Isolation of *NDM-1*-producing multidrug-resistant *Pseudomonas putida* from a paediatric case of acute gastroenteritis, India

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Abstract

Pseudomonas putida is an uncommon opportunistic pathogen, usually susceptible to antimicrobial agents. Data concerning resistance to antimicrobial agents in clinical *P. putida* isolates are limited. To the best of our knowledge we report for the first time the isolation of *NDM-1*-producing multidrug-resistant *P. putida* from a case of acute gastroenteritis. The isolate showed resistance to a wide range of antimicrobials, including fluoroquinolones, third-generation cephalosporins and carbapenems. The isolate also exhibited multiple mutations in the quinolone resistance determining region and showed the presence of *qepA*, bla_{TEM}, bla_{OXA1} and bla_{OXA7} genes. The present study highlights the importance of looking for the relatively rare aetiological agents in clinical samples that do not yield common pathogens.

Keywords: 16S rRNA, diarrhoea, multidrug resistance, mutation, NDM-1, *Pseudomonas putida*

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Introduction

Pseudomonas putida, a non-fermenting Gram-negative bacillus belonging to the fluorescent group of the genus *Pseudomonas* is frequently found in the environment, along with other non-fermenting Gram-negative organisms. Previously thought to be of low pathogenicity [1], there are increasing reports of their emergence as opportunistic human pathogens causing bacteraemia and sepsis in neonatal, neutropenic and cancer patients, as well as in people with urinary tract infections [2,3]. Most *P. putida* are susceptible to antimicrobial agents such as fluoroquinolones, aminoglycosides and carbapenems [4]. In the present study, we report the isolation of an *NDM-1*-producing multidrug-resistant *P. putida* from a 2-month-old female child admitted to a tertiary-care hospital with acute gastroenteritis in Belgaum, South India. To the best of our knowledge this is the first report of isolation of *NDM-1*-producing multidrug-resistant *P. putida* causing acute gastroenteritis.

A female child aged 2 months was admitted to the gastroenteritis ward of a tertiary-care hospital in Belgaum, Karnataka, South India, with symptoms of acute gastroenteritis on 21 June 2013. She had had watery diarrhoea for 3 days along with vomiting for 5 days, showed signs of acute dehydration and had fever of 38.9°C. The fever was intermittent in nature, associated with chills and rigor. The patient was lethargic, restless and had sunken eyes. The patient was put on rehydration therapy, a stool sample was collected for laboratory investigations before administration of any antibiotic at the hospital, and the patient was later empirically treated for acute diarrhoea with oral ciprofloxacin (50 mg) twice daily and metronidazole (25 mg) thrice daily. She recovered and was discharged on 26 June 2013.

The stool sample was processed for isolation and identification of common enteric bacterial pathogens, which include diarrheagenic *Escherichia coli*, *Shigella* sp., *Salmonella* sp., *Vibrio* sp. following WHO 1987, and was subjected to ELISA for rotavirus, RT-PCR for identification of common viral pathogens like norovirus, astrovirus and sapovirus [5]. The sample was also subjected to routine microscopy for detection of various parasites like *Ascaris lumbricoides*, *Giardia lamblia*, *Trichuris trichiura*, hook worm and *Entamoeba histolytica*. The isolate was subjected to identification based on an automated microbial identification system, (Vitec2 Compact; bioMérieux, Marcy l'Etoile, France) which was also used for carrying out Antibiotic sensitivity testing (AST) as per CLSI norms [6]. The identity of the isolate was also confirmed by genotypic-based method of *16S rRNA* gene sequencing as defined earlier [7]. The rabbit ileal loop test was carried out for the isolate essentially as described by Koley *et al* [8]. The volume of the accumulated fluid in

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millilitres and the length of the loop in centimetres were measured, and the extent of the fluid accumulation was expressed as mL/cm. The ileal loop test was performed with positive and negative controls being *Vibrio cholerae* O1 (N16961) and phosphate-buffered saline, respectively.

The isolate was further screened for any mutation in the quinolone resistance determining region following an earlier described protocol (Table 1) [9]. Presence of plasmid-mediated quinolone resistance determinants was screened following standard conditions (Table 1) [10]. The isolate was also subjected to PCR for detection of the presence of various β-lactam resistance genes (bla_{TEM}, bla_{SHV}, $bla_{CTX\text{-}M\text{-}3}$, bla_{OXA1} and bla_{OXA7}) following techniques reported previously (Table 1) [10]. Presence of the *NDM-1* gene was determined by PCR using published primers (Table 1) as described earlier [11]. All PCR products were subjected to nucleotide sequencing in an automatic sequencer (ABI 3130; Applied Biosystems, Foster City, CA, USA) following the manufacturer's instructions. Contig sequences were aligned and edited with SeqScape v2.7 (Applied Biosystems) and compared in BLAST of the NCBI database.

Culture on thiosulphate citrate bile salt sucrose agar and Hektoen enteric agar plates did not yield any isolate whereas on McConkey agar, a pure culture of non-lactose-fermenting colonies appeared. The isolate was identified as *P. putida* by an automated microbial identification system, which was confirmed by *16s rRNA* sequence analysis. The sample was negative for all other bacteria, viruses and parasites tested. Growth of the organism as sole pathogen in the culture media in the absence of any other viral, bacterial or parasitic organism indicates the colonization of the gut of the patient by *P. putida*, probably through suppression of normal microbiota. The *P. putida* isolate showed resistance to a wide range of antimicrobials, including fluoroquinolones, third-generation cephalosporins and carbapenems, according to CLSI breakpoint [6]. The isolate showed a MIC (mg/L) of 32 for Ampicillin-sulbactam (SAM), ≥128 for Ticarcillin (TIC), ≥128 for Piperacillin (PIP), ≥64 for Ceftazidime (CAZ), ≥64 for Ceftriaxone (CRO), ≥64 for Cefepime (FEP), ≥16 for Imipenem (IMP), ≥16 for Meropenem (MEM), ≥16 for Amikacin (AMK), ≥32 for Gentamicin (GEN), ≥16 for Tobramycin (TOB), ≥4 for Ciprofloxacin (CIP), ≥8 for Levofloxacin (LVX), ≥16 for Tetracycline (TET), ≥8 for Tigecycline (TGC) and ≥320 for Co-trimoxazole (CoT).

The isolate was tested and found positive for the production of extended spectrum β-lactamase using the combination disc test using ceftazidime-clavulanic acid (30/10 μg) and ceftriaxone -clavulanic acid (30/10 μg) [10]. The *16S rRNA* gene sequence of this isolate was also compared with other sequences submitted to the NCBI GenBank to understand its genetic relationship with other *P. putida* by neighbour-joining phylogenetic analysis using MEGA 5.2 software [12], which showed 100% similarity to *P. putida* isolated from various other parts of the world (Fig. 1). The isolate resulted in moderate fluid accumulation (0.35 mL/cm), which was higher than the negative control (0.05 mL/cm) and lower than the positive control (1.2 cm/mL).

The isolate exhibited amino acid substitution of T83I and S136A in *gyrA*; E469D in *gyrB*; and T105P, V124A and S136A in

TABLE 1. Details of the primers, amplification temperature and amplicon size used in the study

Sl No.	Gene	Oligonucleotide sequence (5′–3′)	Amplification temp (°C)	Amplicon size (bp)
1	*NDM-1*	ACCGCCTGGACCGATGACCA GCCAAAGTTGGGCGCGGTTG	58°C	264
2	*gyrA*	GACGGCCTGAAGCCGGTGCAC GCCCACGGCGATACCGCTGGA	64°C	417
3	*gyrB*	AGTACTTCGCCGACTTCCT TACAGGCGCGACAGGCGCTT		739
4	*parC*	TCTACGCCATGAGCGAACTGG AGCAGCACCTCGGAA TAGCG		262
6	*qnrA*	ATTTCTCACGCCAGGATTTG GATCGGCAAAGGTTAGGTCA	64°C	516
7	*qnrB*	GATCGTGAAAGCCAGAAAGG ATGAGCAACGATGCCTGGTA		476
8	*qnrC*	GGGTTGTACATTTATTGAATCG CACCTACCCATTTATTTTCA		307
9	*qnrS*	GCAAGTTCATTGAACAGGGT TCTAAACCGTCGAGTTCGGCG		428
10	*aac (6′)-Ib-cr*	TTGCGATGCTCTATGAGTGGCTA CTCGAATGCCTGGCGTGTTT	55°C	482
11	*qepA*	AACTGCTTGAGCCCGTAGAT GTCTACGCCATGGACCTCAC		596
12	bla_{TEM}	GAGTATTCAACATTTTCGT ACCAATGCTTAATCAGTGA	50°C	857
13	bla_{SHV}	TCGCCTGTGTATTATCTCCC CGCAGATAAATCACCACAATG		768
14	$bla_{CTX\text{-}M\text{-}3}$	AATCACTGCGTCAGTTCAC TTTATCCCCCACAACCCAG		701
15	bla_{OXA1}	GCAGCGCCAGTGCATCAAC CCGCATCAAATGCCATAAGTG		198
16	bla_{OXA7}	AGTTCTCTGCCGAAGCC TCTCAACCCAACCAACCC		591