

# Novel *Pseudomonas aeruginosa* Strains Co-Harboring $bla_{\text{NDM-1}}$ Metallo $\beta$ -Lactamase and *mcr-1* Isolated from Immunocompromised Paediatric Patients

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**Background:** The rising resistance to carbapenems in Gram-negative bacteria worldwide poses a major clinical and public health risk. This study aimed to characterise carbapenem- and colistin-resistance genes,  $bla_{\text{NDM-1}}$  and *mcr-1* located on IncX4 plasmid in MDR *Pseudomonas aeruginosa*, isolated from paediatric patients undergoing chemotherapy as a result of leukaemia.

**Methods:** In this study, six carbapenem-resistant strains of *P. aeruginosa* were isolated from two paediatric patients under chemotherapy treatment (1.8 years old female and 2.1 years male) from the Shenzhen Hospital, China, in the year 2019. Isolates were screened for conventional antibiotics such as tobramycin, cefepime, imipenem, and ciprofloxacin in addition colistin by using the broth dilution method. Furthermore, resistance determinants: *mcr-1*,  $bla_{\text{NDM-1}}$ ,  $bla_{\text{KPC-1}}$ , and  $bla_{\text{GES}}$  were screened using PCR and sequencing followed by multi-locus sequence typing. The horizontal gene transfer and location of *mcr-1* and  $bla_{\text{NDM-1}}$  were determined by a liquid mating assay. In addition, Incompatibility type (Inc), PCR-based replicon type, and subgroup (MOB) of plasmid were studied.

**Results:** The screening for conventional antibiotics isolates showed 100% resistance to all the tested antibiotics except tobramycin. All isolates harboured carbapenemase encoding  $bla_{\text{NDM-1}}$ , of which three also had *mcr-1* located on a single IncX4 transferable plasmid. MLST typing revealed that four strains had a novel (new) STs type, while two belonged to ST1966.

**Conclusion:** This study identified for the first time colistin- and carbapenem-resistant MDR *P. aeruginosa* in paediatric patients with leukaemia in Shenzhen, China. It highlights the need for continuous surveillance in high-risk clones of MDR *P. aeruginosa*. Prudent use of antibiotics based on local antimicrobial susceptibility and clinical characteristics can help in reducing mortality in immunocompromised patients.

**Keywords:** *mcr-1*,  $bla_{\text{NDM-1}}$ , *Pseudomonas aeruginosa*, leukaemia patients

## Introduction

*Pseudomonas aeruginosa* is a Gram-negative, rod-shaped, aerobic bacterium and one of the most common causes of infectious disease in paediatric patients, leading to high mortality.<sup>1</sup> It is a very common opportunistic pathogen that can normally be present on the skin of healthy individuals but may cause serious infection in the case of immunocompromised patients with haematological malignancies.<sup>2,3</sup> Carbapenem-resistant *P. aeruginosa* has been classified as a critically important pathogen by the WHO in the year 2017 because of its ability and resilience to develop antimicrobial resistance (AMR) to the most commonly used antimicrobial drugs required for life-threatening infections. Hence, new

antibiotics are urgently required in order to tackle the silent pandemic of AMR.<sup>4</sup> The acquisition of the resistance genes and genetic mutations enables the bacterium to resist antibiotics and survive in an antibiotic enriched environment where the selection pressure for the development of resistant strains is quite high.<sup>5</sup> The resistant *P. aeruginosa* exposure to various antibiotics and the development of cross-resistance between antimicrobials may result in multidrug-resistant (MDR) strains which are not only difficult to treat but may also expand clonally and become endemic to a specific region such as a specific hospital. Such MDR *P. aeruginosa* strains may be transmitted from patient to patient and sometimes lead to outbreaks among patients attending/sharing the same ward.<sup>6</sup> Recent data show that MDR *P. aeruginosa* has a high prevalence rate that varies from 15% to 30% globally in hospital settings.<sup>7</sup> In African countries alone, 13% of severe healthcare-associated infections are caused by MDR *P. aeruginosa*.<sup>8</sup> The ST1966 high-risk clone of *P. aeruginosa* has a high capacity to acquire antibiotic resistance genes and is prevalent in many parts of China, especially in hospital settings (unpublished data). However, no study has reported the occurrence of ST1966 co-harboring New Delhi Metallo-beta-lactamase-1 (NDM-1) and *Klebsiella pneumoniae* carbapenemase (KPC) in *P. aeruginosa*. A previous study has reported a co-existence of *E. coli* with NDM-1 and MCR-1.<sup>9</sup> In addition, there has been no report on the co-existence of *bla*<sub>NDM-1</sub> and plasmid-born colistin resistance *mcr-1* in *P. aeruginosa* in paediatric patients. The occurrence and distribution of colistin resistance in *P. aeruginosa* clones is of huge concern, because of the fatality and limited treatment options and colistin being used as the last choice of drug to treat MDR pathogens.<sup>10</sup> Due to the increased risk of infections caused by NDM-1 and KPC-producing pathogens and the lack of proven local information, the WHO recently pointed out the need to invest in resources to improve the surveillance of antimicrobial-resistant (AMR) clones of bacteria,<sup>11,12</sup> using whole-genome sequencing technology.<sup>13</sup> Hence, it is important to investigate and monitor the emergence of resistant clones and make prudent use of antibiotics in order to treat any life-threatening infections. The present study aimed to evaluate carbapenemase-producing genes and their location in *P. aeruginosa* strains isolated from paediatric patients undergoing chemotherapy for leukaemia in XX hospital in China.

## Methods

The study was conducted according to the guidelines of the Declaration of Helsinki and approved by the Institutional Ethics Committee, Shenzhen Children's Hospital, Reference number: 2018 (013) dated 2018/09/03, which complies with international ethical standards. All experiments were conducted as per the hospital biosafety regulations act.

## Isolation and Identification

In this report, six carbapenem-resistant strains of *P. aeruginosa* were isolated from two paediatric patients (1.8 years old female, and 2.1 years male). Both patients had a history of acute leukaemia and were admitted to our hospital for chemotherapy, from July to August 2019. The demographic and clinical data were recorded on a standardized case form. During treatment, patients developed symptoms such as fever, cough, diarrhoea, and tissue infection, which followed microbiological investigation (Table 1). To identify the pathogens, full blood, urine, peripheral blood, CSF, hydrothorax and stool were cultured on the blood agar and Macconkey's agar. The initially isolated identification was done by using the VITEK®2 compact system (BioMerieux, France), followed by the API20 system and 16sRNA amplification. The sequencing of each purified PCR product was performed by Sangon Biotech, Shanghai. PCR assay was performed to amplify the full length of 16SRNA by using a set of universal primers: F-5'-AGAGTRTGATCMTYGCTWAC-3'; and R-5'-CGYTAMCTTWTACGRCT-3'. PCR reaction volume was 20 µL and it contained 2µL (30ng) of genomic DNA extracted by QIAamp®DNA Kit from QIAGEN (Germany), 10 µL of 2X Master Mix, 0.4 µL (20 pmol) of each forward and reverse primer, and 8.2 µL of nuclease-free water. Thermocycler conditions were set at 94°C for 5 min, followed by 35 cycles of denaturation at 94°C for 30 sec, annealing at 55°C for 30 sec and extension at 72°C for 1min, followed by a final extension at 72°C for 5 min. The PCR water was used as the control while *P. aeruginosa* ATCC 27853 DNA was used as the positive control. The PCR product was run on 1.8% agarose along with a DNA ladder (100Kb). The sequences were analyzed using NCBI BLAST N; <https://blast.ncbi.nlm.nih.gov/Blast>. This isolate was collected as a routine hospital investigation procedure. Only verbal consent was obtained because no personal information was used for research purposes, therefore written consent was not required.

**Table 1** Clinical and Molecular Characterization of *Pseudomonas aeruginosa*

Code	Source	Isolation Ate	ST	PBRT (Conjugants)	MICs Colistin	MICs Imipinime	Antibiotic Resistance Determinant	Therapy	Days Hospitalized	Outcome	Symptoms	Risk Factors	Clonal Relation
SP-19-1051	Blood	30-06-2019	1966	FIS	0.5	32	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>KPC-1</sub></i> , <i>bla<sub>IMP</sub></i>	Piperacillin +cefepime +Amikacin	45	Discharge	Fever	Implanted Port, Ventilator	*
Sp-19-1053	Blood	03-07-2019	1966	FIS	0.5	16	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>KPC-1</sub></i> , <i>bla<sub>IMP</sub></i>	Piperacillin+ Tazobactam+ Cefepime	45	Discharge	Fever, cough, diarrhoea, tissue infection	Implanted Port, Ventilator	*
SP-19-2014	Blood	09-07-2019	New	IncX4	4	32	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>CTX-M-15</sub></i> , <i>bla<sub>KPC-1</sub></i> , <i>bla<sub>GES</sub></i> , <i>mcr-1</i>	Piperacillin+ tazobactam	30	Discharge	Fever, cough, diarrhoea, peritoneal infection	Urethral Catheter, Transverse Colostomy, Retroperitoneal Drain tube	***
SP-19-2015	Blood	10-07-2019	New	FIB, IncX4	4	32	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>KPC-1</sub></i> , <i>mcr-1</i>	Amikacin+ Piperacillin+ Tazobactam sodium	30	Discharge	Fever, cough, diarrhoea, tissue infection	Urethral Catheter, Transverse Colostomy, Retroperitoneal Drain tube	***
SP-19-2016	Blood	12-07-2019	New	HII, IncX4	4	32	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>CMY</sub></i> , <i>mcr-1</i>	Amikacin+ Piperacillin+ Tazobactam sodium	30	Discharge	Fever, cough, diarrhoea, tissue infection	Urethral Catheter, Transverse Colostomy, Retroperitoneal Drain tube	***
SP-19-2019	Blood	14-07-2019	New	Not Detected	1	16	<i>bla<sub>NDM-1</sub></i> , <i>bla<sub>CMY</sub></i>	Amikacin+ Piperacillin+ Tazobactam sodium	30	Discharge	Fever, cough, diarrhoea tissue infection	Urethral Catheter, Transverse Colostomy, Retroperitoneal Drain tube	***

Notes: \*Clone-1; \*\*\*Clone-2.

Abbreviations: ST, sequence type; MICs, minimum inhibitory concentration; PBRT, PCR based replicon typing.

## Antimicrobial Susceptibility

The antimicrobial susceptibility test (AST) was also performed using the VITEK®2 compact system (BioMerieux, France) with ASTGN09 card (software version 9.01) followed by the agar dilution method. Tested antibiotics were Amoxicillin, Tobramycin, Cefepime, Meropenem, Amikacin, Levofloxacin, Ciprofloxacin, Cefoperazone, Ticarcillin, and Colistin. The *P. aeruginosa* ATCC 27853 was used as a quality control strain for the AST. The carbapenemase production among carbapenem-resistant *P. aeruginosa* was confirmed using the modified Hodge test, while Metallo  $\beta$ -lactamase was determined by the MBL E-test.<sup>14</sup> The protocol for both tests was well developed in our infection and drug resistance lab and *E. coli* ATCC 25992 was used as the control strain while characterized strain of *E. coli* from our lab was used as the positive control. Antimicrobial susceptibility excluding colistin results was interpreted according to CLSI guidelines, while colistin resistance was defined according to European Committee on Antimicrobial Susceptibility Testing (EUCAST) clinical breakpoints.<sup>15,16</sup>

## Molecular Characterization

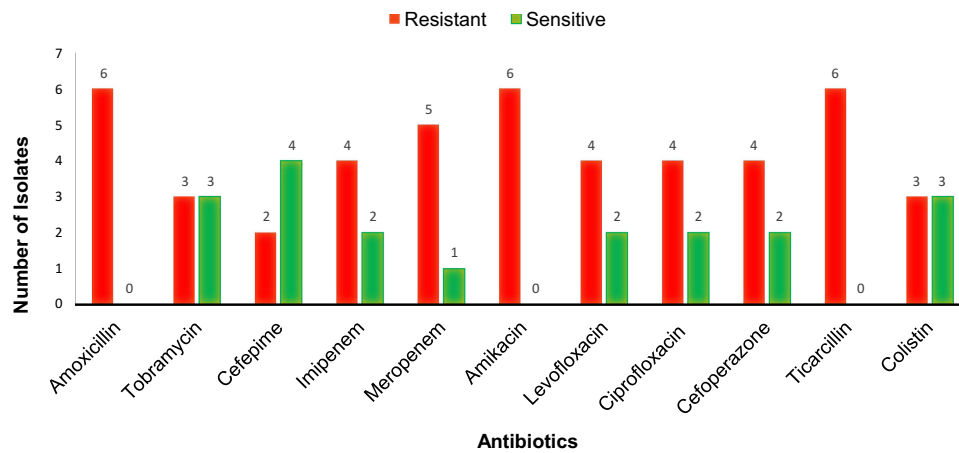
The PCR assays were performed for the detection of  $\beta$ -lactamase-encoding genes including *bla*<sub>CTX-M</sub>, *bla*<sub>TEM</sub>, *bla*<sub>SHV</sub>, *bla*<sub>GES</sub>, *bla*<sub>CARB</sub>, *bla*<sub>PER</sub>, *bla*<sub>IMP</sub>, *bla*<sub>CMY</sub>, *bla*<sub>SME</sub>, *bla*<sub>VIM</sub>, *bla*<sub>OXA</sub>, *bla*<sub>NDM</sub>, and *bla*<sub>KPC</sub>. The primers and protocol were adopted from our lab,<sup>17</sup> whereas the PCR for colistin-resistant genes (*mcr-1* to *mcr-9*) was performed by using primers described previously.<sup>18</sup> The PCR conditions are well established in our laboratory. Thermocycler conditions were set at 94°C for 5 min, followed by 35 cycles of denaturation at 94°C for 30 sec, annealing at 55°C for 30 sec and extension at 72°C for 1 min, followed by a final extension at 72°C for 5 min. The amplified product was run on 1.5% agarose gel with ethidium bromide and extracted using a QIAquick® Gel Extraction Kit by QIAGEN (Germany) followed by sequencing. The Sequences were analysed by BlastN: [https://blast.ncbi.nlm.nih.gov/Blast.cgi?PAGE\\_TYPE=BlastSearch](https://blast.ncbi.nlm.nih.gov/Blast.cgi?PAGE_TYPE=BlastSearch). The clonal relation and MLST typing were performed by amplifying conserved seven housekeeping genes including *acsA*, *aroE*, *guaA*, *mutL*, *nuoD*, *ppsA*, and *trpE*, according to the protocol mentioned on the *P. aeruginosa* MLST website (<https://pubmlst.org/P.aeruginosa/>). The primers for the above housekeeping genes were adopted from the same website. The amplified products were sequenced and Allelic profiles, as well as sequence types, were determined using <https://pubmlst.org/bigsub?db=pubmlstpaeruginosa> seqdefandpage=profilesandscheme\_id=1. The combination of alleles obtained at each locus defined its allelic profile or sequence type (ST). The combinations of alleles did not match with the locus database designated as a “new” ST. While the clonal relatedness was determined by the repetitive PCR (rep-PCR) (Diversilab, Biomerieux). All protocols were adapted from our laboratory.

## Conjugation and Plasmid Characteristics

A mating assay was performed to analyse the horizontal gene transfer of *bla*<sub>NDM-1</sub> and *mcr-1* by using streptomycin-resistant *E. coli* C<sub>600</sub> as the recipient strain. The Luria broth medium for both isolates was mixed and incubated for meeting at 37°C for 24hrs. The successful transconjugants were selected on streptomycin (2000  $\mu$ g/mL) and colistin (4  $\mu$ g/mL) seeded Muller Hinton agar plate. While *bla*<sub>NDM-1</sub> conjugants were selected on streptomycin (2000  $\mu$ g/mL) and meropenem (1  $\mu$ g/mL) seeded Muller Hinton agar plate. Furthermore, transconjugants were tested by PCR assay and sequencing for the presence of *mcr-1* and co-existence of other genes including *bla*<sub>NDM-1</sub>, *bla*<sub>KPC</sub>, and *bla*<sub>CTX-M-15</sub>. The transconjugant frequency was determined as the number of transconjugants divided by the total colony-forming unit (CFU). In addition, the plasmid incompatibility (Inc) group was determined based on PCR-based replicon typing (PBRT). The primer pairs adopted based on the EMBL Gene Databank, for HI1, HI2, II-Iy, X, L/M, N, FIA, FIB, W, Y, P, FIC, A/C, T, FIIA<sub>S</sub>, F, K and B/O replicons results analysed as per described.<sup>19</sup> Degenerate primer MOB typing (DPMT) means sub-groups were determined by using specific primers which were encoded in 33 references release as previously introduced by Alvarado A et al,<sup>20</sup> while protocol was standardised in our laboratory.

## Result

All *P. aeruginosa* isolates were confirmed by API-20 and 16s RNA sequencing. The modified Hodge test shows the inactivation of the carbapenem resulting in positive for carbapenemase production while MBL-E-test results confirmed the production of Metallo-beta-lactamase for all tested strains. The AST results revealed that in all the tested carbapenemase-producing *P. aeruginosa* isolates 6/6, 100% were resistant to amoxicillin, amikacin and ticarcillin; while 60% of these isolates were resistant to other tested antibiotics (Figure 1). About 3/6, 50% were resistant to colistin which is the



**Figure 1** Antibiotic susceptibility testing of clinical isolates *Pseudomonas aeruginosa*.

last resort of choice to treat infection caused by MDR strains. The resistant determinant results showed that all isolates harboured *bla*<sub>NDM-1</sub> among which four co-harboured *mcr-1*. In addition, carbapenemase-producing *P. aeruginosa* were carrying *bla*<sub>KPC-1</sub> (4/6), followed by *bla*<sub>IMP</sub> (3/6), *bla*<sub>GES</sub> (1/6), and *bla*<sub>CMY</sub> (1/6). Colistin use is one of the major factors responsible for the development of colistin resistance but in this study, patients were not treated with colistin (Table 1). In our study, MLST results indicate that 2/6 isolates belonged to *P. aeruginosa* ST1966 high-risk clone, the other four isolates allocated alleles combination were not matched with reported STs in PubMLST database so reported as new STs (Table 2). A total of three successful *mcr-1* carrying transconjugants were obtained while one-SP-19-2019 failed to transfer the *mcr-1* gene (Table 1). Our isolate SP-19-2019 conjugation failure may be due to the location of *mcr-1* on the chromosome. The horizontal gene transfer frequency ranged from  $10^{-3}$  to  $10^{-5}$ . Successful three conjugants co-existence of *bla*<sub>NDM-1</sub> and *mcr-1* which were located in IncX4 plasmid group based on the PBRT results while MOB revealed that plasmid belongs to the P-31 subgroup (Table 1). Moreover, two *bla*<sub>NDM-1</sub> plasmids of transconjugants belong to IncFIS and subgroup P-12. In genotyping, we found that SP-19-1051, 1054 and 2019 were clonally related (>95%). The other three isolates (Sp-19-2014, 2015, 2016) were of the same sequence type. Both patients had invasive infections with colistin-resistant *P. aeruginosa* but were able to recover as a result of antimicrobial therapy and were discharged.

**Table 2** Combination of Alleles with Sequences Type

Locus	Tm	Isolates Number					
		1051	1054	2014	2015	2016	2019
<i>acsA</i>	55	15	15	15	15	15	15
<i>aroE</i>	56	5	5	4	4	4	4
<i>guaA</i>	55	11	11	11	11	11	11
<i>mutL</i>	61	5	5	5	5	5	5
<i>nuoD</i>	58	4	4	4	4	4	4
<i>ppsA</i>	55	3	3	3	3	3	3
<i>trpE</i>	56	141	141	141	141	141	141
	ST	1966	1966	New	New	New	New
	Complex	N/D	N/D	N/D	N/D	N/D	N/D

**Abbreviations:** ST, sequence type; N/D, not detected; Tm, melting temperature.

## Discussion

Infections caused by *P. aeruginosa* are difficult to treat since this bacterium has several mechanisms of resistance and virulence, such as the acquisition of plasmids or integrons.<sup>21</sup> The presence of colistin resistance plasmid-mediated *mcr-1* in *P. aeruginosa* bacteria has drawn global attention because colistin last resort to treat infection caused by MDR pathogens.<sup>22</sup> Therefore, there is an urgent requirement for novel antibiotics with new modes of action or alternative approaches to identifying molecules that can interfere with the process of efflux such as P-glycoprotein.<sup>23</sup> Our previous studies describe the co-production of MCR-1 and NDM-1 in Extended-Spectrum  $\beta$ -Lactamase-Producing *Escherichia coli* ST648 Isolated from a Colonized Patient in China.<sup>9</sup> In the present study, the isolates were resistant to the commonly used antimicrobials to treat Gram-negative infections. Interestingly, we observed that three among six were resistant to colistin, which is the last choice of drug to treat infection caused by MDR Gram-negative bacteria.<sup>24</sup> These antimicrobial susceptibility results are in agreement with other studies that demonstrated the high levels of resistance in *P. aeruginosa* recovered from bloodstream infections.<sup>25</sup> We reported carbapenemase encoding *bla*<sub>NDM-1</sub> from six strains which were the first detected in *P. aeruginosa* in Serbia in the year 2011.<sup>26</sup> In the present study, three strains co-harboured colistin resistance *mcr-1* and carbapenemase encoding *bla*<sub>NDM-1</sub>, to the best of our knowledge; this has not yet been reported. Nevertheless, the co-existence of *mcr-1* and *bla*<sub>CTX-M</sub> has already been reported earlier.<sup>27</sup> In our study, we reported 2/6 *P. aeruginosa* that belong to ST1966, which is similar to a Chinese strain reported in 2011 from the Peking Hospital, Beijing (unpublished data). In addition, previous studies have reported that the novel sequence type strains are evolving, and increasing the mortality rate of *P. aeruginosa* bacteraemia,<sup>28,29</sup> this highlights the need to further investigate this strain by the application of whole-genome sequencing. We have 3/4 *mcr-1* conjugants while one failed, we suppose the location of a gene on the chromosome. The incidence of chromosomally encoded *mcr-1* is rare although it has been reported in several cases.<sup>30</sup> Here, we demonstrate that the *mcr-1* gene is located on IncX4 plasmids, the plasmids that have been reported to play a significant role in the spread of the *mcr-1* gene.<sup>31,32</sup> Although there are limitations to this study, these findings help fill the knowledge gap that has hindered the mechanism of resistance and surveillance for this strain and can be used for further investigation. In addition, we need to consider the resistance mechanisms in MDR *Pseudomonas* as well as the related virulence factor and biofilm production and develop new correlations among these which may help for both diagnostic and therapeutic purposes.<sup>33</sup>

## Conclusion

This is the first report to detect *bla*<sub>NDM-1</sub> and colistin resistance *mcr-1* in MDR *P. aeruginosa* isolates recovered from bloodstream infection in immunocompromised patients during leukaemia treatment. Colistin resistance is emerging in *P. aeruginosa* in a novel sequence type and may have a high-risk clone. The spread of carbapenemases in clinical isolates highlights the need for continuous surveillance for high-risk clones of *P. aeruginosa*, particularly in paediatric patients.

## Abbreviations

NDM, New Delhi Metallo-beta-lactamase; AMR, Antimicrobial-resistant; EUCAST, European Committee on Antimicrobial Susceptibility Testing; PCR, Polymerase Chain Reaction; PBRT, PCR-based replicon typing; MLST, Multi-locus sequence typing (MLST) method and (PFGE); Inc, Incompatibility; STs, Sequence types.

## Ethics Approval and Consent to Participate

The study was approved by the Shenzhen Children's Hospital (Research) ethics committee (Reference number: 2018 (013)).

## Consent to Publish

The clinical isolate samples used in this research were part of the routine Hospital Laboratory procedure. Because we did not use the patient's name or personal information, therefore no written informed consent was required. In addition, verbal consent was taken from the parents' patients but not written.

## Author Contributions

All authors made a significant contribution to the work including conceptualization, study design, execution, acquisition of data, analysis and interpretation, and were involved in drafting, revising and critically reviewing the manuscript. The final version of this manuscript has been approved by all authors and agreed to be accountable for all aspects of the work.

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## Disclosure

The authors declare no competing conflicts of interest in this work.

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