Emergence of carbapenem-resistant and colistin-susceptible Enterobacter cloacae complex co-harboring \( \text{bla}_{\text{IMP}-1} \) and \( \text{mcr-9} \) in Japan

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**Abstract**

**Background:** The spread of Enterobacteriaceae producing both carbapenemases and Mcr, encoded by plasmid-mediated colistin resistance genes, has become a serious public health problem worldwide. This study describes three clinical isolates of Enterobacter cloacae complex co-harboring \( \text{bla}_{\text{IMP}-1} \) and \( \text{mcr-9} \) that were resistant to carbapenem but susceptible to colistin.

**Methods:** Thirty-two clinical isolates of \( E. \) cloacae complex non-susceptible to carbapenems were obtained from patients at 14 hospitals in Japan. Their minimum inhibitory concentrations (MICs) were determined by broth microdilution methods and E-tests. Their entire genomes were sequenced by MiSeq and MinION methods. Multilocus sequence types were determined and a phylogenetic tree constructed by single nucleotide polymorphism (SNP) alignment of whole genome sequencing data.

**Results:** All 32 isolates showed MICs of \( \geq 2 \) \( \mu \)g/ml for imipenem and/or meropenem. Whole-genome analysis revealed that all these isolates harbored \( \text{bla}_{\text{IMP}-1} \), with three also harboring \( \text{mcr-9} \). These three isolates showed low MICs of 0.125 \( \mu \)g/ml for colistin. In two of these isolates, \( \text{bla}_{\text{IMP}-1} \) and \( \text{mcr-9} \) were present on two separate plasmids, of sizes 62 kb and 280/290 kb, respectively. These two isolates did not possess a \( \text{qseBC} \) gene encoding a two-component system, which is thought to regulate the expression of \( \text{mcr-9} \). In the third isolate, however, both \( \text{bla}_{\text{IMP}-1} \) and \( \text{mcr-9} \) were present on the chromosome.

**Conclusion:** The \( \text{mcr-9} \) is silently distributed among carbapenem-resistant \( E. \) cloacae complex isolates, of which are emerging in hospitals in Japan. To our knowledge, this is the first report of isolates of \( E. \) cloacae complex harboring both \( \text{bla}_{\text{IMP}-1} \) and \( \text{mcr-9} \) in Japan.

**Keywords:** Enterobacter cloacae complex, \( E. \) Xiangfangensis, \( E. \) Asburiae, Mcr-9, \( \text{bla}_{\text{IMP}-1} \), Carbapenem resistance, Colistin resistance
Background

The emergence of carbapenemase-producing Enterobacteriaceae (CPE) has become a serious problem in medical settings worldwide [1]. The most frequently detected and globally widespread carbapenemase produced by CPE between the Asian countries are the class B metallo-β-lactamases (MBLs), which include IMP-type, NDM-type, and VIM-type MBLs [2].

Because of the emergence of multidrug-resistant Gram-negative pathogens and the lack of new antibiotics with efficient activities, colistin, a polymyxin-type antibiotic, has been the last resort used to treat CPE infections [3, 4]. Bacteria acquire colistin resistance through chromosomal mutation(s) or plasmid transfer [5]. Chromosome-mediated colistin resistance results from mutation(s) or deletion(s) of two component systems, such as phoPQ and pmrAB, altering the structure of lipopolysaccharides [6, 7]. For example, colistin resistance has been associated with modification of the lipid A moiety in lipopolysaccharide, such as by the addition of 4-amino-4-deoxy-L-arabinose (L-Ara-4 N) and phosphoethanolamine (PEtN) to the anionic phosphate groups of lipid A. These additions reduce the anionic charges on lipid A and its affinity to the cationic colistin, inhibiting membrane destruction resulting from the binding of colistin to lipid A, followed by cell death [8].

To date, various types of plasmid-mediated mobilized colistin-resistance genes, mcr, have been identified, including mcr-1 to mcr-9, with several, including mcr-1, −2, −3, −4, and −6, shown to have PetN transferase activity [6]. The mcr-1 gene was initially detected in isolates of Escherichia coli and Klebsiella pneumoniae obtained from humans and animals in 2015 in China [9], and mcr-9 was initially identified in a clinical isolate of the colistin-susceptible bacterium, Salmonella enterica serotype typhimurium. The amino acid and nucleotide sequences of mcr-9 are closest to those of mcr-3 with similarities of 64.5 and 99.5%, respectively [10]. In this study, mcr-9 was detected in 335 genomes in multiple genera of Enterobacteriaceae. The analysis of mcr-9 promoter region in these genomes showed conserved regions which is likely a recognition sequence for transcription regulator, suggesting that other factors might be involved in full-expression of mcr-9. Of the 335 genomes, 65 had at least one plasmid replicon indicating that mcr-9 can be found extrachromosomally in different species of Enterobacteriaceae [10].

Isolates of E. cloacae complex resistant to both carbapenem and colistin have been reported in several countries, including China [11, 12], France [13], India [14], the USA [15, 16] and Vietnam [17]. One of these, an isolate of E. cloacae complex (Enterobacter hormaechei) co-harboring blaVIM-4 and mcr-9, was first reported in the United States in 2019 [16]. In addition, a colistin-resistant E. hormaechei isolate producing both MCR-9 and NDM-1 was isolated from a patient in China with bloodstream infection in 2019 [11]. This emergence of colistin resistance, particularly in CPE, may result in significant clinical and public health concerns [18, 19].

The study describes three clinical isolates of E. cloacae complex that were resistant to carbapenem but susceptible to colistin. To our knowledge, this is the first report of isolates of E. cloacae complex harboring both blaIMP-1 and mcr-9 in Japan.

Methods

Bacterial strains

Thirty-two clinical isolates of E. cloacae complex, each with minimum inhibitory concentrations (MICs) of ≥2 μg/ml for meropenem and/or imipenem, had been obtained from individual patients at 14 hospitals in eight prefectures throughout Japan from July to October 2018 by BML Biomedical Laboratories R&D Center (Kawagoe, Saitama, Japan).

Drug susceptibility testing

The MICs of antibiotics were determined using a broth microdilution method according to the guidelines of the Clinical and Laboratory Standards Institute (CLSI) [20]. The MICs of colistin were also determined by a broth microdilution using cation-adjusted Muller Hinton broth and 96-well microtiter plates (Kohjin Bio, Co., Ltd. Saitama, Japan) according to the guidelines of the European Union Committee for Antimicrobial Susceptibility Testing (EUCAST) [21].

Whole genome sequencing

DNA was extracted from each E. cloacae complex isolate using DNeasy Blood and Tissue kits (Qiagen, Tokyo, Japan). A Nextera XT DNA library was prepared from each extracted DNA sample. Each DNA library of was sequenced on the MiSeq system (Illumina) to obtain short reads with 300-bp paired-end reads. MiSeqRun was performed using Nextera XT Index Kit v2 and MiSeq Reagent Kit v3. DNA Libraries for MiniON (Oxford Nanopore Technologies, Oxford, UK) were prepared from three isolates (A2483, A2504 and A2563) using Ligation Sequencing Kits 1D (SQK-LSK109) to yield long contigs. The long read generated by MiniON were assembled using Canu v1.7.1 and polished with the short reads generated by MiSeq using Pilon v1.22. The nucleotide sequences of plasmids and chromosomes carrying blaIMP-1 and mcr-9 were compared with similar sequences using BLAST and visualized by In silico MolecularCloning. Ver.7 genomic edition (https://www.insilicobiology.co.jp/).

Bacterial species were identified by analyses of average nucleotide identity (ANI) [22] and digital DNA-DNA
hybridization (dDDH) [23] of whole genome sequences. The seven type strains used as reference species included Enterobacter asburiae (ATCC35953T), E. cloacae (ATCC13047T), E. hormaechei (ATCC49162T), Enterobacter kobei (DSM13645T), Enterobacter ludwigi (EN-119T), Enterobacter nimpressurialis (DSM18955T) and Enterobacter xiangfangensis (LMG27195T). In silico multilocus sequence typing (MLST) was assigned by PUBMLST database (https://pubmlst.org/databases/). Acquired antibiotic resistance genes were identified using the ResFinder 3.2 tool (https://cge.cbs.dtu.dk/services/ResFinder/) from the Center for Genomic Epidemiology (CGE).

Phylogenetic analysis based on SNPs

Single nucleotide polymorphisms (SNPs) in the 32 isolates were identified by aligning whole-genome sequencing data of these isolates with the genomic sequences of the E. xiangfangensis reference isolate LMG27195 (GenBank accession no. CP017183.3), using the CSI Phylogeny 1.4 tool (https://cge.cbs.dtu.dk/services/CSI-Phylogeny/) from CGE. A phylogenetic tree was constructed using Fig Tree (version 1.4.4) and a maximum likelihood phylogenetic tree (http://tree.bio.ed.ac.uk/software/figtree/).

Results

Phenotypic and genotypic properties of carbapenem-non-susceptible isolates

Drug susceptibility of carbapenem-non-susceptible isolates

The MICs of the 32 clinical isolates of E. cloacae complex are shown in Table 1. All were susceptible to amikacin and colistin, but resistant to cefazidime. Of these 32 isolates, 25 were resistant to aztreonam, 15 were resistant to ciprofloxacin, and 12 were resistant to tigecycline. Of the all 32 isolates, 28 isolates were resistant to imipenem and/or meropenem with MICs ≥4 μg/ml, whereas the remaining 4 were intermediate to imipenem and/or meropenem with MICs ≥2 μg/ml (Table S1). There are no isolates susceptible to both imipenem and meropenem (Table S1).

Whole genome sequences of carbapenem-non-susceptible isolates

Whole genome sequencing of the 32 isolates of E. cloacae complex showed that, based on ANI and dDDH analyses, 31 were E. xiangfangensis and one was E. asburiae. MLST analysis revealed that 13 isolates (40.6%) belonged to sequence type (ST) 78; 10 (31.2%) to ST133; two each (6.3%) to ST175 and ST1196; and one each (3.1%) to ST62, ST93, ST418, and ST484. The ST for one isolate could not be determined because its housekeeping genes did not match those of current STs. A phylogenetic tree of these 32 isolates revealed four major clades, with clades I, II, III and IV consisting of 14, 2, 10 and 6 isolates, respectively (Fig. 1). Clade I consisted of isolates belonging to ST78 and the non-typeable isolate, clade II of isolates belonging to ST418 and ST484, clade III of isolates belonging to ST133 and clade IV of isolates belonging to ST1196, ST175, ST93 and ST62. These isolates harbored various genes associated with drug resistance (additional file: Table S1). All 32 isolates harbored blaIMP-1, with three also harboring mcr-9 (Table S1).

Phenotypic and genotypic properties of isolates harboring both blaIMP-1 and mcr-9

Bacterial identification and drug susceptibility

Of three isolates co-harboring blaIMP-1 and mcr-9, two, A2483 and A2504, were E. xiangfangensis and one, A2563, was E. asburiae (Table S1). The A2483 and A2504 strains obtained in a hospital belonged to ST1199, whereas the A2563 strain obtained in another hospital belonged to ST484. The two hospitals located in the same prefecture in Japan. The drug susceptibility profiles of the two E. xiangfangensis isolates were identical to each other, with both A2483 and A2504 being resistant to aztreonam, cefazidime, imipenem, meropenem and tigecycline, and susceptible to amikacin, ciprofloxacin and colistin (Table 2). The E. asburiae isolate was resistant to cefazidime and imipenem, had intermediate resistance to meropenem, but was susceptible to the other drugs tested including colistin (Table 2).

Whole genome sequences of isolates harboring both blaIMP-1 and mcr-9

As shown in Table 3, E. xiangfangensis A2483 contained a chromosome of 5,024,985 bp with a GC content of 55.24% and two plasmids of 61,594 bp and 288,696 bp, respectively. E. xiangfangensis A2504 contained a chromosome of 4,934,510 bp with a GC content of

| Table 1 | MIC values of 32 clinical E. cloacae complex isolates |
|----------|-----------------------------------------------------|
| Antimicrobial Agents | Breakpoint for resistance (μg/ml) | Range | No. of resistant isolates (%) | MIC data (μg/ml) | MIC<sub>50</sub> | MIC<sub>90</sub> |
| Amikacin | ≥24 | 0 | 0.5 to 4 | 1 | 2 |
| Aztreonam | ≥16 | 25 (78.1%) | <0.25 to 256 | 32 | 256 |
| Cefazidime | ≥16 | 32 (100%) | 256 | 512 |
| Ciprofloxacin | ≥4 | 15 (46.9%) | <0.25 to 64 | 4 | 16 |
| Colistin<sup>a</sup> | >2 | 0 | 0.03 to 2 | 0.25 | 0.5 |
| Imipenem | ≥24 | 18 (56.25%) | <0.25 to 16 | 4 | 8 |
| Meropenem | ≥24 | 18 (56.25%) | 0.5 to 16 | 4 | 16 |
| Tigecycline<sup>a</sup> | >0.5 | 12 (37.5%) | <0.25 to 4 | 0.5 | 1 |

<sup>a</sup>Breakpoints for antimicrobial resistance were determined according to CLSI guidelines

<sup>b</sup>Breakpoint for Colistin and Tigecycline was determined according to EUCAST guidelines
54.70% and two plasmids of 61,594 bp and 276,927 bp, respectively. The whole genome sequences of A2483 were very close to those of A2504 with similarities of 100% (98% query coverage) on the chromosome, 100% for the 62-kbp plasmid and 100% (96% query coverage) for the 289-kbp plasmid.

E. asburiae A2563 contained a chromosome of 4,934,510 bp with a GC content of 55.80% and one plasmid 115,246 bp in size (Table 3). In addition to blaIMP-1 and mcr-9, these isolates harbored several other genes associated with drug resistance, including aac (6′)-IIc, blaACT-6, blaACT-7, fosA and sul1 (Table 3). blaACT genes are the intrinsic AmpC encoding genes of Enterobacter cloacae complex species.

Location of mcr-9 and its genetic environments

The mcr-9 gene was present on the 289-kbp IncHI2 plasmid of A2483 and the 277-kbp IncHI2 plasmid of A2504, but was present on the chromosome of A2563 (Table 3). The two plasmids harboring mcr-9, pA2483mcr-9 on A2483 and pA2504mcr-9 on A2504, had the same GC content of 46.30%, and contained open reading frames (ORFs) of 360 and 358, respectively (Fig. 2. (a)). The nucleotide sequences of these plasmids were identical to each other, except for a genetic region with 11,770 bp, from nucleotide (nt) 146,310 to nt 158,080, in the 277-kbp plasmid. The mcr-9 gene on the chromosome of A2563 was detected at nt ~129 Mb.

The genetic environments of mcr-9 in the A2483 and A2504 plasmids were identical to each other, with mcr-9 located in a ~30 kb region surrounded by two insertion sequences encoding an IS5-like element (IS903 family transposase; Fig. 3). The region upstream of mcr-9 included renR (encoding a Ni/Co-binding transcriptional repressor), pcoS (encoding a two-component sensor histidine kinase) and pcoE (encoding a copper-binding protein). The region downstream of mcr-9 included wbuC (encoding a cupin fold metalloprotein) but no genes encoding the two-component system qseC-qseB, which has been associated with the expression of mcr-9 [24]. Insertion sequences were not detected in the region downstream of mcr-9 on the A2563 chromosome. The region upstream of mcr-9 was renR-pcoS-DpcoE, whereas the region downstream of mcr-9 was wbuC-qseC-qseB (Fig. 3). The A2483 and A2504 plasmids showed 83% query coverage and 99.97% identity to the IncHI2 plasmid, pME-1a (GenBank accession no. NZ_CP041734.1), in E. hormaecheii, a strain isolated in 2019 from a pediatric inpatient in the USA (Fig. 3) [16]. Three IncHI2 plasmids were

| Isolates/Antimicrobial Agents | MIC (μg/ml) |
|------------------------------|-------------|
|                              | AMK | AZT | CAZ | CIP | CST | IPM | MPM | TIG |
| E. xiangfangensis A2483      | 1   | 128 | >512| 1   | 0.125| 4   | 8   | 1   |
| E. xiangfangensis A2504      | 1   | 128 | >512| 1   | 0.125| 4   | 8   | 1   |
| E. asburiae A2563            | 0.5 | 0.25| 128 | 0.5 | 0.125| 8   | 2   | 0.5 |

*Breakpoints for antimicrobial resistance were determined according to CLSI guidelines

*Breakpoints for Colistin and Tigecycline was determined according to EUCAST guidelines
identified with similar sequences, pCTXM9_020038 from E. hormaechei isolated in China in 2018 (83% query and 99.97% identity; GenBank accession no. CP013724), pRH-R27 from Salmonella enterica Infantis in Germany in 2015 (82% query coverage and 99.99% identity; GenBank accession no. LN555650), and pMCR-SCNJ07 from E. hormaechei isolated in China in 2019 (80% query and 99.99% identity; GenBank accession no. MK933279) (Fig. 3).

**Location of bla IMP-1 and its genetic environment**

The bla IMP-1 gene was present on the 62-kbp plasmids of A2483 and A2504 and on the chromosome of A2563 (Table 3). The two plasmids harboring bla IMP-1, pA2483imp-1 from A2483 and pA2504imp-1 from A2504, had the same GC content of 47.40% and identical nucleotide sequences (Fig. 2 (b)). In these plasmids, bla IMP-1 was located in a class I integron containing intI bla IMP-1-aac (6′)-Iic-qacEΔ1-sul1. The bla IMP-1 gene on the chromosome of A2563 was present in the same class I integron (Fig. 2 (c)). The same class I integron containing bla IMP-1 and aac (6′)-Iic was detected in the bacteria E. asburiae NUH15_ECL035_1 (GenBank accession no. AP013886.1), Enterobacter cloacae NUH15_ECL020 (GenBank accession no. AP013866.1) and Enterobacter asburiae NUH12_ECL030 (GenBank accession no. AP019383.1), all of which were isolated in Japan in 2019. The two pA2483imp-1 and pA2504imp-1 showed 84% query coverage and 97.75% identity with the plasmid pJJ1886.4 (GenBank accession no. CP006788.1), which was detected in the USA and did not contain a class I integron or any other resistance genes [25].

**Discussion**

The mcr-9 gene may be silently spreading in Enterobacteriaceae throughout the world. The prevalence of mcr-9 is unclear because this gene is not actually related to colistin resistance, as it may be silent or inducible in clinical isolates of Enterobacteriaceae. For example, an isolate of E. hormaechei harboring mcr-9 did not express its gene product [16]. This isolate was susceptible to colistin, likely because the two-component system genes qseCB were lacking from the region downstream of mcr-9. In contrast, another isolate of E. hormaechei that harbored and expressed mcr-9 was found to be resistant to colistin and to have the two-component system genes in the region downstream of mcr-9 [11]. The expression of mcr-9 is mediated by the two-component system QseCB and can be induced by subinhibitory concentrations of colistin [24]. At least 11 mcr-9-positive IncHI2 plasmids have been detected by Blast, with six having and five lacking the two-component system genes [11].

The two-component QseCB system, consisting of a sensor (qseC) and a response regulator (qseB), plays an essential role in the expression of mcr-9 [24]. Our finding, that the isolate A2563 harbored mcr-9 along with the two-component system genes qseCB but was susceptible to colistin suggests that other, as yet undetermined, genes or molecules may regulate mcr-9 expression. The pA2480mcr-9 and pA2504mcr-9 had similar structures to those of pME-1a and pCTXM9_020038, as they lacked qseCB. This two-component system was transcribed as an operon, with the QseB promoter binding to low- and high-affinity binding sites located – 500 to – 10 bp at upstream of qseB [26]. The nucleotide sequence of this region in A2563 was 100% identical to that of the QseB promoter (– 500 to +1 bp) in pMCR-SCNJ07, which confers resistance to colistin [11], suggesting that the QseB promoter in A2563 may be repressed by an as yet undetermined mechanism [26]. Four plasmids, pME-1a, pCTXM9_020038, pRH-R27 and pMCR-SCNJ07, had the conserved gene structure, renR-pcoS-pcoE-IS-5, upstream of mcr-9. Whereas, the chromosome of A2563 had the same conserved gene structure, but with a 53-bp deletion in pcoE (ΔpcoE), suggesting that the deleted region may be associated with mcr-9 expression. Further studies are necessary to determine the mechanism for regulation of mcr-9 expression in Enterobacteriaceae.

To our knowledge, it is the first report describing a bacterial isolate harboring mcr-9 on its chromosome,
indicating that mcr-9 may have been inserted into the chromosome by mobile elements. Several Enterobacteriaceae isolates from animals and humans have reported the chromosomal location of mcr-1 and mcr-2 [27–33]. The mcr-1 was detected on the chromosomes of two colistin-resistant E. coli strains isolated from swine in 2012 in China [28], and on the chromosome of an E. coli ST410 strain harboring blacTX-M-15 isolated from a sample of turkey meat in 2013 in Germany [27]. The chromosomal integration of mcr-1 was also detected in a clinical strain of E. coli ST156 harboring blaNDM-5 isolated from a bile sample in 2015 in China [29], in E. coli isolated from food production animals in 2011–2016 in Poland [32], and in E. coli isolated from veal calves in 2016 in the Netherlands [30]. Chromosomes carrying mcr-1 were detected in Enterobacteriaceae from environmental water sources in 2017 in China [33]. Moreover, the mcr-2 gene (mcr-6.1) was detected on the chromosome of a strain of Moraxella isolated from a pig in 2014–2015 in Great Britain [31]. These studies support the mobility characterization of mcr genes across different genetic elements and insertion of the plasmid-variant of mcr into chromosome could lead to higher prevalence of colistin resistance among Enterobacteriaceae species.

The direct origin of the mcr-9 on the chromosome of A2563 is unclear. However, the genetic environments of the mcr-9 and qseCB genes in A2563 are similar to those of pMCR-SCNJ07 from E. hormaechei in China in 2019 (GenBank accession no. MK933279), pRH-R27 from Salmonella enterica Infantis in Germany in 2015 (GenBank accession no. LN555650), pT5282-mphA from E. cloacae in China in 2012 (GenBank accession no. KY270852), pN1863-HI2 from E. cloacae in China in 2017 (GenBank accession no. MF344583), pSE15-SA01028 from S. enterica subsp. enterica in Germany in 2018 (GenBank accession no. NZ_CP026661) and p707804-NDM from Leclercia adecarboxylata in China in 2018 (GenBank accession no. MH909331). These 7 strains carrying plasmids with mcr-9 in China and Germany did not harbor blaIMps, but blaNDCs or blAVIMs [11].

The plasmids pA2483mcr-9 and pA2504mcr-9 had the same backbone as the plasmid plJ1886_4 (GenBank accession no CP006788.1), which had been isolated in the USA. The 55,956 bp plasmid plJ1886_4, which was smaller in length than the 61,594 bp plasmids pA2483mcr-9 and pA2504mcr-9, lacked a class I integron carrying blalMP1 (intI1-blalMP1-aac (6')-Ile-gacE Δ1-suI). The E. cloacae EN3600 plasmid (GenBank accession no. CP035638.1) carrying blalMP1 also had the same backbone as pJ1886_4, with 83% coverage and 96.8% identity. These findings indicate that plJ1886_4 has spread globally and captured drug-resistance genes and that this plasmid functions as a carrier of acquired drug-resistance genes.
**Conclusion**

In conclusion, this study describes the characterization of the complete genomes of three clinically obtained isolates of carbapenem-resistant and colistin-susceptible *E. cloacae* complex harboring both *bla*<sup>IMP-1</sup> and *mcr-9* from different hospitals in Japan. *Enterobacteriaceae* harboring both *bla*<sup>IMP-1</sup> and *mcr-9* may become a healthcare problem, suggesting the need for steps to prevent their further dissemination.

**Supplementary information**

Supplementary information accompanies this paper at https://doi.org/10.1186/s12879-020-05021-7.

**Acknowledgements**

Not applicable.

**Authors’ contributions**

PK and TK created the research data and wrote the draft of the manuscript. SI and KK determined phenotypic properties of isolates. SO and TT determined genetic properties of the isolates. SW, MS and MO collected the clinical isolates. All authors read, made significant edits to the first version, and approved the final manuscript.

**Funding**

This study was supported by grants from the Japan Society for the Promotion of Science (grant numbers 18K07120 (T.T.), 18K07121 (T.K.) and 19K02023 (T.K.), the Research Program on Emerging and Re-emerging Infectious Diseases from the Japan Agency for Medical Research and Development [grant number 19k0108061h0302 (T.K.)], and the Asahi Group Holdings, Ltd. [grant number AM227CH501 (S.W.)]. We declare no conflict of interest. P.K. received an international student scholarship from Juntendo University (grant number 6119002).

**Availability of data and materials**

All the data necessary to reproduce the results can be found in the manuscript’s tables; the calculations can be reproduced using the manuscript’s appendixes. The genome sequence of strain A2563 has been deposited into GenBank under the accession number of AP022628. The complete nucleotide sequence of pA2483mcr-9, p2483imp-1, pA2504mcr-9, and p2504imp-1 have been deposited into GenBank under the accession numbers of LCS32224, LCS32225, LCS32226 and LCS32227, respectively.

**Ethics approval and consent to participate**

The study protocol was approved by the ethics committee of Juntendo University (number 809), and by the Biosafety Committee, Juntendo University (approval number BSL2/29–1). Individual informed consent was waived by the ethics committee listed above because this study used currently existing sample collected during the course of routine medical care and did not pose any additional risks to the patients. Informed consent about study participation was officially announced by mail and poster. All patient data were anonymized prior to the analysis.

**Consent for publication**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

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Received: 15 January 2020 Accepted: 8 April 2020 Published online: 16 April 2020

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