

Colistin resistance in Parisian inpatient faecal *Escherichia coli* as the result of two distinct evolutionary pathways

Anne Sophie Bourrel¹†, Laurent Poirel^{2,3}†, Guilhem Royer^{1,4,5}†, Mélanie Darty¹, Xavier Vuillemin¹, Nicolas Kieffer^{2,3}, Olivier Clermont⁴, Erick Denamur^{4,6}, Patrice Nordmann^{2,3} and Jean-Winoc Decusser^{1,4*}
on behalf of the IAME Resistance Group‡

¹Laboratoire de Bactériologie et d'Hygiène Hospitalière, CHU Henri Mondor, Assistance Publique - Hôpitaux de Paris, Créteil, France ;
²Laboratoire Européen Associé INSERM, Emerging Antibiotic Resistance in Gram-Negative Bacteria, Emerging Antibiotic Resistance Unit, Medical and Molecular Microbiology, Faculty of Science and Medicine, University of Fribourg, Fribourg, Switzerland; ³National Reference Centre for Emerging Antibiotic Resistance (NARA), Fribourg, Switzerland; ⁴IAME, UMR1137 INSERM, Université Paris Diderot, Université Paris Nord, Emerging Antibiotic Resistance in Gram-Negative Bacteria, Paris, France; ⁵LABGeM, Génomique Métabolique, Genoscope, Institut François Jacob, CEA, CNRS, Univ Evry, Université Paris-Saclay, Evry, France; ⁶Laboratoire de Génétique Moléculaire, Hôpital Bichat, Assistance Publique - Hôpitaux de Paris, Paris, France

*Corresponding author. Department of Microbiology, Assistance Publique - Hôpitaux de Paris, University Hospital Henri Mondor, 94000 Créteil, France. Tel: +33 1 49 81 49 36; Fax: +33 1 49 81 28 39; E-mail: jean-winoc.decusser@aphp.fr

†Contributed equally.

‡Other members are listed in the Acknowledgements section.

Background: Beyond plasmid-encoded resistance (*mcr* genes) prevalence in strain collections, large epidemiological studies to estimate the human burden of colistin-resistant *Escherichia coli* gut carriage are lacking.

Objectives: To evaluate the prevalence of colistin-resistant *E. coli* carriage in inpatients and decipher the molecular support of resistance and the genetic background of the strains.

Methods: During a 3 month period in 2017, we prospectively screened patients in six Parisian hospitals for rectal carriage of colistin-resistant *E. coli* using a selective medium, a biochemical confirmatory test and MIC determination. WGS of the resistant strains and their corresponding plasmids was performed.

Results: Among the 1217 screened patients, 153 colistin-resistant *E. coli* strains were isolated from 152 patients (12.5%). The *mcr-1* gene was identified in only seven isolates (4.6%) on different plasmid scaffolds. The genetic background of these MCR-1 producers argued for an animal origin. Conversely, the remaining 146 colistin-resistant *E. coli* exhibited a phylogenetic distribution corresponding to human gut commensal/clinical population structure (B2 and D phylogroup predominance); 72.6% of those isolates harboured convergent mutations in the PmrA and PmrB proteins, constituting a two-component system shown to be associated with colistin resistance.

Conclusions: We showed that the occurrence at a high rate of colistin resistance in human faecal *E. coli* is the result of two distinct evolutionary pathways, i.e. the occurrence of chromosomal mutations in an endogenous *E. coli* population and the rare acquisition of exogenous *mcr-1*-bearing strains probably of animal origin. The involved selective pressures need to be identified in order to develop preventative strategies.

Introduction

Colistin is among the very few last-resort antibiotics in the armamentarium for treating infections caused by MDR Gram-negative bacteria. The emergence and spread of plasmid-encoded colistin resistance among Enterobacteriaceae is therefore of great concern, threatening the clinical relevance of the polymyxin antibiotic family.¹ The occurrence of colistin resistance, and more particularly transferable colistin resistance, in *Escherichia coli* is of paramount

importance. Currently, *E. coli* ranks first as the bacterial species responsible for community and hospital infections worldwide. Beyond the report of infection cases, the extent of the human reservoir for colistin-resistant Enterobacteriaceae remains unclear. Most of the epidemiological data originated from strain collections that were selected either according to their resistance to another antimicrobial class (mainly isolates producing ESBLs or carbapenemases) or according to their clinical origin (e.g. bloodstream isolates).²⁻⁴ Consequently the prevalence of intestinal carriage of acquired

colistin resistance in humans in Europe remains unknown, mostly because of the lack of an adequate screening medium. In the present work we performed a prospective multicentre study using a screening medium to: (i) measure the prevalence of colistin-resistant human faecal *E. coli*; (ii) decipher the molecular mechanisms mediating colistin resistance; and (iii) identify the genetic background of colistin-resistant *E. coli* isolates.

Methods

Study design and material

We performed a prospective multicentre non-interventional study among six University hospitals ('Assistance Publique-Hôpitaux de Paris') geographically distributed in the Paris area [two in Paris *intra muros* and four in the suburbs; see Figure S1 (available as Supplementary data at JAC Online) for details], France, namely the ColiRED study. During a 3 month period in 2017 (January-March), each adult patient screened at hospital admission because of risk factors for carriage of XDR bacteria (such as carbapenemase-producing Enterobacteriaceae) or at an ICU admission for MDR bacteria (i.e. ESBL-producing Enterobacteriaceae) was additionally screened for carriage of colistin-resistant *E. coli* using Superpolymyxin[®] agar plates (ELITECH Microbiology, Signes, France) according to the recommendations of the manufacturer. Regarding the French guidelines, patients at risk for carriage of eXDR bacteria (emerging extensively drug-resistant bacteria, i.e. VRE or carbapenemase-producing Enterobacteriaceae) were direct repatriates as inpatients from an overseas health institution for a hospital stay, patients with a history of hospitalization abroad within the previous year, patients with past medical history of eXDR carriage or patients with a history of contact with eXDR patients.⁵ Only one single rectal swab (Eswab[®], Copan, Italia) per patient was included. In order to prevent the emergence of adaptive resistance, we limited the use of selective medium to the initial screening step; picking and streaking of subsequent colonies were performed on non-selective medium. Basic clinical data such as gender, age, dates of hospital admission and of specimen collection, and history of admission in the preceding year in overseas hospitals were recorded.

Ethics

Informed consent was obtained from each participant; this study was approved by the Ethics Committee - Institutional Review Board from Henri Mondor Hospital (IRB Mondor 20180607 Decousser).

Antimicrobial susceptibility testing

After collection, the swabs were immediately grown on the screening medium. Each different morphotype of colonies was identified at the species level using MALDI-TOF MS (Andromas[®], Beckman Coulter, Villepinte, France). The colistin resistance of *E. coli* isolates was confirmed by using the Rapid Polymyxin NP[®] test (Elitech Microbiology, Signes, France) according to the recommendations of the manufacturer. MICs of colistin were determined for each strain by the broth microdilution method (BMD) according to the EUCAST recommendations (http://www.eucast.org/ast_of_bacteria/mic_determination/?no_cache=1). Antimicrobial susceptibility testing of the other antimicrobial compounds was performed using the disc diffusion method according to the EUCAST recommendations (http://www.eucast.org/ast_of_bacteria/disk_diffusion_methodology/). The list of antimicrobial compounds that were tested is reported in the Supplementary data. Carbapenemase production was tested using the Rapidec Carba NP test[®] (bioMérieux, Craponne, France).

WGS and analysis

The whole genomes of all *E. coli* isolates exhibiting an MIC >2 mg/L were sequenced using the Illumina technology and analysed with an in-house

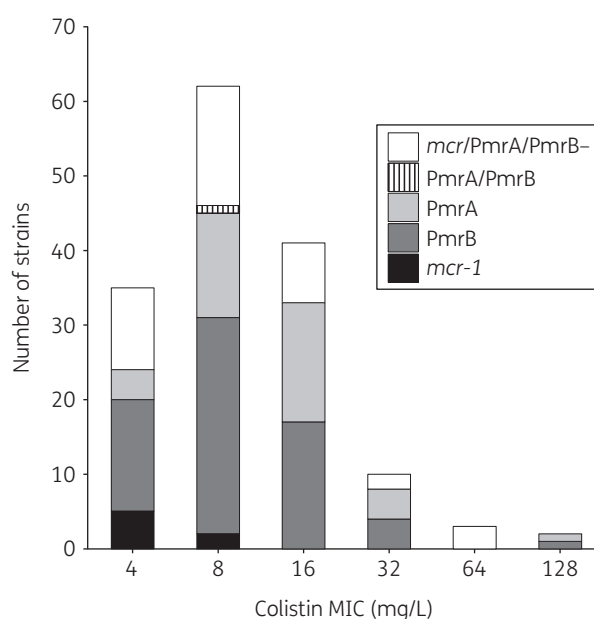


Figure 1. Colistin MIC distribution for the 153 colistin-resistant *E. coli* strains. MICs were determined using the broth microdilution method. The MICs for the seven *mcr-1*-positive colistin-resistant *E. coli* were 4 mg/L ($n=5$) and 8 mg/L ($n=2$) (indicated in black). The MIC distribution of the PmrA and PmrB mutations is indicated in different shades of grey. The strain mutated in both PmrA and PmrB is indicated by the hatched lines. The strain without additional *mcr* gene and no mutation in PmrA or in PmrB is indicated in white.

bioinformatics pipeline (see Table S1 and Table S2 for details). Briefly, after SPAdes assembly, the chromosome sequences were ordered against reference genomes with Ragout to enhance the assembly quality.^{6,7} The phylogroup was determined *in silico* using the ClermonTyping method, and read-based determination of MLST through the Warwick scheme and serotype were performed with SRST2.^{8,9} The resistome, the virulome and the plasmid type were defined by BlastN approach with Abricate (<https://github.com/tseemann/abricate>) using the ResFinder database, a custom database, and PlasmidFinder, respectively.^{10,11} The plasmid sequences of each isolate were also predicted by PlaScope.¹² The contigs carrying *mcr* genes were annotated using Prokka (v1.13.3) to explore the genetic environment.¹³ Phylogenetic history of the strains was reconstructed using SNPs of the core genome. Briefly, SNP calling was performed for each strain against *E. coli* K-12 MG1655 with Snippy (v4.0) (<https://github.com/tseemann/snippy>). Then, only core SNPs were conserved and used to construct a maximum likelihood phylogenetic tree using FastTree (v2.1.9) with the general time reversible evolution model and a gamma distribution of rates across sites.¹⁴ All the genomic data are publicly available through the bio-project PRJEB28020.

Plasmid sequencing and identification of plasmid-mediated colistin resistance determinants

To confirm the data of plasmid sequences obtained with WGS of the natural isolates, we performed conjugation assays for all the *mcr*-positive strains using *E. coli* J53 as the recipient strain, followed by PCR-based replicon typing as described to identify the genetic background of the *mcr*-bearing plasmids.¹⁵ The whole genome of each *E. coli* J53 transconjugant was also sequenced by using the Illumina technology and assembled with SPAdes. Then, *E. coli* J53 transconjugants were aligned against the *E. coli* J53 complete reference sequence using Quast 4.5 with standard parameters.^{16,17}

Table 1. Main characteristics of the MCR-1-producing colistin-resistant *E. coli* isolates

Isolate	Hospital (name)	Hospitalization abroad history ^a	ST Warwick	ST Pasteur	Phylogenetic group	Serotype	Colistin resistance genotype	Incompatibility group	ESBL	Ampicillin resistance genotype	Tetracycline resistance genotype	Sulphonamide/trimethoprim resistance genotype	Phenicol resistance genotype	Intestinal pathogenic <i>E. coli</i> virulence gene	Extra-intestinal <i>E. coli</i> virulence gene
370D	Avicenne	none	7056	869	clade I	Onovel27: H45	<i>mcr-1</i>	IncX4	-	<i>bla</i> _{TEM-1b}	<i>tet</i> (B), <i>tet</i> (M)	<i>sul2</i> , <i>sul3</i> , <i>dfrA12</i>	<i>cmiA1</i>	-	<i>ompT</i>
436A	Lariboisière	none	8154	842	B1	O54:H8	<i>mcr-1</i>	IncX4	-	-	<i>tet</i> (A)	<i>sul2</i> , <i>dfrA17</i>	-	-	<i>iha</i> , <i>ireA</i> , <i>ompT</i>
635A	Avicenne	Thailand	189	809	A	O177:H26	<i>mcr-1.9</i>	IncX4	-	<i>bla</i> _{TEM-1b}	<i>tet</i> (A), <i>tet</i> (M)	<i>sul1</i> , <i>sul3</i> , <i>dfrA12</i>	<i>catA2</i>	<i>eae</i>	<i>traT</i>
925B	Louis Mourrier	none	10	2	A	O89:H10	<i>mcr-1</i>	IncI2	-	-	-	-	-	-	-
933A	Avicenne	none	48	843	A	O6:H16	<i>mcr-1</i>	IncHI2	-	<i>bla</i> _{TEM-1b}	<i>tet</i> (A)	<i>sul2</i> , <i>dfrA14</i>	<i>catA1</i> , <i>flor</i>	-	<i>iha</i> , <i>iucC</i> , <i>iroN</i> , <i>traT</i>
1057A	Henri Mondor	Cambodia	219	870	E	O114:H34	<i>mcr-1</i>	IncHI2	-	<i>bla</i> _{TEM-1b}	<i>tet</i> (A), <i>tet</i> (M)	<i>sul2</i> , <i>sul3</i>	<i>cmiA1</i> , <i>flor</i>	-	<i>iron</i> , <i>traT</i>
1263A	Henri Mondor	South Korea	7329	881	B1	Onovel9: H21	<i>mcr-1</i>	IncI2	-	<i>bla</i> _{TEM-1b}	<i>tet</i> (A)	<i>sul1</i> , <i>sul2</i> , <i>dfrA1</i>	-	-	<i>hra</i> , <i>iron</i> , <i>traT</i> , <i>ompT</i>

^aDuring the previous year.

Unaligned contigs (i.e. plasmid sequences) were analysed with ResFinder and PlasmidFinder.

Prediction of the impact of amino acid substitutions of PmrA and PmrB proteins

Functional effects of missense mutations observed in PmrA and PmrB were predicted with SIFT, PolyPhen-2 and PROVEAN software as proposed in Desroches *et al.*¹⁸ Mutations were considered as impacting the protein when they were predicted by at least two of these as damaging/deleterious. Additionally, we looked for mutations in *pmrA* and *pmrB* genes that modified the length of the protein.¹⁹

Mutations in other genes involved in chromosomal colistin resistance in other bacteria

We looked for mutations in *Klebsiella pneumoniae mgrB* and *ctrB* orthologous genes in *E. coli* strains that have been reported in colistin-resistant *K. pneumoniae*.¹ Regarding *mgrB*, which is also named *yobG*, we tested its presence and putative associated mutation in the corresponding protein using the online available reference sequence (<https://www.uniprot.org/uniprot/P64512>). Regarding *ctrB*, we looked for genes presenting at least 30% similarity in terms of protein counterpart.

Data management and statistical analysis

Clinical data were collected into a Microsoft Excel 2010 database that was password protected. The data management was approved by the Commission Nationale de l'Informatique et des Libertés (#2201069 v 0). The Mann-Whitney test for between-group comparisons with Bonferroni correction was used to compare colistin MICs according to resistance mechanisms. All tests were two-sided and the level of statistical significance was set at 0.05.

Results

Population characteristics

During the study period, 1217 patients originating from the six hospitals were prospectively screened. The median age was 65 years and the male/female ratio was 1.49. The reason for screening was the presence of risk factors for MDR/XDR carriage at ICU admission (1111/1217, 91.3%) or the presence of risk factors for XDR carriage at hospital admission (106/1217, 8.7%).

Prevalence of colistin-resistant strains

Most of the included patients were hospitalized <48h before screening (817/1217, 67.1%). The screening test was positive for 178 patients from which 184 *E. coli* isolates identified by MALDI TOF were recovered. The rapid confirmation test for colistin resistance was positive for 165 patients, with a total of 168 positive strains (Figure S2). The determination of MICs identified 153 *E. coli* strains with an MIC >2 mg/L considered as colistin-resistant *E. coli* from 152 patients (Figure 1 and Figure S2). According to the BMD results, the specificities of the Superpolymyxin[®] screening agar and the Rapid Polymyxin NP[®] test observed in our study were therefore 83.2% and 91.1%, respectively (Figure S2). The 16 isolates for which negative results were obtained with the confirmation test, as well as the 15 isolates with an MIC of colistin ≤2 mg/L were all negative for *mcr* genes (data not shown). Overall, the prevalence of colistin-resistant *E. coli* carriage was 12.5% (152/1217).

Table 2. Phylogenetic group/subgroup distribution of the 146 *mcr*-negative colistin-resistant *E. coli* with their resistance and virulence characteristics

Phylogenetic group/subgroup	Strains, n (%)	PmrA/PmrB mutation ^a , n (%)	ESBL, n (%)	Ampicillin resistance, n (%)	Tetracycline resistance, n (%)	Sulphonamide and/or trimethoprim resistance, n (%)	Phenicol resistance, n (%)	Intestinal pathogenic <i>E. coli</i> virulence gene score, M (min-max)	Extra-intestinal <i>E. coli</i> virulence gene score, M (min-max)
A	17 (11.6)	10 (58.8)	2 (11.8)	9 (52.9)	5 (29.4)	5 (29.4)	2 (11.8)	0.1 (0-1)	2.8 (0-6)
B1	12 (8.2)	10 (83.3)	1 (8.3)	6 (50.0)	4 (33.3)	6 (50.0)	1 (8.3)	0.3 (0-2)	2.6 (0-9)
CC87	3 (25.0)	3 (100.0)	0	3 (100.0)	3 (100.0)	3 (100.0)	1 (33.3)	0.3 (0-1)	7.3 (6-9)
B2	68 (46.6)	48 (70.6)	11 (16.2)	36 (52.9)	20 (29.4)	26 (38.2)	4 (5.9)	0.0 (0-1)	10.4 (3-17)
I ST131	14 (20.6)	12 (85.7)	9 (64.3)	14 (100.0)	9 (64.3)	8 (57.1)	2 (14.3)	0.1 (0-1)	8.8 (5-12)
I non-ST131	3 (4.4)	3 (100.0)	0	2 (66.7)	2 (66.7)	1 (33.3)	0	0.0 (0-0)	7.3 (6-9)
II	17 (25)	9 (52.9)	0	8 (47.1)	2 (11.8)	7 (41.2)	1 (5.9)	0.0 (0-0)	13.2 (7-17)
III	1 (1.5)	1 (100.0)	0	0	0	0	0	0.0 (0-0)	12.0 (12-12)
IV	2 (2.9)	2 (100.0)	0	0	0	0	0	0.0 (0-0)	9.0 (6-12)
V	1 (1.5)	1 (100.0)	0	1 (100.0)	0	1 (100.0)	0	0.0 (0-0)	12 (12-12)
VI	4 (5.9)	3 (75.0)	0	0	0	0	0	0.0 (0-0)	11.8 (11-12)
VII	11 (16.2)	4 (36.4)	0	9 (81.8)	4 (36.4)	5 (45.5)	0	0.1 (0-1)	9.1 (6-13)
IX	7 (10.3)	6 (85.7)	1 (14.3)	1 (14.3)	1 (14.3)	2 (28.6)	1 (14.3)	0.0 (0-0)	11.9 (10-14)
UA	8 (11.8)	7 (87.5)	1 (12.5)	1 (12.5)	2 (25.0)	2 (25.0)	0	0.0 (0-0)	8.6 (3-13)
C	4 (2.7)	4 (100.0)	0	3 (75.0)	4 (100.0)	3 (75.0)	3 (75.0)	0.3 (0-1)	8.3 (7-10)
D	29 (19.9)	24 (82.8)	4 (13.8)	23 (79.3)	12 (41.4)	18 (62.1)	4 (13.8)	0.2 (0-2)	5.9 (0-11)
CGA	20 (69.0)	16 (80.0)	2 (10.0)	17 (85.0)	11 (55.0)	13 (65.0)	2 (10.0)	0.0 (0-0)	7.2 (1-11)
non-CGA	9 (31.0)	8 (88.9)	2 (22.2)	6 (66.7)	1 (11.1)	5 (55.6)	2 (22.2)	0.6 (0-2)	3.1 (0-6)
E	3 (2.1)	3 (100.0)	1 (33.3)	1 (33.3)	0 (0.0)	1 (33.3)	0	0.0 (0-0)	3.3 (1-5)
F	12 (8.2)	7 (58.3)	1 (8.3)	5 (41.7)	2 (16.7)	6 (50.0)	1 (8.3)	0.1 (0-1)	7.3 (4-12)
Clade V	1 (0.7)	0	0	0	0	0	0	0.0 (0-0)	0.0 (0-0)
Total	146 (100.0)	106 (72.6)	20 (13.7)	83 (56.8)	47 (32.2)	65 (44.5)	15 (10.3)	0.1 (0-2)	7.5 (0-17)

^aMissense mutations previously reported to be associated with colistin resistance as well as mutations modifying the length of the protein were considered. For the sake of clarity, strain 130A belonging to ST131 and exhibiting two mutations (one in PmrA and one in PmrB) was scored with a unique mutation.

Prevalence and characteristics of *mcr*-positive strains

Seven out of the 153 colistin-resistant *E. coli* isolates (4.6%) (Figure 1 and Figure S2) recovered from four out of the six participating centres and from seven distinct patients were positive for the *mcr-1* gene. No isolate was found to be positive for other known *mcr* genes (*mcr-2* to *mcr-8*) (Table 1). Although the number of colistin-resistant *mcr*-positive *E. coli* was low ($n=7$), these strains seem to be more frequent in the group of patients screened at hospital admission compared with the group of patients screened at ICU admission (data not shown). Three out of the seven patients were hospitalized abroad during the previous year (Thailand, Cambodia and South Korea), and six out of the seven patients had been hospitalized for <48h. The plasmid support of the *mcr-1* gene was determined using the PlaScope pipeline and confirmed in all except one isolate by *in vitro* and *in silico* analyses of *E. coli* transconjugants. The plasmids carrying the *mcr-1* gene possessed different backbones, including IncX4, IncI2 and IncHI2 (Table 1 and Table S3). Those plasmid types have been reported in association with the *mcr-1* gene in previous studies.¹ The observed *mcr-1* genetic environments were also typical, with the presence of the *pap2* gene downstream, the presence of *parA/pir* and *nikB* genes in the IncX4 and IncI2 plasmids, respectively, and the infrequent

presence of ISAp1 (Figure S3).²⁰ Among the participating patients, 0.6% (7/1217) carried a colistin-resistant *E. coli* that harboured *mcr-1*. With the exception of isolate 925B, all the *mcr-1*-positive *E. coli* belonged to different phylogenetic groups (A, B1, E or *Escherichia* clade I) and STs frequently reported in animals, but rarely in humans (Figure 2).²¹⁻²⁵ A single isolate (635A) exhibited the *eae* gene coding for intimin and can be considered as an atypical enteropathogenic *E. coli*. All the *mcr-1*-positive *E. coli* isolates possessed a very limited number of extra-intestinal virulence genes (Table 1). Furthermore, all *mcr-1*-positive *E. coli* except one (isolate 925B) harboured resistance determinants to antibiotics frequently used in veterinary medicine, i.e. tetracycline, sulphonamide/trimethoprim and phenicols (Table 1). All isolates remained susceptible to expanded-spectrum cephalosporins and carbapenems. No mutation previously associated with colistin resistance was identified in the chromosomal genes (*pmrA*, *pmrB*, *phoP*, *phoQ* and *yobG*; see below) of those MCR producers.^{1,26}

Prevalence and characteristics of colistin-resistant *mcr*-negative strains

Among the 146 colistin-resistant *E. coli* isolates that did not carry any *mcr* gene, missense mutations in the *pmrA* or *pmrB* gene,

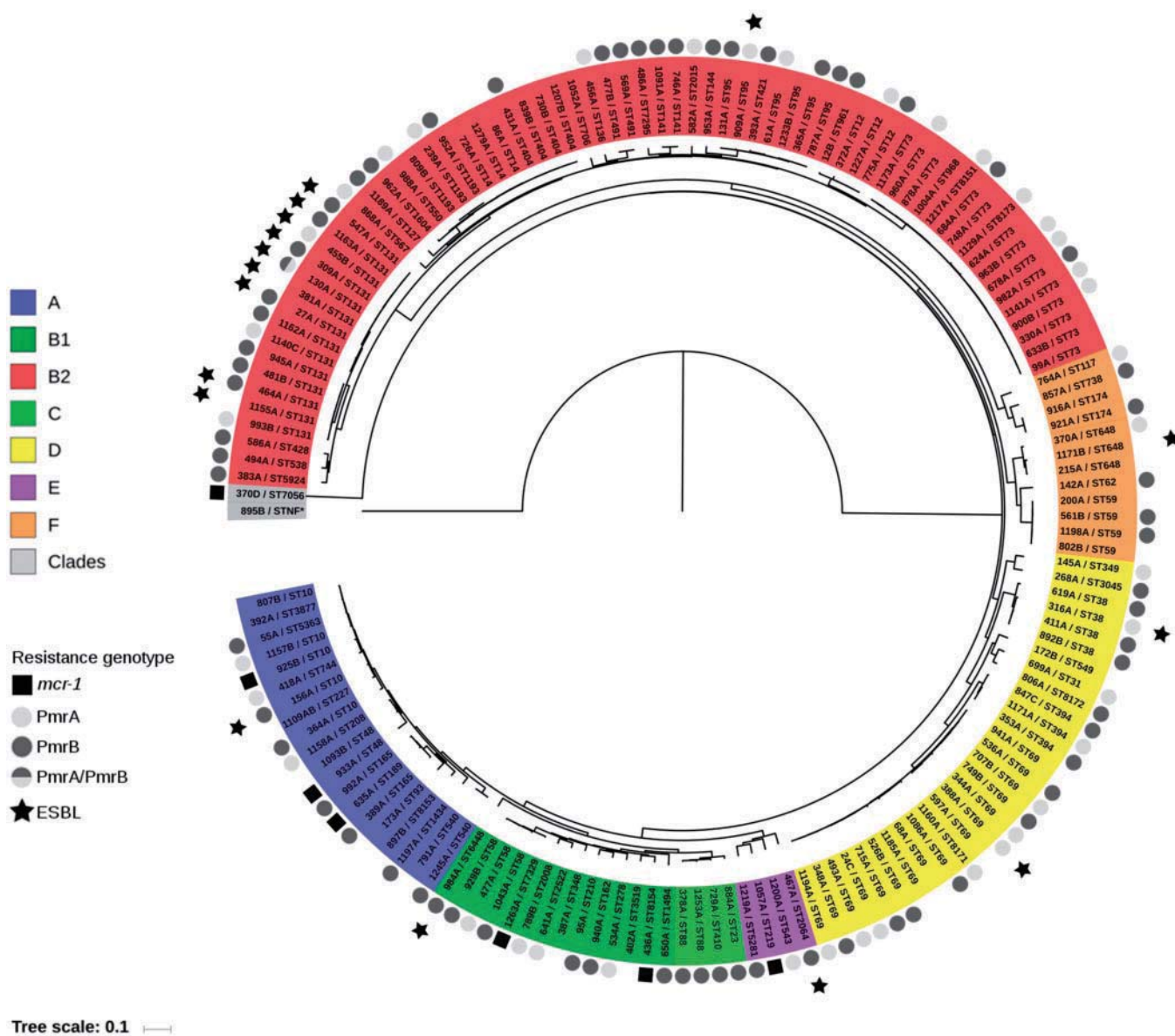


Figure 2. Maximum likelihood tree of the 153 colistin-resistant *E. coli* strains reconstructed from 422324 core SNPs. The tree was rooted on strain 895B (*Escherichia* clade V). The strains are indicated by their ID and ST (Warwick scheme). The main *Escherichia* phylogroups (A, B1, B2, C, D, E and F) and the *Escherichia* clades are represented by colours. The presence of *mcr* genes, PmrA/PmrB mutations (missense or altering the length of the protein) and production of an ESBL are indicated outside the strain ID circle according to the given code. The scale at the bottom represents genetic distances in nucleotide substitutions per site. This figure appears in colour in the online version of JAC and in black and white in the print version of JAC.

coding for a two-component system associated with colistin resistance, were identified in 39 (26.7%) and 55 (37.7%) isolates, respectively (Figure 1).^{27–32} The mutations were mostly localized in two (G53 and R81) and five (L10, C84, P94, E121 and A159) amino acid hot spots in PmrA and PmrB, respectively (Figure 3 and Table S4); these positions were shown to be critical for colistin resistance.¹ All except one of these mutations (strain 806A) were predicted as impacting the protein (Table S4). In addition, we identified one and 12 mutations in *pmrA* and *pmrB* genes, respectively, modifying the length of the protein: four truncating mutations (nonsense $n=2$, frameshift $n=2$), three amino acid

duplications (2–10 amino acids) and six deletions (1–10 amino acids) (Table S5). Of note, the strain 130A (ST131 O25b: H4) exhibiting the amino acid deletion in PmrA also possessed a missense mutation in PmrB (colistin MIC=8 mg/L). Most of the mutations occurred in specific domains of the proteins: the REC domain (residues 1–112) for PmrA and the HAMP (residues 92–114) and HisKA (residues 145–205) domains for PmrB (Figure 3). The strains bearing these mutations belonged to various phylogenetic groups and STs (Table 2 and Figure 2). The frequent occurrence of PmrA/PmrB mutations (missense or modified length) targeting specific domains in distinct clones argues for multiple independent

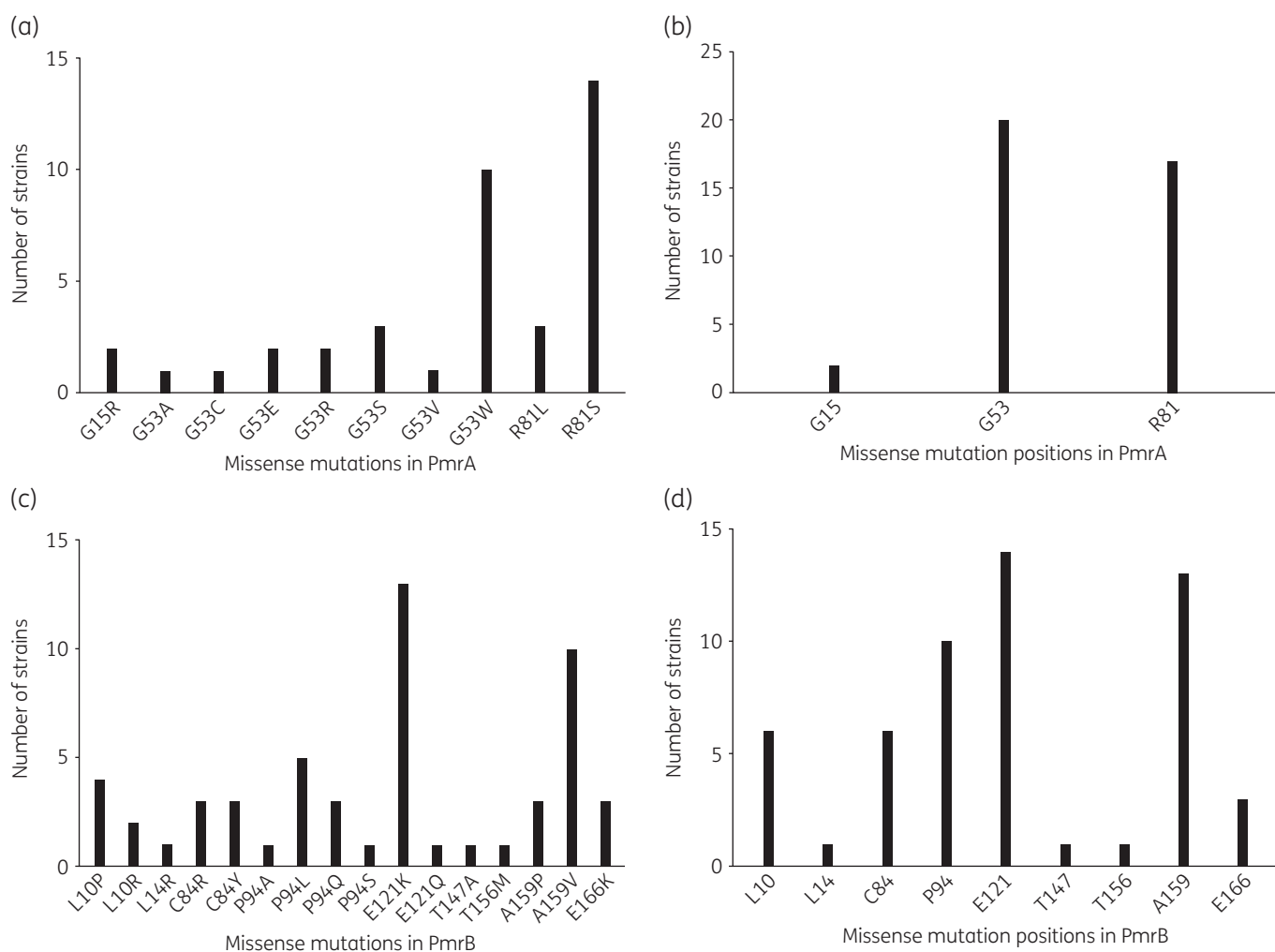


Figure 3. Amino acid sequence missense mutations in PmrA and PmrB with their locations in the 146 colistin-resistant *mcr*-negative *E. coli* strains. (a and b) and (c and d) correspond to PmrA and PmrB proteins, respectively. (a and c) correspond to the different mutations identified, whereas (b and d) represent the location of the mutations in the proteins. The one-letter code for the amino acid designation is used.

mutational events and a convergent evolution. No mutation reported in association with colistin resistance was observed in the PhoP/PhoQ proteins.

Regarding *yobG*, the orthologue of *mgrB* in *E. coli*, we did not identify any mutation or IS insertion that modified the length of the protein. We found only mutations that corresponded to polymorphisms associated with the phylogroup (the V8A mutation in strains from the B2 phylogroup, V8A and V12A mutations in the strain from clade V, and the I41L mutation in the seven strains belonging to STc95). We did not identify a *crrB* orthologue in *E. coli*; this must be a *K. pneumoniae*-specific gene.

The *mcr*-negative isolates mainly belonged to the phylogenetic group B2. Their distribution within the seven main phylogenetic groups actually mirrored that of the overall commensal/clinical isolates previously identified in the Paris area (Table 2 and Figure 2).^{14,33} Those isolates possessed a limited number of intestinal pathogenic genes, but a high rate of extra-intestinal pathogenic genes. According to the resistome analysis, a rate of 11.6% of ESBL carriers was found, and 32.2%, 44.5% and 10.3% of the

isolates showed resistance to tetracycline, sulphonamide/trimethoprim and phenicols, respectively (Table 2). According to the results of the disc diffusion antimicrobial susceptibility testing, the percentage of expanded-spectrum cephalosporin- or carbapenem-resistant strains among *pmrA/B* mutants was 11.3% and 0%, respectively. Of note, comparison of colistin MICs according to resistance mechanisms (i.e. *mcr-1*, PmrA mutation, PmrB mutation, unknown) revealed a significantly lower MIC for *mcr-1*-bearing strains versus PmrA- or PmrB-mutated strains [Mann-Whitney $P=0.00084$ (Bonferroni corrected $P=0.0083$) and Mann-Whitney $P=0.0078$ (Bonferroni corrected $P=0.0083$), respectively]. Regarding the PmrA- and/or PmrB-mutated and PmrA/B non-mutated strains, no difference in terms of phylogenetic groups, MIC₅₀, MIC₉₀ and MIC range was reported (data not shown).

Discussion

The occurrence of colistin-resistant *E. coli* prevalence in humans is a pivotal issue, as the reasons for their selection remain intriguing

in many instances and the extent of their dissemination remains largely unknown. In order possibly to prevent their further selection and spread, their prevalence must be evaluated globally (i.e. the gut carriage) without considering other antibiotic resistance traits, since many of those isolates do not exhibit additional resistance mechanisms. Evaluating the rate of faecal carriage with colistin-resistant Gram-negatives is a challenging process owing to: (i) the interfering effect of intrinsically resistant bacterial species; (ii) some inoculum effect associated with colistin resistance; (iii) the issues of colistin stability and availability in solid agar medium; and (iv) the phenotypic expression of colistin resistance that may be inducible or lead to low-level resistance, especially for *mcr*-associated resistance.¹ To achieve this goal, we used an agar screening medium, corresponding to the commercial product derived from a previously published home-made medium.³⁴ Using such a technical approach, we focused on a well-defined test population.

Surprisingly, we showed that the prevalence of patients colonized by colistin-resistant *E. coli* was very high (12.5%) compared with published studies.³⁵ The contribution of plasmid-mediated MCR-mediated resistance was, however, quite low (4.6%). In 2017, two single-centre studies based on a selective agar culture failed to identify any plasmid-mediated colistin-resistant *E. coli* in France and Switzerland among 653 and 1144 specimens, respectively.^{35,36} When excluding naturally resistant species (such as *Hafnia alvei*, *Proteus* spp., *Morganella* spp., *Providencia* spp., and *Serratia* spp.), the prevalence of colistin-resistant Enterobacteriaceae ranged from 0.8% to 1.5%, with no identification of *mcr-1*-positive strains. A Dutch study focusing on the *mcr-1* prevalence among travellers through a metagenomic approach found only a single MCR-producing colistin-resistant *E. coli* carrier out of 122 healthy humans before travel.³⁷ Here in France, a study focusing on the prevalence of *mcr-1* in patients attending a tertiary care hospital through a real-time PCR approach identified a 0.35% prevalence rate (576 tested patients).³⁸ In China, several studies focused on the carriage of MCR-producing strains. Zhong *et al.*³⁹ reported a 6.2% *mcr-1* prevalence from inpatients and outpatients of three hospitals, mainly corresponding to MDR *E. coli*. This prevalence was 1.6% in another study and rose to 9.8% when hospitalized children were targeted.^{40,41} In Hong Kong, the prevalence was 2.0% in a mixed population of healthy subjects and inpatients.⁴²

In our study, the prevalence of plasmid-mediated colistin-resistant *E. coli* was 0.5%. This prevalence could be underestimated by the use of a selective screening medium and the well-known existence of colistin-susceptible *mcr*-positive strains.¹ Among the seven carriers, three were hospitalized in Asia in the previous year. Although we did not know how many *mcr* non-carriers presented such risk factors, our work therefore could support that travelling to Asia and hospitalization abroad might be considered as risk factors for acquisition of MCR-producing Enterobacteriaceae.⁴³ The genetic supports and environments of the *mcr* genes identified in our study were very similar to those reported in different parts of the world, with three different plasmid types identified belonging to commonly identified incompatibility groups (Table 1 and Figure S3).^{4,20}

Interestingly, for all except one isolate, the genetic backgrounds of those MCR-1-producing colistin-resistant *E. coli* were particular since they corresponded to strains usually identified in animals (Table 1). The *Escherichia* clade I isolate, exhibiting the

– – + – genotype using the Clermont method, actually corresponds to STc2715 (Warwick scheme) and is found in 73 strains of the Enterobase database (<https://enterobase.warwick.ac.uk/species/index/ecoli>), all from animal (mainly livestock) and environmental origins.⁴⁴ Similarly, ST219 (isolate 1057A) and ST189 (isolate 635A) were identified in 14 and 51 strains of the Enterobase database, respectively, almost all originating from poultry and livestock. Isolate 1263A belonged to the ST58/ST155 complex recently reported as being of animal origin and spreading in humans.²⁴ Noticeably, strains belonging to this clonal complex have a high capacity for acquiring and disseminating resistance genes, thus making the identification of the *mcr-1* gene in such a clone even more worrisome.²⁴ Isolate 436A belonged to a very rare B1 phylogroup ST and B1 strains are mostly of animal origin, and isolate 933A belonged to the phylogroup A ST48, frequently associated with animals (383 strains identified in the Enterobase database, mainly from livestock and poultry) and considered as zoonotic.^{21,25} Finally, the phylogroup A ST10 isolate (925B) might be of either human or animal origin.

Altogether, these observations are in line with a recent study including >300 *mcr-1*-positive *E. coli* that showed a predominance of STc10 (including ST48), ST189 and ST155, almost all of animal origin.⁴ In combination with their genetic background, the resistance gene armamentarium of those strains (except 925B) supports their animal origin. Indeed, a high rate of genes encoding resistance to chloramphenicol (*floR*, *catA* and *cml*), tetracycline (*tet* family) and/or sulphonamide/trimethoprim (*sul* and *dfr*) was found among MCR-1 producers in our study (Table 1) with those resistance genes corresponding to antibiotics widely used in veterinary practice. The subsequent acquisition of an *mcr*-bearing plasmid by *E. coli* strains belonging to a highly virulent human clone (i.e. ST131) as reported by Wang *et al.*⁴¹ does not contradict this hypothesis.

Beyond identification of plasmid-mediated colistin resistance, the second relevant finding obtained through the present study was the unexpected high prevalence of *mcr*-negative colistin-resistant *E. coli* carriage (12.0%). As opposed to MCR-1-producing colistin-resistant *E. coli*, the genetic background (phylogenetic group to which it belongs, presence of extra-intestinal genes) as well as the pattern of antibiotic resistance (production of ESBL, low level of resistance to tetracycline, sulphonamide/trimethoprim and phenicols) of the other colistin-resistant *E. coli* actually point to a human origin.^{22,33,45} The identification of missense mutations at specific positions in the PmrA/PmrB two-component system, resulting sometimes from different nucleotide substitutions (Figure 3 and Table S4), as well as mutations modifying the length of PmrB (Table S5), in variable genetic backgrounds (Figure 2), indicates a convergent evolution, a strong hallmark of selection.⁴⁶ Moreover most of the mutations occurred in specific domains of the proteins: the REC domain (residues 1–112) for PmrA and the HAMP (residues 92–141) and HisKA (residues 145–205) domains for PmrB (Figure 3).

Altogether, these findings clearly indicate two evolutionary paths leading to colistin resistance in human faecal *E. coli*, corresponding to a minority of plasmid-encoded MCR-1-producing isolates of animal origin (probably exposed to colistin use) and a vast majority of isolates of human origin, mostly exhibiting resistance through chromosomally encoded mechanisms.¹ The selective pressure for such a high level of non-plasmid-encoded colistin

resistance remains to be identified. In France, considering that colistin therapeutic use is limited to a last resort antimicrobial option in hospitalized patients with a history of XDR carriage and very rarely for treating infections due to *E. coli*, the selective pressure is definitely very low, if present at all. As most of the included patients were hospitalized <48h before screening (817/1217, 67.1%), the colistin consumption in the French community is a relevant factor: the corresponding values were 0.018 DDDs/1000 inhabitants/day in 2016 and 2017 (<https://ecdc.europa.eu/en/anti-microbial-consumption/database/distribution-by-antimicrobial-group>). So, we can consider that the selective pressure due to colistin consumption in the community setting was quite low in France compared with other countries such as the UK (0.104 DDDs/1000 inhabitants/day in 2017). The most probable explanation resides in a coincidental evolution as has been shown in *E. coli* grown at high temperature that became resistant to rifampicin by parallel mutations within *rpoB*, the gene encoding the beta subunit of RNA polymerase, despite never being in contact with rifampicin.⁴⁷ An alternative selective pressure leading to colistin resistance could be the exposure in the gut to non-antibiotic human-targeted drugs. It has been recently reported that such drugs can render *E. coli* resistant to antibiotics by mechanisms involving membrane modifications, among others.⁴⁸ Also, exposure to chlorhexidine has been shown to select mutations conferring colistin resistance in *K. pneumoniae*.⁴⁹

Our study presents several limitations. Firstly, we focused on *E. coli* although other enterobacterial species were reported as acquiring colistin resistance, particularly *K. pneumoniae*. Secondly, we did not perform an exhaustive analysis of the clinical history of the 1217 patients, including previous antimicrobial consumption or short periods of travel outside the country. Our study population, namely patients admitted to a healthcare facility and putatively harbouring some risk factors for MDR bacteria colonization, could have artificially increased the prevalence of colistin-resistant *E. coli*; an additional study focusing on a healthy population would be needed to confirm our findings. Thirdly, the low number of MCR-1-producing colistin-resistant *E. coli* forced us to balance our conclusions on this population of strains, although consistent with a larger previously published study.⁴ Lastly we focused on chromosomal mutations that have been previously proved to be responsible for colistin resistance in *E. coli*, *Salmonella* spp. and *K. pneumoniae* or that modified the length of the protein; the screening, identification and confirmation of the role of new mutations is a relevant, but independent, work. This broad and non-targeted prospective investigation of colistin resistance in *E. coli* constitutes a fully fledged work that is currently underway. Nevertheless, the strength of our work is supported by a robust methodological approach including non-biased multicentre rectal sampling, a marketed culture-based screening method, a reference confirmation test and WGS analysis of all the colistin-resistant isolates.

Conclusions

This is the first multicentre study focusing on the human faecal carriage prevalence of colistin-resistant and *mcr*-positive *E. coli* in Europe. Although the prevalence of *mcr*-positive *E. coli* carriage was low, the high rate of colistin-resistant, but *mcr*-negative, *E. coli* is of concern. The genetic background study of these two

populations of colistin-resistant *E. coli* clearly indicated two distinct evolutionary pathways as the origin of this phenomenon. The selective pressures leading to this pattern of emergence need to be identified to allow the development of preventative strategies.

Acknowledgements

Other members of the IAME Resistance Group

Matthieu LAFAURIE, Béatrice BERCOT (St Louis Hospital), Violaine WALEWSKI, Mathilde LESCAT, Etienne CARBONNELLE (Avicenne-Jean-Verdier Hospitals), Fateh OUSSER, Nadia IDRI, Jean-Damien RICARD, Luce LANDRAUD, (Louis Mourier Hospital), Mathieu LE DORZE, Hervé JACQUIER, Emmanuelle CAMBAU (Lariboisière Hospital), Raphaël LEPEULE and Camille GOMART (Henri-Mondor Hospital).

Funding

This work was partially supported by the Laboratoire Européen Associé INSERM ‘Emerging Antibiotic Resistance in Gram-Negative Bacteria’ and by the Swiss National Science Foundation (project FNS-407240_177381). G. R. was supported by the Assistance Publique - Hôpitaux de Paris [poste d’accueil AP-HP - CEA (Commissariat à l’énergie atomique et aux énergies alternatives) 2016–2018]. E. D. was partially supported by the ‘Fondation pour la Recherche Médicale’ (Equipe FRM 2016, grant number DEQ20161136698).

The funders had no role in the study design, data collection, data analysis, data interpretation or writing of the report.

Transparency declarations

Conflicts of interest: none to declare.

The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Supplementary data

Supplementary data, including Figures S1 to S3 and Tables S1 to S5, are available at JAC Online.

References

- 1 Poirel L, Jayol A, Nordmann P. Polymyxins: antibacterial activity, susceptibility testing, and resistance mechanisms encoded by plasmids or chromosomes. *Clin Microbiol Rev* 2017; **30**: 557–6.
- 2 Jayol A, Poirel L, Dortet L et al. National survey of colistin resistance among carbapenemase-producing Enterobacteriaceae and outbreak caused by colistin-resistant OXA-48-producing *Klebsiella pneumoniae*, France, 2014. *Euro Surveill* 2016; **21**: pii=30339.
- 3 Lupo A, Saras E, Madec JY et al. Emergence of *bla*_{CTX-M-55} associated with *fosA*, *rmtB* and *mcr* gene variants in *Escherichia coli* from various animal species in France. *J Antimicrob Chemother* 2018; **73**: 867–72.
- 4 Matamoros S, van Hattem JM, Arcilla MS et al. Global phylogenetic analysis of *Escherichia coli* and plasmids carrying the *mcr-1* gene indicates bacterial diversity but plasmid restriction. *Sci Rep* 2017; **7**: 15364.
- 5 Lepelletier D, Berthelot P, Lucet JC et al. French recommendations for the prevention of ‘emerging extensively drug-resistant bacteria’ (eXDR) cross-transmission. *J Hosp Infect* 2015; **90**: 186–95.

- 6 Bankevich A, Nurk S, Antipov D *et al.* SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 2012; **19**: 455–77.
- 7 Kolmogorov M, Raney B, Paten B *et al.* Ragout—a reference-assisted assembly tool for bacterial genomes. *Bioinformatics* 2014; **30**: i302–9.
- 8 Beghain J, Bridier-Nahmias A, Le Nagard H *et al.* ClermonTyping: an easy-to-use and accurate *in silico* method for *Escherichia* genus strain phylotyping. *Microb Genom* 2018; **4**: e000192.
- 9 Inouye M, Dashnow H, Raven LA *et al.* SRST2: rapid genomic surveillance for public health and hospital microbiology labs. *Genome Med* 2014; **6**: 90.
- 10 Zankari E, Hasman H, Cosentino S *et al.* Identification of acquired antimicrobial resistance genes. *J Antimicrob Chemother* 2012; **67**: 2640–4.
- 11 Carattoli A, Zankari E, García-Fernández A *et al.* In silico detection and typing of plasmids using PlasmidFinder and plasmid multilocus sequence typing. *Antimicrob Agents Chemother* 2014; **58**: 3895–903.
- 12 Royer G, Decousser JW, Branger C *et al.* PlaScope: a targeted approach to assess the plasmidome of *Escherichia coli* strains. *Microb Genom* 2018; doi: 10.1099/mgen.0.000211.
- 13 Seemann T. Prokka: rapid prokaryotic genome annotation. *Bioinformatics* 2014; **30**: 2068–9.
- 14 Price MN, Dehal PS, Arkin AP. FastTree: computing large minimum evolution trees with profiles instead of a distance matrix. *Mol Biol Evol* 2009; **26**: 1641–50.
- 15 Carattoli A, Bertini A, Villa L *et al.* Identification of plasmids by PCR-based replicon typing. *J Microbiol Methods* 2005; **63**: 219–28.
- 16 Yi H, Cho YJ, Yong D *et al.* Genome sequence of *Escherichia coli* J53, a reference strain for genetic studies. *J Bacteriol* 2012; **194**: 3742–3.
- 17 Gurevich A, Saveliev V, Vyahni N *et al.* QUAST: quality assessment tool for genome assemblies. *Bioinformatics* 2013; **29**: 1072–5.
- 18 Desroches M, Royer G, Roche D *et al.* The odyssey of the ancestral *Escherichia coli* strain through culture collections: an example of allopatric diversification. *mSphere* 2018; **3**: e00553-17.
- 19 Phan MD, Nhu NTK, Achard MES *et al.* Modifications in the *pmrB* gene are the primary mechanism for the development of chromosomally encoded resistance to polymyxins in uropathogenic *Escherichia coli*. *J Antimicrob Chemother* 2017; **72**: 2729–36.
- 20 Wang Q, Sun J, Li J *et al.* Expanding landscapes of the diversified *mcr-1*-bearing plasmid reservoirs. *Microbiome* 2017; **5**: 70.
- 21 Escobar-Páramo P, Le Menach A, Le Gall T *et al.* Identification of forces shaping the commensal *Escherichia coli* genetic structure by comparing animal and human isolates. *Environ Microbiol* 2006; **8**: 1975–84.
- 22 Massot M, Daubié AS, Clermont O *et al.* Phylogenetic, virulence and antibiotic resistance characteristics of commensal strain populations of *Escherichia coli* from community subjects in the Paris area in 2010 and evolution over 30 years. *Microbiology* 2016; **162**: 642–50.
- 23 Mercat M, Clermont O, Massot M *et al.* *Escherichia coli* population structure and antibiotic resistance at a buffalo/cattle interface in Southern Africa. *Appl Environ Microbiol* 2015; **82**: 1459–67.
- 24 Skurnik D, Clermont O, Guillard T *et al.* Emergence of antimicrobial-resistant *Escherichia coli* of animal origin spreading in humans. *Mol Biol Evol* 2016; **33**: 898–914.
- 25 Dissanayake DR, Octavia S, Lan R. Population structure and virulence content of avian pathogenic *Escherichia coli* isolated from outbreaks in Sri Lanka. *Vet Microbiol* 2014; **168**: 403–12.
- 26 Jeannot K, Bolard A, Plésiat P. Resistance to polymyxins in Gram-negative organisms. *Int J Antimicrob Agents* 2017; **49**: 526–35.
- 27 Quesada A, Porrero MC, Téllez S *et al.* Polymorphism of genes encoding PmrAB in colistin-resistant strains of *Escherichia coli* and *Salmonella enterica* isolated from poultry and swine. *J Antimicrob Chemother* 2015; **70**: 71–4.
- 28 Sun S, Negrea A, Rhen M *et al.* Genetic analysis of colistin resistance in *Salmonella enterica* serovar Typhimurium. *Antimicrob Agents Chemother* 2009; **53**: 2298–305.
- 29 Froelich JM, Tran K, Wall D. A *pmrA* constitutive mutant sensitizes *Escherichia coli* to deoxycholic acid. *J Bacteriol* 2006; **188**: 1180–3.
- 30 Nordmann P, Jayol A, Poirel L. Rapid detection of polymyxin resistance in Enterobacteriaceae. *Emerg Infect Dis* 2016; **22**: 1038–43.
- 31 Olaitan AO, Li J. Emergence of polymyxin resistance in Gram-negative bacteria. *Int J Antimicrob Agents* 2016; **48**: 581–2.
- 32 Cannatelli A, Giani T, Aiezza N *et al.* An allelic variant of the PmrB sensor kinase responsible for colistin resistance in an *Escherichia coli* strain of clinical origin. *Sci Rep* 2017; **7**: 5071.
- 33 Clermont O, Couffignal C, Blanco J *et al.* Two levels of specialization in bacteraemic *Escherichia coli* strains revealed by their comparison with commensal strains. *Epidemiol Infect* 2017; **145**: 872–82.
- 34 Nordmann P, Jayol A, Poirel L. A universal culture medium for screening polymyxin-resistant Gram-negative isolates. *J Clin Microbiol* 2016; **54**: 1395–9.
- 35 Saly M, Jayol A, Poirel L *et al.* Prevalence of faecal carriage of colistin-resistant Gram-negative rods in a university hospital in western France, 2016. *J Med Microbiol* 2017; **66**: 842–3.
- 36 Zurfluh K, Stephan R, Widmer A *et al.* Screening for faecal carriage of MCR-producing Enterobacteriaceae in healthy humans and primary care patients. *Antimicrob Resist Infect Control* 2017; **6**: 28.
- 37 von Wintersdorff CJ, Wolffs PF, van Niekerk JM *et al.* Detection of the plasmid-mediated colistin-resistance gene *mcr-1* in faecal metagenomes of Dutch travellers. *J Antimicrob Chemother* 2016; **71**: 3416–9.
- 38 Terveer EM, Nijhuis RHT, Crobach MJT *et al.* Prevalence of colistin resistance gene (*mcr-1*) containing Enterobacteriaceae in feces of patients attending a tertiary care hospital and detection of a *mcr-1* containing, colistin susceptible *E. coli*. *PLoS One* 2017; **12**: e0178598.
- 39 Zhong LL, Phan HTT, Shen C *et al.* High rates of human fecal carriage of *mcr-1*-positive multidrug-resistant Enterobacteriaceae emerge in China in association with successful plasmid families. *Clin Infect Dis* 2017; **66**: 676–85.
- 40 Hu YY, Wang YL, Sun QL *et al.* Colistin resistance gene *mcr-1* in gut flora of children. *Int J Antimicrob Agents* 2017; **50**: 593–7.
- 41 Wang Y, Tian GB, Zhang R *et al.* Prevalence, risk factors, outcomes, and molecular epidemiology of *mcr-1*-positive Enterobacteriaceae in patients and healthy adults from China: an epidemiological and clinical study. *Lancet Infect Dis* 2017; **17**: 390–9.
- 42 Chan WS, Au CH, Ho DN *et al.* Prospective study on human fecal carriage of Enterobacteriaceae possessing *mcr-1* and *mcr-2* genes in a regional hospital in Hong Kong. *BMC Infect Dis* 2018; **18**: 81.
- 43 Pires J, Bernasconi OJ, Hauser C *et al.* Intestinal colonisation with extended-spectrum cephalosporin- and colistin-resistant Enterobacteriaceae in HIV-positive individuals in Switzerland: molecular features and risk factors. *Int J Antimicrob Agents* 2017; **49**: 519–21.
- 44 Clermont O, Christenson JK, Denamur E *et al.* The Clermont *Escherichia coli* phylo-typing method revisited: improvement of specificity and detection of new phylo-groups. *Environ Microbiol Rep* 2013; **5**: 58–65.
- 45 Jolivet S, Vaillant L, Poncin T *et al.* Prevalence of carriage of extended-spectrum β -lactamase-producing enterobacteria and associated factors in a French hospital. *Clin Microbiol Infect* 2018; **24**: 1311–4.
- 46 Tenaillon O, Rodríguez-Verdugo A, Gaut RL *et al.* The molecular diversity of adaptive convergence. *Science* 2012; **335**: 457–61.
- 47 Rodríguez-Verdugo A, Gaut BS, Tenaillon O. Evolution of *Escherichia coli* rifampicin resistance in an antibiotic-free environment during thermal stress. *BMC Evol Biol* 2013; **13**: 50.

- 48** Maier L, Pruteanu M, Kuhn M *et al.* Extensive impact of non-antibiotic drugs on human gut bacteria. *Nature* 2018; **555**: 623–8.
- 49** Wand ME, Bock LJ, Bonney LC *et al.* Mechanisms of increased resistance to chlorhexidine and cross-resistance to colistin following exposure of *Klebsiella pneumoniae* clinical isolates to chlorhexidine. *Antimicrob Agents Chemother* 2016; **61**: e01162-16.
- 50** Ondov BD, Treangen TJ, Melsted P *et al.* Mash: fast genome and metagenome distance estimation using MinHash. *Genome Biol* 2016; **17**: 132.
- 51** Cock PJ, Antao T, Chang JT *et al.* Biopython: freely available Python tools for computational molecular biology and bioinformatics. *Bioinformatics* 2009; **25**: 1422–3.
- 52** Joensen KG, Scheutz F, Lund O *et al.* Real-time whole-genome sequencing for routine typing, surveillance, and outbreak detection of verotoxigenic *Escherichia coli*. *J Clin Microbiol* 2014; **52**: 1501–10.
- 53** Chen L, Zheng D, Liu B *et al.* VFDB 2016: hierarchical and refined dataset for big data analysis—10 years on. *Nucleic Acids Res* 2016; **44**: D694–7.
- 54** Clermont O, Olier M, Hoede C *et al.* Animal and human pathogenic *Escherichia coli* strains share common genetic backgrounds. *Infect Genet Evol* 2011; **11**: 654–62.
- 55** Poirer L, Kieffer N, Brink A *et al.* Genetic features of MCR-1-producing colistin-resistant *Escherichia coli* isolates in South Africa. *Antimicrob Agents Chemother* 2016; **60**: 4394–7.