RESEARCH ARTICLE

Isolation of five Enterobacteriaceae species harbouring $\text{bla}_{\text{NDM-1}}$ and $\text{mcr-1}$ plasmids from a single paediatric patient

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Abstract

In Argentina, NDM metallo-β-lactamase was first reported in 2013. By now, it has disseminated throughout the country in diverse Gram negative bacteria. Here, we report the case of a paediatric patient that underwent a 1-year hospitalisation due to erythrodermic psoriasis. During his stay, five isolates were obtained from rectal swabs (rs) or blood culture (bc) suspicious of carbapenemase production: a K. quasipneumoniae subsp. quasipneumoniae (rs), Citrobacter freundii (rs), Escherichia coli (bc), Enterobacter cloacae (rs), and a Serratia marcescens (bc). The isolates were studied with broth microdilution, biparental conjugation and plasmid and whole genome sequencing (Illumina). All isolates harbour an 138,998-bp type 1 IncC plasmid that carried $\text{bla}_{\text{NDM-1}}$, $\text{bla}_{\text{TEM}}$, $\text{bla}_{\text{CMY-6}}$, $\text{rmtC}$, $\text{aac(6')-Ib}$-cr1, and $\text{sul1}$ resistance genes. Additionally, the $\text{bla}_{\text{NDM-1}}$-plasmids contained ISKpn8 an insertion sequence previously described as associated only to $\text{bla}_{\text{KPC}}$. One isolate, a colistin-resistant $E$. coli, also carried a $\text{mcr-1}$-containing an IncI2 plasmid, which did not harbour additional resistance. The whole genome of K. quasipneumoniae subsp. quasipneumoniae isolate was fully sequenced. This isolate harboured, additionally to $\text{bla}_{\text{NDM-1}}$, three plasmid-mediated quinolone resistance genes: $\text{qnrB4}$, $\text{qnrB52}$ and $\text{aac(6')-Ib-cr1}$. The E. cloacae isolate also harboured $\text{qnrA1}$. These findings alert to the underestimated horizontal dissemination of multidrug-resistant plasmids limiting treatment options with last resort antimicrobials.

Introduction

New Delhi metallo-β-lactamase (NDM) is a plasmid-borne carbapenemase that severely limits treatment options against gram-negative pathogens. The $\text{bla}_{\text{NDM-1}}$ gene was initially identified in Klebsiella pneumoniae and Escherichia coli isolates but later it was reported in many Gram
negative species like *Klebsiella oxytoca*, *Proteus mirabilis*, *Enterobacter cloacae*, *Citrobacter freundii*, *Providencia* spp., *Acinetobacter* spp., *Pseudomonas aeruginosa* and *Vibrio cholerae* [1]. Some of these bacteria are carried as gut flora and are commonly found in the environment, providing reservoirs for future infections [2].

The *bla*\textsubscript{NDM-1} gene is carried on plasmids usually carrying additional resistance genes that compromise antimicrobial treatment. Plasmids associated to the dissemination of *bla*\textsubscript{NDM-1} vary greatly in attributes such as size, gene content, organization and incompatibility group or replicon type (e.g. IncA, IncC, IncF types, IncL/M, IncN, IncX, and IncH) [3, 4].

The plasmids harbouring *mcr-1* have been identified in multidrug-resistant *Enterobacteriaceae* isolates, some of which co-produced carbapenemases such as KPC, VIM or NDM [5].

Here, we report five *bla*\textsubscript{NDM-1}-producing *Enterobacteriaceae* species isolated from the same patient, one of which co-produced MCR-1. We also characterised the genetic elements that harbour *bla*\textsubscript{NDM-1} and *mcr-1*, and the high similarities found indicate the presence of one *bla*\textsubscript{NDM-1} plasmid circulating among these different species.

**Case study**

In February 2014, a 4-year-old child suffering from erythrodermic psoriasis with previous hospitalisations was admitted to the hospital due to edematous-ascitic syndrome. After 124 days of hospitalisation the patient died as a result of septic shock. During the patient’s hospitalisation, five *Enterobacteriaceae* species suspicious of carbapenemase production due to imipenem inhibition halos ≤22mm were isolated. As a consequence, he was treated with standard doses of colistin and tigecycline. The timeline of isolation is as follows: at day 39 of hospitalisation, *K. pneumoniae* M17277 was isolated from a rectal swab, later identified as *K. quasipneumoniae* subsp. *quasipneumoniae*; at day 71, *C. freundii* M17394 from a rectal swab; at day 74, *E. coli* M17386 from a blood culture; at day 117, *E. cloacae* M17464 from a rectal swab and finally, at day 123, *Serratia marcescens* M17468 was obtained from a blood culture. Rectal swabs were periodically obtained as part of the hospital’s surveillance programme to detect KPC-producing *K. pneumoniae*. These isolates were sent to the National Reference Laboratory on Antimicrobial Resistance (NRLAR) for further characterisation.

In addition to the clinical situation described above, the patient was also infected and colonized with several microorganisms obtained from rectal swabs, blood cultures or catheter tips as follows: *Staphylococcus aureus*, *Staphylococcus haemolyticus*, *Enterococcus faecalis*, *Enterococcus faecium*, *Acinetobacter baumannii*, *P. aeruginosa*, and *Candida parapsilosis*. In addition, *Candida glabrata*, *C. albicans* and *Toxocara* spp. were isolated from urine cultures. The patient was treated according to the infecting microorganism with: clindamycin, ceftriaxone, vancomycin, cephalotin, trimethoprim-sulfamethoxazole, colistin, tigecycline, rifampicin, meropenem, minocycline, linezolid, amikacin, gentamicin, ampicillin-sulbactam, teicoplanin, and metronidazol (due to *Giardia lamblia*). In addition, the patient was treated with immunosuppressors and antimycotics, as albendazol and fluconazol to treat the mycotic and parasitic infections. Unfortunately, exact dates of isolation of the mentioned isolates are missing in the hospital records.

**Materials and methods**

**Microbiological identification, antimicrobial susceptibilities and PCR screening for resistance genes.**

Isolates were identified using conventional biochemical tests. All Gram negatives, oxidase negative isolates were tested for *Enterobacteriaceae* species considering lactose fermentation in...
CLDE media and the following tests: TSI (Triple Sugar Iron), SIM (Sulfide Indole Motility), Citrate media, LIA (Lysine Iron Agar) and MIO (Motility, Indole, Ornithine). These media were purchased in Laboratorios Britania S.A., Argentina. Species identification were confirmed with matrix-assisted laser desorption ionization time of flight (MALDI-TOF) mass spectrometry (Bruker, Germany). Carbapenemase activity was screened with: Triton-Hodge Test, Blue-Carba Test and Carba-NP-Direct test (in house), [6–8]. Screening of metallo-β-lactamase activity was done by synergy between a meropenem (10 μg) disc and Ethylenediaminetetraacetic acid/Sodium Mercaptoacetate (EDTA/SMA) (Laboratorios Britania S.A., Argentina).

Antibiotic susceptibility profiles were determined by agar dilution according to the guidelines and interpretation criteria of the Clinical and Laboratory Standards Institute (CLSI 2018). MICs were determined for imipenem, meropenem, ceftazidime, cefotaxime, cefepime, aztreomycin, ciprofloxacin, amikacin, gentamicin, fosfomycin, tigecycline, minocycline and colistin.

PCRs were performed using the primers listed in S1A Table to detect the resistance genes: \textit{bla}_{PER}, \textit{bla}_{CTX-M}, \textit{bla}_{KPC}, \textit{bla}_{CMY-2/7}, \textit{bla}_{VIM}, \textit{bla}_{IMP}, \textit{bla}_{NDM}, \textit{mcr}-1, \textit{rmtC}, \textit{aac(6')-Ib-cr} and \textit{qnr}-type (C, D, A, B, S and E) (S1A Table).

**Biparental conjugation**

The agar mating method was used to conjugate clinical isolates with E. coli J53 sodium azide-resistant (Az\textsuperscript{R}) as recipient [9]. \textit{bla}_{NDM-1} transconjugants were selected in meropenem (containing 0.2 μg/ml) Luria-Bertani plates plus sodium azide (200 μg/ml). E. coli M17386 harbouring \textit{bla}_{NDM-1} and \textit{mcr}-1 was conjugated to a susceptible Salmonella spp. strain M1744 (S1B Table). Transconjugants were identified using Salmonella-Shigella agar plates supplemented with meropenem (0.2 μg/ml) or colistin (2 μg/ml). Subsequently, Salmonella transconjugants producing NDM or MCR were separately conjugated with E. coli J53 Az\textsuperscript{R} as explained above, and the retrotranscojugants were selected as explained above.

**Plasmid characterisation**

Plasmid content was analysed by S1 nuclease (Promega, Southampton, UK) digestion followed by PFGE, as described [10]. Plasmids were blotted onto a positively charged nylon membrane (Bio-Rad, Hercules, USA) and hybridised with a \textit{bla}_{NDM-1} probe (ECL Direct Nucleic Acid Labelling And Detection, Buckinghamshire, England).

\textit{bla}_{NDM-1}\textsuperscript{-} and \textit{mcr}-1-harbouring plasmids were fully characterised as follows. Plasmids were extracted from transconjugants with Qiagen Large-Construct kit (Qiagen, Hilden, Germany) and sequenced on a MiSeq sequencer with MiSeq Reagent Kit v3 (Illumina, USA).

Plasmid characterisation was performed with the CLC Genomics Workbench software (CLC bio, Qiagen) and gaps were closed by PCR using the primers depicted in S1 Table, followed by Sanger sequencing. Open reading frames were predicted and annotated by Prokka v1.12 [11], and manual curation in Artemis [https://www.sanger.ac.uk/science/tools/artemis] and BLAST search [https://blast.ncbi.nlm.nih.gov/Blast.cgi]. ResFinder and PlasmidFinder were used to identify resistance genes and plasmid incompatibility groups, respectively [https://cge.cbs.dtu.dk/services/]. The insertion sequences were identified using ISfinder [https://www-is.biotoul.fr].

Whole genome sequencing (WGS) and phylogenetic analysis of M17277

Whole bacterial DNA was extracted with QIAcube, using the QIAGEN\textsuperscript{R} DNA Mini Kit (Qiagen) and sequenced on a MiSeq sequencer. Reads were de novo assembled using VelvetOptimiser v2.2.5 [12]. Automated annotation was done with Prokka [11].
Phylogenetic analysis of M17277 was done with a WGS dataset from 60 type-strains and isolates of K. pneumoniae, K. quasipneumoniae subsp. quasipneumoniae, K. quasipneumoniae subsp. similipneumoniae and Klebsiella variicola (namely, the Kleb-dataset). This dataset comprised: two draft genomes of the type-strains of K. quasipneumoniae subsp. quasipneumoniae and K. quasipneumoniae subsp. similipneumoniae, namely 01A030\textsuperscript{T} and 07A044\textsuperscript{T}, respectively [13], and genome assemblies from 58 isolates selected from a collection of 288 isolates previously characterised [14]. These 58 isolates comprised all K. quasipneumoniae subsp. quasipneumoniae and K. variicola available in the Holt’s database (1, 19 and 18 isolates, respectively), and 20 isolates randomly selected from the subset of K. pneumoniae isolates of the Holt’s database (S3 Table) [14]. The multifasta files of the type strains 01A030\textsuperscript{T} and 07A044\textsuperscript{T} were downloaded from NCBI accession numbers CCDF01000000 and CBZR010000000, respectively, and annotated with Prokka [11]. The annotated assemblies of M17277 and the Kleb-dataset were used in a pan-genome analysis with Roary [15] to generate a core-gene alignment (concatenated genes present in $\geq99\%$ of the genomes with $\geq95\%$ of nucleotide identity). This core-genome alignment was used to generate a single-nucleotide polymorphism (SNP) alignment with SNP-sites v2.3.2 [16] used to construct a maximum likelihood phylogenetic tree with RAxML v8.2.8 [17] under the generalised time reversible model (GTRCAT) and bootstrapping with 1,000 replicates. To support the re-classification of K. quasipneumoniae subsp. quasipneumoniae we used a genome-to-genome distance approach using the online tool at https://ggdc.dsmz.de/ggdc.php [18].

Nucleotide sequence accession number
Plasmid sequences were submitted to GenBank under accession no. MH995506 (TC-17394), MH995507 (TC17277), MH995508 (TC-17464), MK123267 (RT-17386), MK123268 (TC-17468). K. quasipneumoniae subsp. quasipneumoniae M17277 assembly was submitted under BioProject PRJNA499048 and BioSample: SAMN10340875, GenBank RZJN00000000.

Results
Species identification, antimicrobial susceptibilities and resistance genes
The five species were originally identified as K. pneumoniae M17277, E. coli M17386, C. freundii M17394, E. cloacae M17464 and S. marcescens M17468. M17277 was one of the first bla\textsubscript{NDM-1}-harbouring clinical isolates belonging to the genus Klebsiella that had been sent to the NRLAR (2014). K. pneumoniae M17277 was identified as K. pneumoniae by biochemical techniques and MALDI-TOF.

The screening tests for carbapenemase activity were positive for all isolates and showed synergy between a carbapenem disc and EDTA/SMA, indicating the production of a metallo-β-lactamase.

Full resistance phenotype to aminoglycosides (gentamicin and amikacin) was observed in all isolates. Additionally, K. pneumoniae M17277, E. coli M17386, C. freundii M17394, E. cloacae M17464 were resistant to all β-lactams including carbapenems and aztreonam while S. marcescens M17468 was susceptible to this last drug (Table 1). E. coli M17386 was resistant to colistin. Three isolates were resistant to minocycline, two to ciprofloxacin, and two to tigecycline. All isolates remained susceptible to fosfomycin (Table 1).

In line with those results, PCR assays showed that all the clinical isolates carried bla\textsubscript{NDM-1} and rmtC while E. coli M17386 additionally carried mcr-1. The plasmid mediated quinolone resistance genes PMQRs detected in K. pneumoniae M17277 were qnrB4 and qnrB52 and are described in the text below. E. cloacae M17464 harboured qnrA1 which could explain the resistance to ciprofloxacin detected.
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Abbreviations used: R, resistant; I, intermediate; S, susceptible; ND, Not determined; TC, transconjugant; RT, retrotransconjugant; KQN, K. quasipneumoniae subsp. quasipneumoniae; ECO, E. coli, CFR, C. freundii; ECL, E. cloacae; SMA, S. marcescens.

<sup>a</sup>MICs were interpreted according to CLSI 2018 guidelines.

<sup>b</sup>Serratia spp. are naturally resistant to colistin.

<sup>c</sup>Fosfomycin, tigecycline and colistin MICs were interpreted following the breakpoints of EUCAST.

E. coli J53-Az<sup>b</sup>, biparental conjugation acceptor strain, azide resistant.

RT were obtained from ECO-M17386, RT-NDM-M17386 harbours bla<sub>NDM</sub> plasmid and RT-MCR-M17386 harbours a bla<sub>NDM-1</sub>/mcr plasmid.

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Horizontal transfer of \( bla_{NDM-1} \) and \( mcr-1 \)-harbouring plasmids

All \( bla_{NDM-1} \)-harbouring plasmids were transferred by conjugation and the susceptibility profiles of transconjugants are shown in Table 1. Together with carbapenem resistance mediated by NDM-1, high level resistance to aminoglycosides was co-transferred mediated by RmtC. Resistance to cephalosporins was transferred mediated by CMY-6. On the contrary, resistance to ciprofloxacin, aztreonam, tigecycline and minocycline was not transferred to transconjugant strains, demonstrating that these resistance determinants may be encoded in the chromosome, in a non-mobile plasmid or in another plasmid that was not transferred.

\( E. \) coli-M17386 harboured both \( bla_{NDM-1} \) and \( mcr-1 \). Therefore sequential conjugations were performed to demonstrate that both genes were located on separate conjugative plasmids. In consequence, we obtained two retrotransconjugants that each expressed NDM-1 (RT-NDM-M17386) or MCR-1 (RT-MCR-M17386) (Table 1).

Whole plasmid characterisation

All isolates harboured 2 to 4 plasmids (48.5–470 kb) as observed by S1 nuclease-PFGE (S1A Fig). The \( bla_{NDM-1} \) probe hybridised with one band of ca. 138-kb in all clinical isolates and their respective transconjugants (S1B Fig). Full sequencing showed that all isolates harboured a plasmid of 138 kb carrying \( bla_{NDM-1} \). The \( bla_{NDM-1} \) Plasmids from \( K. \) pneumoniae M17277, \( E. \) coli M17386 and \( C. \) freundii M17394, were 100% identical while those from \( E. \) cloacae M17464 and \( S. \) marcescens M17468 showed 3 and 10 SNPs, respectively, with the former three, and 7 SNPs between them. The plasmid of isolate \( K. \) pneumoniae M17277 (pKQN17277) was selected for the subsequent analysis. pKQN17277 showed maximal identity (99.0%) with pKP1-NDM-1 (acc. no. KF992018) a type 1 IncC obtained from a clinical isolate of \( K. \) pneumoniae in 2010 in Australia from a patient arrived from India [19]. Interestingly, besides 3 SNPs, the unique difference between both plasmids was the presence of the insertion sequence IS\( Kpn8 \) (acc. no. EF382672) in pKQN17277 (Fig 1) not associated to any resistance gene [20]. Moreover, all the NCBI entries that showed 98% cover query and around 99% identity with pKQN17277 did not harbour IS\( Kpn8 \). Given the high similarity with pKP1-NDM-1, the plasmids characterised here harboured the main hallmarks of type 1 IncC plasmids (Fig 1A and 1B) including rhs1 (unknown biological function) [21], orf1832 and ARI-A, which contained rmtC and \( bla_{NDM-1} \) [21] (Fig 1B). This last gene was located within a truncated TnI25 structure previously reported [4] (Fig 1B). Additionally, these plasmids harboured \( bla_{CMY-6} \) associated to IS\( Ecpl \), which was located outside ARI-A (Fig 1). On the other hand, pKQN17277 was not related to the \( bla_{NDM-1} \)-harbouring plasmid of \( K. \) quasipneumoniae CCBH16302 strain reported in Brazil which was isolated in 2014 (ca. 346 kb, acc.no. MDCA00000000) [22]. Moreover, this last plasmid did not contain rmtC and harboured \( bla_{NDM-1} \) associated with Tn3000 [17].

Full sequencing of the \( mcr-1 \)-harbouring plasmid of \( E. \) coli M17386 (pMCR-M17386) showed that this gene was located in a ca. 61-kb IncI2 plasmid, which was 100% identical to pMCR-M15224 (acc. no. KY471309) isolated from a clinical isolate of \( C. \) amalonaticus in 2016 from the same paediatric hospital than the case reported here [Faccone, 2018 #4738]. No other antimicrobial resistance gene was identified in pMCR-M17386.

Phylogenetic analysis of the isolate M17277

M17277 was one of the first \( bla_{NDM-1} \)-harbouring clinical isolates belonging to the genus \( Klebsiella \) that had been sent to the NRLAR (2014). M17277 was identified as \( K. \) pneumoniae by biochemical techniques and MALDI-TOF. However, these methodologies did not discriminate between \( K. \) pneumoniae, \( K. \) quasipneumoniae and \( K. \) variicola [13]. Therefore, M17277
was subjected to WGS and pan-genome analysis in the context of a global collection (S2 Table) [14]. The phylogenetic tree (S3 Table) showed that the isolate M17277 clustered with 100% bootstrap support with the type strain 01A030\textsuperscript{T} and the unique isolate of \textit{K. quasipneumoniae} (Fig 2). In order to validate the species identification, we determined the overall genomic similarity between \textit{K. quasipneumoniae} M17277 and the type strains 01A030\textsuperscript{T} and 07A044 with a genome-to-genome distance calculation. The results confirmed same species between M17277 and 01A030\textsuperscript{T} (0.95\% G+C) and distinct species between M17277 and 07A044 (1.19\% G+C) (see S4 Table). We determined the overall genomic similarity by calculating the genome-to-genome distance between and the type strains 01A030 and 07A044. The results obtained confirmed that M17277 is the same species than the type strain 01A030 with a difference in \% G+C: 0.95 and distinct species compared with 07A044 with a difference in \% G+C: 1.19 (details in S4 Table).

M17277 showed ciprofloxacin resistance (Table 1). However, the analysis of \textit{gyrA}, \textit{gyrB}, \textit{parC} and \textit{parE}, compared to the corresponding figures of \textit{K. quasipneumoniae} subsp. \textit{quasipneumoniae} \textsuperscript{01A030}, did not show quinolone resistance mutations. Interestingly, M17277 harboured three plasmid-mediated quinolone resistance genes. The \textit{qnrB} alleles \textit{qnrB4} and \textit{qnrB52}, were found in contigs of 14,325 and 2,947 nucleotides, respectively, that showed maximal identity with the plasmids pYNKP001\textsuperscript{-}dftrA (99.98\% identity) of \textit{Raoultella ornitholytica} YNK001 and pQnrB52\textsuperscript{-}020046 (100\% identity) of \textit{K. pneumoniae} SCKP020046, respectively (acc. no. KY270853 and CP028782, respectively). Both \textit{qnrB} alleles were located in the variable region 2 of complex class 1 integrons (associated with ISCR1), showing the typical immediate genetic environment of \textit{qnrB} alleles, i.e., bordered by the genes \textit{sapA} and \textit{pspF} [24].
In addition, aac(6')-Ib-cr1 was found in a 3,542-nucleotide contig that encompassed the cas-sette region of complex class-1 integron [aac(6')-Ib-cr1, arr-3, dfrA27 and aadA16], with 100% identity with 26 plasmids from nine enterobacterial species, including K. quasipneumoniae.

Discussion

Here, we report the case of a paediatric patient with erythrodermic psoriasis hospitalised due to edematous-ascitic syndrome. The patient was treated with multiple antibiotics including carbapenems and colistin. Notwithstanding the treatments provided, he was infected and colonised with multiple microorganisms including five NDM-1-producing Enterobacteriaceae, one co-producing MCR-1. Unfortunately, during 2014, parenteral fosfomycin (all five strains were susceptible to fosfomycin) was not available in that hospital.

The isolate M17277, originally typified as K. pneumoniae, was found to be K. quasipneumoniae subsp. quasipneumoniae. This species was first proposed in 2014 [13] and, therefore, the current epidemiological information is still scarce [14]. K. pneumoniae was divided through phylogenetic analysis into three species named K. pneumoniae, Klebsiella quasipneumoniae and K. variicola [13, 14]. In turn, K. quasipneumoniae was subdivided into two subspecies, namely, K. quasipneumoniae subsp. quasipneumoniae and K. quasipneumoniae subsp. similipneumoniae (phylogroups KpIIA and KpIIB, respectively) [13, 14]. In 2017, Aires et al. reported the first observation of an NDM-1-producing K. quasipneumoniae, isolated from a
rectal swab in Brazil [22]. M17277 was also isolated from a rectal swab and becomes the first report on NDM-1-producing *K. quasipneumoniae* subsp. *quasipneumoniae* from Argentina. Besides *bla*_{NDM-1}, this isolate also harboured *qnrB4* and *qnrB52* that are epidemiologically relevant since they were not previously reported in Argentina [25]. The presence of these genes, in addition to *aac(6')-Ib-cr1*, explains, at least in part, the high ciprofloxacin resistance level observed in M17277 (Table 1).

Up to date, there are four reports that involve seven cases of co-carriage of unrelated bacteria with fully sequenced *bla*_{NDM} plasmids [4, 26–28]. In two of these cases, *bla*_{NDM} was located in different plasmids associated with different mobile genetic elements [4, 26]. In other two cases, *bla*_{NDM} was harboured by different plasmids but within the same mobile genetic element [4, 27]. Finally, in the remaining three cases, *bla*_{NDM} was located in identical plasmids (IncC or IncN groups) [4, 28], a situation similar to that reported here. A/C plasmids are now considered IncA and IncC groups [21]. These were the earliest broad host range plasmids to be associated with antibiotic resistance [29]. IncC was divided into type 1 and type 2 plasmids defined by the backbone and resistance island features [29]. Most IncC type 1 plasmids are known to disseminate *bla*_{NDM} within an antibiotic resistance island known as ARI-A [29, 30].

Plasmids highly identical to those found herein were only reported in Australia [19] and USA [31]. Interestingly, the Argentinian plasmids harboured IS{Kpn}8, an insertion sequence that up to date has only been associated to *bla*_{KPC}, environment in Argentina and Asia [20]. Moreover, we recently characterised a *C. amalonaticus*-producing *bla*_{NDM-1} and *mcr-1.5* in plasmids highly similar to those reported here, isolated in 2016 and recovered from the same pediatric hospital than the case reported here [32]. Therefore, our analyses contribute to show that *bla*_{NDM} was and continues to be disseminated through several mechanisms worldwide and across species barriers.

The first three *bla*_{NDM-1} plasmids isolated were identical, therefore, it is likely that *K. quasipneumoniae*-M17277 colonized the patient and passed pKQN17277 to a colonizing *mcr-1*-harbouring *E. coli* strain giving rise to *E. coli*-M17386, which later caused a systemic infection. Similarly, *C. freundii*-M17394 could have acquired pKQN17277 by conjugation from previously colonizing *Enterobacteriaceae*. The SNPs detected in the *bla*_{NDM-1} plasmids of *E. cloacae*-M17464 and *S. marcescens*-M17468 suggest independent acquisitions in these isolates, but more data is necessary to corroborate these assumptions.

Colistin is a last resort drug that, in particular, has been reclassified by the World Health Organization as “Highest Priority Critically Important Antimicrobials” to human medicine, in an attempt to optimize its clinical use to treat serious human infections considering also the emergence of plasmid encoded colistin resistance gene *mcr-1* [33]. In this study, we report an NDM-1-producing *E. coli*-M17386 that was additionally resistant to colistin due to the co-expression of MCR-1 [23]. The co-production of NDM-type and MCR-type enzymes has already been reported in *E. coli* and other *Enterobacteriaceae* species from hospitalized patients [5], healthy and sick food animals like broiler chickens and swine as well as from the farm environment like the slaughterhouse and sewage [34].

**Conclusions**

Health institutions worldwide have to affront the lack of effective antimicrobial treatments against multidrug-resistant bacteria and the spread of promiscuous plasmids that are able to disseminate resistance determinants into infecting or colonising unrelated bacterial species. In this case study, we showed the intra-patient dissemination of a *bla*_{NDM-1} harbouring plasmid among different *Enterobacteriaceae* species including the emerging pathogen *K. quasipneumoniae* subsp. *quasipneumoniae* [25]. As a consequence, active surveillance measures and strict
institutional policies are imperative to determine the local prevalence and prevent further dissemination.

Supporting information

S1 Fig. (a) Estimation of plasmid content and size of clinical isolates and transconjugants. Nuclease S1-PFGE of DNA plugs was performed to estimate plasmid content and size of the studied isolates. The five clinical isolates and transconjugants harboured between 2 to 4 plasmids (48.5–470 kb). (b) S1 nuclease-PFGE and Southern blot with blaNDM-1 probe of blaNDM-1 transconjugants

S1 Table. (a) List of primers used. (b) Salmonella spp. M1744 origin and MIC.

S2 Table. Metadata and accession numbers of the 58 isolates included in the Kleb-dataset selected from the Holt’s database [1]. The phylogroups KpI, KpIIA, KpIIB and KpIII correspond to K. pneumoniae, K. quasipneumoniae subsp. quasipneumoniae, K. quasipneumoniae subsp. similipneumoniae and K. variicola, respectively.

S3 Table. Stats of the pan-genome analysis of the isolate M17277 and the Kleb-dataset. The alignment of concatenated core genes (highlighted in yellow) resulted in a dataset of 61 taxa with 1,702,827 positions. The subsequent selection of the SNP sites from that core-genome alignment resulted in a SNP alignment of 61 taxa with 206,622 positions, which was used to construct the maximum likelihood phylogenetic tree.

S4 Table. Gene to gene distance results.

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