nine ribosomal RNA (rRNA genes). The strain SINDOREI genome was profiled as a circular map, exhibiting CDSs, virulent genes, GC plot, and GC skew (Figure 1).

Phylogenetic analysis of members of the genus *Stenotrophomonas*

Firstly, a 16S rRNA phylogenetic tree of strain SINDOREI and other available type strains of genus *Stenotrophomonas* were constructed using the maximum-likelihood method (Supplementary Figure S1). Strain SINDOREI was clustered to the branch of *S. acidaminiphila* with a bootstrap consistency of 99%. In addition, the 16S rRNA identities between SINDORE and other *S. acidaminiphila* strains was over 99.68%, indicating the strain SINDOREI belongs to *S. acidaminiphila*.

The phylogenomic tree was constructed and revealed that strain SINDOREI has the closest relationship with strain JCM13310 and strain T25-65, which was consistent with the 16S rRNA tree (Figure 2). Whole genome comparison identified the least SNPs/Indels (11,650) between SINDOREI and T25-65 (Supplementary Table S8). Meanwhile, the ANI and dDDH values between *S. acidaminiphila* SINDOREI and *S. acidaminiphila* T25-65 are up to 99.9% and 94.1%, respectively (Figure 3; Supplementary Table S9).

Comparative genome analysis of *Stenotrophomonas acidaminiphila*

Six genomes of *S. acidaminiphila* strains were employed for the comparative genome analysis (Table 1). Only SINDOREI and SUNEO were isolated from clinical specimens. The largest genome size was from ZAC14D2_NAIMI4_2 (4,138,397 bp), followed by strain SINDOREI. There were 2,506 orthogroups found in each strain (Figure 4), which represented the set of non-redundant core genes of six strains (Figure 5). The second highest orthogroups (292 orthogroups) were uniquely observed in SINDOREI, JCM13310, T25-65, T0-18 and ZAC14D2_NAIMI4_2, except SUNEO. Further analysis of pair-wise comparisons demonstrated that strain SINDOREI share 3,166, 3,105, 3,084, 3,028, and 2,691 orthogroups with T0-18, JCM 13310, T25-65, ZAC14D2_NAIMI4_2 and SUNEO, respectively (Supplementary Figure S2). Strain ZAC14D2_NAIMI4_2 had the most strain-specific genes (293 genes) as shown in Figure 5A, followed by strain JCM 13310 (216 genes). The ratio of specific genes in strain ZAC14D2_NAIMI4_2 were 3,166, 3,105, 3,084, 3,028, and 2,691 orthogroups with T0-18, JCM 13310, T25-65, ZAC14D2_NAIMI4_2 and SUNEO, respectively (Supplementary Figure S2). Strain ZAC14D2_NAIMI4_2 had the most strain-specific genes (293 genes) as shown in Figure 5A, followed by strain JCM 13310 (216 genes). The ratio of specific genes in strain ZAC14D2_NAIMI4_2 were 3,166, 3,105, 3,084, 3,028, and 2,691 orthogroups with T0-18, JCM 13310, T25-65, ZAC14D2_NAIMI4_2 and SUNEO, respectively (Supplementary Figure S2). Strain ZAC14D2_NAIMI4_2 had the most strain-specific genes (293 genes) as shown in Figure 5A, followed by strain JCM 13310 (216 genes). The ratio of specific genes in strain ZAC14D2_NAIMI4_2 were 3,166, 3,105, 3,084, 3,028, and 2,691 orthogroups with T0-18, JCM 13310, T25-65, ZAC14D2_NAIMI4_2 and SUNEO, respectively (Supplementary Figure S2). Strain ZAC14D2_NAIMI4_2 had the most strain-specific genes (293 genes) as shown in Figure 5A, followed by strain JCM 13310 (216 genes). The ratio of specific genes in strain ZAC14D2_NAIMI4_2 were 3,166, 3,105, 3,084, 3,028, and 2,691 orthogroups with T0-18, JCM 13310, T25-65, ZAC14D2_NAIMI4_2 and SUNEO, respectively (Supplementary Figure S2).

Virulence factors associated with infections

Virulence factors are components, produced by bacterial cells, which generally cause damages to the host by increasing adhesion, facilitating colonization and invasion into eukaryotic cells, escaping the host immune responses and providing the essential nutrient (Casadevall and Pirofski, 2009). The presence of various virulence genes was investigated in the strain SINDOREI genome and the comparison of virulence genes between related strain genomes was conducted. The results revealed that the virulence genes in *S. acidaminiphila* genomes are classified into 11 categories according to virulence factor (VF) category, including motility (flagella), adherence (Type IV pili and non-pili adhesins), biofilm formation, immune modulation [lipooligosaccharides (LOS), capsule, and lipopolysaccharide (LPS)], antimicrobial activity/competitive advantage, stress survival, nutritional/
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metabolic factor, regulation, exotoxin, effector delivery system, and others. The virulence genes of \textit{S. acidaminiphila} genomes was shown in Figure 6 and the details were listed in Supplementary Table S11. The genomes of \textit{S. acidaminiphila} strains included the same VF categories, but the presence and distribution of genes were different. It is worth noting that the \textit{narGH} operon only found in strain SINDOREI is responsible for nitrate reductase, which is identified as an important virulence factor for many bacterial infections, such as \textit{Mycobacterium tuberculosis} and \textit{Pseudomonas aeruginosa} (Palmer et al., 2007; Sohaskey and Modesti, 2009).

Biofilm formation, which is a mixture of cells, polysaccharides, nucleic acids, lipids and proteins, provides resistance to various antimicrobial drugs and to host immune defense of bacterial virulence. Biofilm-related infections represent more than 60% of all microbial infections in humans. In the genome of strain SINDOREI, various virulence factors are involved in the biofilm formation according to the previously reported including LPS.
(Huang et al., 2006; Pompilio et al., 2011; Zhuo et al., 2014; Madi et al., 2016), flagella formation (Di Bonaventura et al., 2007; Kang et al., 2015), type IV pili (Strom and Lory, 1993; Giltner et al., 2012), and purine biosynthesis (Kang et al., 2015). Details are listed in Table 2. These genes associated with biofilm formation exist in the genome of strain SINDOREI, suggesting strain SINDOREI harbor the basic characteristic of biofilm formation.

Comparative analysis of antibiotic resistance genes

Antimicrobial susceptibility test revealed that strain SINDOREI is resistant to six antibiotics including TMP/SMX, ciprofloxacin, ofloxacin, cefepime, ceftazidime, and aztreonam (Table 3), which are classified as sulfonamide antibiotic, fluoroquinolone antibiotic and β-lactam antibiotic. At the same time, strain SINDOREI is sensitive to meropenem, piperacillin tazobactam, sulbactam cefoperazone, amikacin, and tigecycline.

The comparison of resistance-related genes in strains SINDOREI, SUNEO, JCM 13310T, T25-65, T0-18 and ZAC14D2_NAIMI4 was performed (Table 4). The results showed that S. acidaminiphila possesses similar antibiotic resistance genes. A total of 24 key genes including encoded genes (14 genes) and efflux pumps (10 genes) contribute to the multidrug resistance in strain SINDOREI. The 24 genes are involved in resistance to TMP/SMX (Toleman et al., 2007; Hu et al., 2011), fluoroquinolone antibiotic and β-lactam antibiotic. At the same time, strain SINDOREI is sensitive to meropenem, piperacillin tazobactam, sulbactam cefoperazone, amikacin, and tigecycline.

Homologues of the sul genes in Stenotrophomonas acidaminiphila genomes

The sul genes encoding variants of the dihydropteroate synthase are responsible for resistance to TMP/SMX in many bacteria (Huang et al., 2018; Domínguez et al., 2019). There are two sul genes (sul1 and sul2 genes) in Strain SINDOREI. The amino acid sequences of homologues to the sul genes in Stenotrophomonas genomes were employed to construct the phylogenetic tree. Four distinct groups were generated (Figure 7), and Sul1 and Sul2 in SINDOREI belong to different groups. The distribution of sul1 and sul2 genes in strain SINDOREI, T0-18 and T25-65 may promote the resistance to TMP/SMX. The amino acid sequence of Sul1 in strain SINDOREI shows 100% identity with that of strains JCM13310T, ZAC14D2_NAIMI4_2 and T25-65. The identity sequences suggest that these elements are ancient and acquired long with the use of antibiotics. According to the genome sequence of strain SINDOREI, we noted that Is6 and Tn3 family transposase were located immediately upstream of sul2 and Is6 family transposase was located downstream of sul2, suggesting that sul2 gene was acquired from horizontal gene transfer.
Discussion

In this study, the genomes of six *S. acidaminiphila* strains, including SINDOREI, SUNE0, JCM13310T, T25-65, T0-18, and ZAC14D2_NAIMI4_2, were employed to perform the comparative analysis. Strain SINDOREI was isolated from the blood of a patient with sepsis. Meanwhile, *S. acidaminiphila* was the sole organism cultured from this patient. Antimicrobial susceptibility test reveals that strain SINDOREI is resistant to TMP/SMX, ciprofloxacin, ofloxacin, cefepime, ceftazidime, and aztreonam. The results suggested that *S. acidaminiphila* is an emerging opportunistic pathogen with environmental origin.

Based on the analysis of six genomes, strain SINDOREI shows highest ANI and dDDH values with T25-65 and shared lowest ANI and dDDH values with SUNE0. Two thousand five hundred six orthogroups were found in six genomes, though only 2,691 orthogroups was shared between strain SINDOREI and SUNE0. The clinical isolates, strain SINDOREI and SUNE0, shows relatively high strain-level diversity compared to other strains. Further work is still needed for getting more isolates to profile a comprehensive picture of the population of *S. acidaminiphila*.

We explored the virulence features in strain SINDOREI, and the nitrate reductase enzyme operon (*narGH*) was also found. However, *narth* gene was only found in strain JCM 13310T, T25-65, T0-18 and ZAC14D2_NAIMI4_2. As reported previously, the nitrate reductase enzyme is important in the pathogenesis of many bacteria and a *narG* knockout mutant can cause reduced virulence and reduce lung damage in severe combined immunodeficiency mice (Fritz et al., 2002). Biofilm formation is
FIGURE 4
Groups of orthologous and paralogous genes (i.e., orthogroups) identified in the six Stenotrophomonas acidaminiphila strains used in this study. The vertical bars show the number of orthogroups exclusive to the strains marked as lower dots in the matrix. Horizontal bars represent the total number of genes in each strain.

FIGURE 5
Comparison of the gene contents in Stenotrophomonas acidaminiphila. (A) Flower plot diagram showing the core genes and specific genes. (B) Distribution of COG functional annotations of SINDOREI specific genes.
also an important pathogenesis of bacteria such as *S. maltophilia*. Various genes associated with biofilm formation were found in strain SINDOREI which were considered as the basic characteristics of biofilm formation involved in pathogenesis of infections. Biofilm can be attached to various abiotic surfaces and tissues (Flores-Treviño et al., 2019). This specific structure provides up to 1,000 times more resistance to antimicrobial drugs (Mah, 2012; Olsen, 2015) and contributes to respiratory diseases (Costerton et al., 1999; Pompilio et al., 2010). New antimicrobial strategies (antibiofilm strategies) were used to treat *Stenotrophomonas* infections. Biofilm assay should be conducted in further study to explore biofilm’s correlations with virulence. Therefore, present ongoing studies about strain SINDOREI are limited and valuable to in-depth mining.

**Investigation of the presence of virulence genes is important to explain the genetic mechanisms of multidrug resistance.** In MDR strain SINDOREI, 24 genes involved in the resistance to a broad array of antimicrobial agents were analyzed. Strain SINDOREI has all the target genes mainly encoding antibiotic inactivating enzymes and multidrug efflux pumps, which is an important reason for the resistance to antibiotics. Meanwhile, the unique and redundant resistance genes occurred in strain SINDOREI. GES-1 (GES-type β-lactamase), *aadA3* (streptomycin 3′-adenylyltransferase), *qacE* (antibiotic efflux of disinfecting agents), and *cmlA5* (antibiotic efflux of phenicol antibiotic) associated with the resistance to β-lactam, aminoglycoside, disinfecting agents and phenicol are exclusive to strain SINDOREI. The phylogenetic analysis of homologues to *sul* genes indicated that there are two types of *sul* genes (*sul1* and *sul2*) in strain SINDOREI. The presence of redundant genes (*sul1* and *sul2*) in strain SINDOREI, should be responsible for the increased resistance to TMP/SMX. The presence of IS6 and Tn3 family transposase located upstream and downstream of the *sul2*, suggesting that acquisition of resistance genes is a relevant mechanism for strain SINDOREI antibiotic resistance.

**Conclusion**

In this study, we reported the world’s first case of fatal infection with *S. acidaminiphila* and obtained the high-quality genome of clinical isolated MDR strain SINDOREI. The comparative
### TABLE 2. Genes involved in biofilm formation in *Stenotrophomonas acidaminiphila* SINDOREI.

| Genes | SINDOREI gene locus | Activity | Function | References |
|-------|---------------------|----------|----------|------------|
| **Polysaccharides** | | | | |
| wbtL | JNIIIPNH_00556 | Lipopolysaccharides | Glucose-1-phosphate thymidyl transferase | Huang et al. (2006), Pompilio et al. (2011), Zhuo et al. (2014), and Madi et al. (2016) |
| rfbC | JNIIIPNH_00557 | Biosynthesis involved in | dTDP-4-dehydrorhamnose 3,5-epimerase | |
| spgM | JNIIIPNH_00560 | Biofilm formation | Phosphoglucomutase/phosphomannomutase | |
| **Flagella** | | | | |
| fliG | JNIIIPNH_01883 | Flagellar basal body rod protein | Twitching motility | Di Bonaventura et al. (2007) and Kang et al. (2015) |
| fliH | JNIIIPNH_01884 | Flagellar basal body L-ring protein precursor | Twitching motility | |
| fliI | JNIIIPNH_01923 | Flagellar biosynthesis protein | | |
| fliA | JNIIIPNH_01926 | Flagellar switch protein | | |
| **Fimbriae** | | | | |
| pilU | JNIIIPNH_00520 | Type IV pilus formation | Twitching motility | Strom and Lory (1993) and Giltner et al. (2012) |
| pilU | JNIIIPNH_00971 | associated with biofilm | | |
| pilZ | JNIIIPNH_00959 | Type 4 fimbrial biogenesis | | |
| pilT | JNIIIPNH_00970 | Twitching motility | | |
| pilH | JNIIIPNH_02554 | Twitching motility | | |
| pilH | JNIIIPNH_02624 | Twitching motility | | |
| pilG | JNIIIPNH_02625 | Two-component response regulator | | |
| pilB | JNIIIPNH_02689 | Type IV fimbrial biogenesis | | |
| tapC | JNIIIPNH_02705 | Type IV fimbrial assembly | | |
| tapD/pilD | JNIIIPNH_02706 | Prepilin peptidase | | |
| pilM | JNIIIPNH_02777 | Type IV plus inner membrane | | |
| tapU | JNIIIPNH_00971 | Twitching ATPase | | |
| **Other** | | | | |
| purD | JNIIIPNH_03072 | Purine biosynthesis involved in biofilm formation | Phosphoribosylamine-glycine ligase | Kang et al. (2015) |
| purC | JNIIIPNH_03118 | | Phosphoribosylaminomimidazolesuccinocarboxamide | |
| purI | JNIIIPNH_02993 | | Phosphoribosylformylglycinamidine synthase | |

Genomes analysis suggested a unique gene (narG) and key genes involved in biofilm formation in strain SINDOREI played an important role in pathogenesis of infections. Antimicrobial susceptibility test revealed that stain SINDOREI were resistant to TMP/SMX, ciprofloxacin, ofloxacin, cefepime, ceftazidime, and aztreonam. The presence of redundant Sul1 and Sul2 from two distinct groups and the exclusive determinants GES-1, *aad*A3, *qac*L and *cml*A5 exist in SINDOREI, which can explain the mechanisms of strain's multidrug resistance and afford potential therapeutic strategies for pathogen infections.

**Data availability statement**

The genome sequence data of strain SINDOREI has been deposited in National Genomics Data Center (NGDC, [https://ngdc.cnbc.ac.cn/](https://ngdc.cnbc.ac.cn/)) and accession numbers is PRJCA009493, as mentioned in Table 1 ([CNCB-NGDC Members and Partners., 2022](https://ngdc.cnbc.ac.cn/)).

**Ethics statement**

Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

**Author contributions**

YZ, PX, and WC made contributions to the acquisition of clinical data. QY, MZ, AZ, and WW contributed to laboratory
### TABLE 3 Antimicrobial susceptibility test of *Stenotrophomonas acidaminiphila* SINDOREI, SUNEO and JCM 13310<sup>β</sup>.

| Class                     | Antibiotics                             | SINDOREI | SUNEO | JCM 13310<sup>β</sup> |
|---------------------------|-----------------------------------------|----------|-------|------------------------|
| Sulfonamide antibiotic    | Trimethoprim/Sulfamethoxazole           | ≥64/304, R | 80 (4/76), R | ≥2/38, S |
| Fluoroquinolone antibiotic| Ciprofloxacin                           | ≥4, R    | ≤0.25, S | ≤1, S |
|                           | Ofloxacin                               | ≥8, R    | ≤1, S  | ≤1, S |
| β-lactam antibiotic       | Cefalotin                               | ≥32, R   | ≤1, S  | > 32, R |
|                           | Cefepime                                | ≥16      |       |           |
|                           | Ceftriaxone                             | ≥64, R   | ≤1, S  | ≤4, S |
|                           | Amoxicillin                             | > 16, R  |       |           |
|                           | Piperacillin                             | ≤16, S   |       |           |
|                           | Aztreonam                               | ≥64, R   |       |           |
|                           | Imipenem                                | ≥16, R   |       | > 8, R    |
|                           | Meropenem                               | ≤0.25, S |       |           |
| β-lactam combination agents| Amoxicillin clavulanic acid              | > 16, R  |       |           |
|                           | Ampicillin Sulbactam                    | ≤2, S    |       |           |
|                           | Piperacillin tazobactam                 | ≤16, S   |       |           |
|                           | Subactam cepolperazone                  | ≤8, S    |       |           |
| Aminoglycoside antibiotic | Amikacin                                | 16, S    | 16, S  | ≤8, S |
|                           | Gentamicin                              | 2, S     | ≤4, S  |           |
| Tetracycline derivative   | Tigecycline                             | ≤0.5, S  | ≤0.5, S |           |

(Continued)

### TABLE 4 The antibiotic resistance genes among the *Stenotrophomonas acidaminiphila* strains.

| Genes               | SINDOREI | SUNEO | ZAC14D2_NAIMI4_2 | T25-65 | T0-18 | JCM13310<sup>β</sup> |
|---------------------|----------|-------|------------------|--------|-------|------------------------|
| Trimethoprim/sulfamethoxazole resistance gene | sulk | JNIIIPNH_01405 | B1L07_06465 | AOT14_RS07185 | F0P98_RS10870 | F0P95_RS03720 | ABB33_13125 |
| sulk2               | JNIIIPNH_01524 | –     | –                | –      | –     | –                      |
| dfkA                | JNIIIPNH_02899 | –     | AOT14_RS14395 | F0P98_RS03500 | F0P95_RS17150 | –                      |
| Fluoroquinolone resistance gene | qnr | JNIIIPNH_03313 | B1L07_15000 | AOT14_RS17045 | F0P98_RS16470 | F0P95_RS02400 | –                      |
| β-lactam resistance gene | L1 | JNIIIPNH_02567 | B1L07_11340 | AOT14_RS12805 | F0P98_RS05130 | F0P95_RS15250 | ABB33_10340 |
| L2                   | JNIIIPNH_01026 | B1L07_04670 | AOT14_RS05330 | F0P98_RS12890 | F0P95_RS08575 | ABB33_03240 |
| GES-1                | JNIIIPNH_00474 | –     | –                | –      | –     | –                      |
| ampR                | JNIIIPNH_01025 | B1L07_04665 | AOT14_RS05345 | F0P98_RS12895 | F0P95_RS08570 | ABB33_02800 |
| ampN                | JNIIIPNH_00207 | B1L07_01060 | AOT14_RS02770 | F0P98_RS10115 | F0P95_RS04783 | ABB33_12865 |
| ampG                | JNIIIPNH_00208 | B1L07_01065 | AOT14_RS02775 | F0P98_RS11150 | F0P95_RS04790 | ABB33_12870 |
| ampD                | JNIIIPNH_02390 | B1L07_01310 | AOT14_RS11925 | F0P98_RS06020 | F0P95_RS14375 | ABB33_06015 |
| mrcA                | JNIIIPNH_00728 | B1L07_01315 | AOT14_RS13800 | F0P98_RS04120 | F0P95_RS16525 | ABB33_04710 |
| mrdA                | JNIIIPNH_00607 | B1L07_02815 | AOT14_RS03340 | F0P98_RS14735 | F0P95_RS06700 | –                      |
| Aminoglycoside resistance gene | adaA3 | JNIIIPNH_01525 | –     | – | – | – | – |
| Efflux pump         | smeDEF RND system | | | | |
| sme D               | JNIIIPNH_01709 | B1L07_07555 | AOT14_RS08230 | F0P98_RS14800 | F0P95_RS06630 | ABB33_10360 |
| sme E               | JNIIIPNH_01710 | B1L07_07560 | AOT14_RS08235 | F0P98_RS14795 | F0P95_RS06635 | ABB33_10365 |
| sme F               | JNIIIPNH_01712 | B1L07_07570 | AOT14_RS08245 | F0P98_RS05900 | F0P95_RS14485 | ABB33_10375 |

(Continued)
TABLE 4 (Continued)

| Genes          | SINDOREI     | SUNEO       | ZAC14D2_NAIMI4_2 | T25-65 | T0-18 | JCM13310 |
|----------------|--------------|-------------|------------------|--------|-------|----------|
| smeOP·ToIC RND system |             |             |                  |        |       |          |
| tolC           | JNIIIPNH_00706 | B1LO7_03300 | AOT14_RS03825    | F0P98_RS14240 | F0P95_RS07185 | ABB33_09990 |
| pcm            | JNIIIPNH_00707 | B1LO7_03305 | AOT14_RS03830    | F0P98_RS14235 | F0P95_RS07190 | ABB33_09985 |
| smeO           | JNIIIPNH_00709 | B1LO7_03315 | AOT14_RS03840    | F0P98_RS04655 | F0P95_RS16015 | ABB33_09950 |
| smeP           | JNIIIPNH_00710 | B1LO7_03320 | AOT14_RS03845    | F0P98_RS14220 | F0P95_RS07205 | ABB33_09945 |
| SMR efflux pump |             |             |                  |        |       |          |
| qacL           | JNIIIPNH_01527 | –           | –                | –      | –     | –        |
| MFS efflux pump |             |             |                  |        |       |          |
| cmlA5          | JNIIIPNH_01526 | –           | –                | –      | –     | –        |
| tetC           | JNIIIPNH_01532 | –           | –                | F0P98_RS10910 | F0P95_RS03755 | –        |

**FIGURE 7**

The phylogenetic tree based on the amino acid of sul genes in the members of genus Stenotrophomonas. The numbers present on the branches of the tree represent the bootstrap values (based on 1,000 replicates).
work. DW, HX, and YX conducted the clinical work. YZ and DL drafted the manuscript. ZJ revised the manuscript critically for important intellectual content and had given final approval of the version to be published. All authors contributed to the article and approved the published version.

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Conflict of interest

DL was employed by Hugobiotec Co. Ltd.

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Supplementary material

The Supplementary material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2022.989259/full#supplementary-material

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