We investigated unusual carbapenemase-producing *Enterobacter cloacae* complex isolates (n = 8) in the novel sequence type (ST) 873, which caused nosocomial infections in 2 hospitals in France. Whole-genome sequence typing showed the 1-year persistence of the epidemic strain, which harbored a *bla*<sub>VIM-4</sub> ST1-IncHI2 plasmid, in 1 health institution and 2 closely related strains harboring *bla*<sub>CTX-M-15</sub> in the other. These isolates formed a new subgroup in the *E. hormaechei* metacluster, according to their *hsp60* sequences and phylogenomic analysis. The average nucleotide identities, specific biochemical properties, and pangenomic and functional investigations of isolates suggested isolates of a novel species that had acquired genes associated with adhesion and mobility. The emergence of this novel *Enterobacter* phylogenetic lineage within hospitals should be closely monitored because of its ability to persist and spread.

Controlling the dissemination of carbapenemase-producing *Enterobacteriaceae* (CPE) is challenging because carbapenems are among the few antimicrobial drugs that can be used to treat severe infections in this family (1,2). Tzouvelekis et al. calculated the mortality rate of primary bacteremia involving CPEs without active therapy to be 54% (3). Thus, CPEs may carry the threat of a return to the pre–antimicrobial drug era.

The *Enterobacter cloacae* complex (ECC) has become the third most common species among CPEs in France (4). ECCs are not dominated by any single genotype (5), and only certain subspecies/species have previously been associated with infections and nosocomial outbreaks (6–8). The accurate identification of species and subspecies within the ECC is therefore needed to monitor outbreaks and infections.

The identification of species and subspecies within the ECC is challenging, and even more problematic because routine bacterial identification methods based on biochemical tests or matrix-assisted laser desorption/ionization time-of-flight mass spectrometry are yet unable to distinguish between them (9–13). In a seminal work, Hoffmann and Roggenkamp defined 13 genetic clusters (I–XIII) of the ECC on the basis of *hsp60* gene sequences and assigned them to species and subspecies (14). Recently, Chavda et al. extended the number of clusters in the ECC to 18 phylogenomic groups (A–R) by analyzing core single-nucleotide polymorphisms (SNPs) in 390 whole genomes (15).

Using whole-genome sequencing (WGS) approaches, we investigated a cluster of nosocomial carbapenemase-producing ECC isolates collected over a 13-month period in a university hospital in France. The results suggest a double-string diffusion mechanism involving the emergence of both a carbapenemase-encoding plasmid and an ECC phylogenetic lineage not previously described.

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1These authors contributed equally to this article.
Materials and Methods

Clinical and Epidemiologic Survey
The patients were admitted to the Edouard Herriot Hospital in Lyon, France, where a prospective surveillance of CPEs has been implemented since 2012. Case-patients were defined as persons hospitalized after CPE was diagnosed in ≥1 clinical sample during January 12, 2014–December 31, 2015 (online Technical Appendix Table 1, https://wwwnc.cdc.gov/EID/article/24/8/18-0151-Techapp1.pdf). We investigated contact patients and performed an environmental study (online Technical Appendix).

Bacteria Isolation and Phenotypic Characterization
We isolated CPE on a chromogenic medium, chromID CARBA (bioMérieux, Marcy l’Etoile, France). We preincubated environmental CPE specimens for 24 h at 36°C in Trypticase soy broth (TSB; bioMérieux) supplemented with 0.5 mg/L ertapenem. The clinical (n = 7) and environmental (n = 2) isolates were identified by mass spectrometry (VitekMS; bioMérieux). We used the API 50CH system (bioMérieux) for biochemical testing and assessed antimicrobial susceptibility according to the EUCAST guidelines (http://www.eucast.org/). We detected carbapenemase by using the RAPIDEC CARBA-NP test (bioMérieux) (16). As previously described, we determined the sequence of bla<sub>vim-4</sub> by using the Sanger method (17). We performed conjugation experiments at 25°C as previously described (18) and plasmid size determination by pulsed-field gel electrophoresis (19). Biofilm formation and cell adhesion assays are described in the Technical Appendix.

WGS and Genome Assembly
We determined the whole-genome sequences of strains C45 and C309 by using a hybrid de novo assembly of 2 × 150-bp paired-end reads generated by using sequencing technology by Illumina (San Diego, CA, USA) and long reads generated by using Pacific Biosciences technology (Menlo Park, CA, USA). We determined WGS of the other strains by using a de novo assembly of 2 × 150-bp paired-end reads. We performed the assemblies by using SPAdes (20), mapped the reads by using the Burrows-Wheeler aligner (BWA) (21), and polished the assembly by using Pilon (22). The raw data were deposited in EMBL as project PRJEB22398 and the assemblies as LT991954–60. We report further analysis processes in the online Technical Appendix.

Results

Emergence of Carbapenemase-Producing ECC strains
During January 12, 2014–December 31, 2015, a total of 320 positive cultures for ECC with antibiogram were identified in the Edouard Herriot Hospital in Lyon. Each of 7 (2.2%) ECCs recovered from 7 patients (designated P1–7) produced a carbapenemase (Table 1). These isolates were resistant to penicillins and combinations of penicillins, reacting by releasing β-lactamase inhibitors, oxyiminocephalosporins, and ertapenem (Table 2). PCR and sequencing showed the presence of the carbapenemase-encoding gene bla<sub>vim-4</sub> in the 7 isolates. During January 11, 2013–November 30, 2014, no positive cultures for ECC with antibiogram showed bla<sub>vim-4</sub>. These results suggest an epidemic spread of VIN-4–producing ECC strains in the hospital.

Temporal and Spatial Links of the Carbapenemase-Encoding ECC Cases
The attack rate of CPE was 0.7/10,000 hospital stays during January 12, 2014–December 31, 2015, compared with 0.0/10,000 hospital stays during January 11, 2013–November 30, 2014 (p = 0.008; Figure 1). The all-causes crude mortality rate among patients with ECC isolates was 43% (n = 3). We provide additional clinical data in the online Technical Appendix. None of the patients had a recent history of travel or hospitalization in foreign countries. Only 2 patients (P6 and P7) were hospitalized in the same unit at the same time. Five patients (P1, P2, P3, P4, and P7) had undergone surgery in the same operating room but at different dates and with different operating teams. We identified CPE in urine samples of 5 patients (P1–P5) that were drawn from the patients’ urinary catheters. CPE was isolated from a skin sample that we excised from a necrotic ulcer from P6 and in a sample of operative peritoneal fluid from P7. Patient P1 was simply colonized, whereas patients P2–P7 were infected. All patients had ≥1 negative samples from the same clinical site during hospitalization before colonization/infection by ECC (Table 1). These findings showed no clear-cut epidemiologic or temporal links between the VIN-4 ECC case-patients, except that surgical procedures were performed in the same operating room on 5 of the 7 patients.

Environmental Investigations
Putative sources previously described in other settings, such as handwashing sinks (23) and endoscopes (24), were not assumed to be a source of bla<sub>vim-4</sub> ECC because surveillance samples were negative during the study period. In addition to the 102 contact patients, we screened 65 persons during the first 5 episodes (illnesses in P1–P5) without identifying any secondary cases. For the last episode, which involved P6 and P7, we screened 125 of 160 contact patients; all were negative. These findings support a key role for 1 or multiple environmental reservoirs in the nosocomial transmission of bla<sub>vim-4</sub> ECC to patients. However, transmission by healthcare workers (HCWs) cannot be completely ruled out. We did not screen HCWs
Table 1. Case descriptions of VIM-4–producing Enterobacter cloacae complex nosocomial infections in outbreak involving carbapenemase-producing strains, Lyon, France, January 12, 2014–December 31, 2015*  

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>P1</th>
<th>P2</th>
<th>P3</th>
<th>P4</th>
<th>P5</th>
<th>P6</th>
<th>P7</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient no.</td>
<td>67/F</td>
<td>72/M</td>
<td>64/F</td>
<td>69/M</td>
<td>87/M</td>
<td>84/F</td>
<td>82/M</td>
</tr>
<tr>
<td>Hospitalization duration</td>
<td>10 d</td>
<td>Kidney transplant</td>
<td>106 d</td>
<td>Peripheral arterial disease</td>
<td>52 d</td>
<td>Septic shock</td>
<td>26 d</td>
</tr>
<tr>
<td>Purpose of hospitalization</td>
<td>Transplant, medicine</td>
<td>Surgery, medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
</tr>
<tr>
<td>Type of ward</td>
<td>Transplant, medicine</td>
<td>Surgery, medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
<td>Medical, ICU</td>
</tr>
<tr>
<td>Antimicrobial therapy before diagnosis†</td>
<td>VAN, OFL, AMX</td>
<td>CTR, VAN, AMI, MEM,† CTR</td>
<td>CTR, VAN, TZP, AMI, CLI, IMP†</td>
<td>AMX, VAN</td>
<td>AMC</td>
<td>CTR, GEN</td>
<td>CTR, OFL, AMX, PTZ, GEN</td>
</tr>
<tr>
<td>Urinary catheter</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Intubation</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Central venous catheter</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>In-hospital death: delay from admission, d, and etiology</td>
<td>Yes: 106, septic shock from respiratory system</td>
<td>Yes: 52, septic shock, undetermined origin</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Yes: 36, septic shock, peritonitis with hemorrhage</td>
<td></td>
</tr>
<tr>
<td>Last negative sample,‡ delay from admission, d</td>
<td>Urinary; 7</td>
<td>Urinary; 61</td>
<td>Urinary; 1</td>
<td>Urinary; 20</td>
<td>Rectal swab; 2</td>
<td>Necrotic skin; 39</td>
<td>Urinary; 2</td>
</tr>
<tr>
<td>First positive sample</td>
<td>Urinary</td>
<td>Urinary</td>
<td>Urinary</td>
<td>Urinary</td>
<td>Urinary</td>
<td>Necrotic skin</td>
<td>Peritoneal fluid</td>
</tr>
<tr>
<td>Delay from admission, d</td>
<td>32</td>
<td>76</td>
<td>41</td>
<td>29</td>
<td>20</td>
<td>49</td>
<td>28</td>
</tr>
<tr>
<td>Delay from last negative sample, d</td>
<td>25</td>
<td>15</td>
<td>40</td>
<td>9</td>
<td>18</td>
<td>10</td>
<td>26</td>
</tr>
<tr>
<td>ST</td>
<td>ST873</td>
<td>ST873</td>
<td>ST118</td>
<td>ST783</td>
<td>ST118</td>
<td>ST783</td>
<td>ST110</td>
</tr>
<tr>
<td>Isolate identification</td>
<td>C45</td>
<td>C46</td>
<td>C48</td>
<td>C310</td>
<td>E14</td>
<td>E16</td>
<td>C47</td>
</tr>
</tbody>
</table>

*AMI, amikacin; AMX, amoxicillin; AMC, amoxicillin-clavulanate; CLI, clindamycin; CTR, ceftriaxone; CTX, cefotaxime; ETP, ertapenem; GEN, gentamicin; ICU, intensive care unit; IMP, imipenem; MET, metronidazole; MEM, meropenem; OFL, ofloxacin; PTZ, piperacillin/tazobactam; ST, sequence type; VAN, vancomycin.
†Carbapenems.
‡For detection of VIM-4–producing Enterobacter cloacae complex.

for possible CPE carriage, but no particular HCW was involved in care of all CPE cases.

The data, including the temporal distribution of the cases over 13 months and the molecular characterization of the isolates, suggested intermittent transmission of human or environmental origin. We therefore implemented environmental screening in 3 rooms occupied by patients P3 and P7, including the beds, mattress covers, and shared equipment, in June and August 2016. These rooms were investigated because they had been occupied ≥1 time by patients with VIM-4–producing ECC (data not shown). Analysis of the environmental samples after the discharge of patients with VIM-4–producing ECC showed that those collected from a radiator and the mattress cover in 1 patient’s room (P7) were contaminated by VIM-4–producing ECC isolates (E14 and E16). The antibedsore mattresses were used in different rooms for several patients. The incidence of CPE was reduced by discarding the mattress covers after the patients known to be VIM-4–producing ECC carriers were discharged from the hospital (data not shown).

Table 2. Key features of clinical VIM-4–producing Enterobacter cloacae complex isolates in nosocomial outbreak involving carbapenemase-producing strains, Lyon, France, January 12, 2014–December 31, 2015*  

<table>
<thead>
<tr>
<th>Isolate</th>
<th>Species</th>
<th>ST</th>
<th>hsp60 cluster (phylogenome)</th>
<th>Size of assembled genomes, bp</th>
<th>MICs, mg/L</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>ETP</td>
<td>IPM</td>
</tr>
<tr>
<td>C45</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,290,194</td>
<td>2</td>
</tr>
<tr>
<td>C46</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,257,311</td>
<td>2</td>
</tr>
<tr>
<td>C48</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,260,873</td>
<td>2</td>
</tr>
<tr>
<td>C310</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,254,482</td>
<td>2</td>
</tr>
<tr>
<td>E14</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,251,662</td>
<td>2</td>
</tr>
<tr>
<td>E16</td>
<td>E. cloacae complex</td>
<td>873</td>
<td>NA (S)</td>
<td>5,250,845</td>
<td>2</td>
</tr>
<tr>
<td>C47</td>
<td>E. cloacae cluster III</td>
<td>118</td>
<td>III (D)</td>
<td>5,083,854</td>
<td>2</td>
</tr>
<tr>
<td>C308</td>
<td>E. cloacae cluster III</td>
<td>118</td>
<td>III (D)</td>
<td>4,998,377</td>
<td>2</td>
</tr>
<tr>
<td>C309</td>
<td>E. hormaechei steigerwaltii</td>
<td>110</td>
<td>VIII (B)</td>
<td>5,200,769</td>
<td>4</td>
</tr>
</tbody>
</table>

*AAT, aztreonam; CAZ, ceftazidime; CTX, cefotaxime; ETP, ertapenem; FEP, IPM, imipenem; MEM, meropenem; NA, not applicable; ST, sequence type.
These data suggest that the transmission of ECCs by contact with mattress covers could be 1 of the key causative factors, especially for the last episode (P6–P7).

**Multiclonal Spread of the VIM-4 Carbapenemase**

We sequenced the genomes of isolates to assess the molecular links between the VIM-4 ECC cases (Table 2). Six isolates (C45, C46, C48, C310, E14, and E16) formed a clonal cluster designated clone A, in which core genome SNP analysis showed a diversion of <10 SNPs and wgMLST. (Figure 2).

Isolate C309 belonged to ST110, isolates C47 and C308 to ST118, and the 6 isolates of clone A to ST873 (\textit{dnaA}:85/\textit{fusA}:63/\textit{gyrB}:101/\textit{leuS}:103/\textit{pyrG}:96/\textit{rplB}:6/\textit{rpoB}:53). By screening a collection of 30 ECCs isolated in France during the same period, we identified 2 ST873 isolates (CNR1568 and CNR1569) containing the extended-spectrum \(\beta\)-lactamase–encoding gene \textit{bla}_{\text{CTX-M-15}} in the teaching hospital of Caen. These related isolates differed by 28 core genome SNPs and 27 loci (Figure 2). The clustering of the ECC isolates was also apparent in the analysis of the antimicrobial resistance gene contents (Figure 3). Overall, these data show a multiclonal spread of \textit{bla}_{\text{VIM-4}} ECCs, the predominance of ECC genotype ST873 among \textit{bla}_{\text{VIM-4}} ECC, and the interregional spread of this ECC lineage.
VIM-4–Encoding Plasmids

To explore possible links between clone A and the other VIM-4–encoding isolates, we investigated the plasmid contents and the transferability of bla_{VIM-4} by conjugation. The transfer of ertapenem resistance into *Escherichia coli* C600 was successful; all of the 9 isolates were at room temperature. Hybridization of plasmids with a specific probe revealed the location of bla_{VIM-4} on conjugative plasmids of ≈300 kb (n = 6) in the clone A isolates, C47, and C308 and of ≈245 kb (n = 1) in isolate C309. In all isolates, analysis of the assembled genomes showed the presence of an ST1-IncHI2 replicon encoding bla_{VIM-4}.

The plasmid from isolate C45 designated pC45-VIM4 formed a circular 299,117-bp sequence (Figure 4, panel A). The antimicrobial drug resistance genes were located in a ≈50-kb region (bases 97,253–154,784); bla_{VIM-4} gene was included in a 16-kb Tn21-like transposon designated Tn6540 (bases 97,253–113,368). Tn6540 comprised a class 1 integron including bla_{VIM-4} as the first gene cassette, followed by aac(6′)-Ib, dfrA1b, Aant(3′), and smr2. Seven heavy metal resistance loci were also encoded by pC45-VIM4: the tellurite resistance genes terY3Y2XY1W (bases 65,568–69,439) and terZABCDEF (bases 76,028–82,281); the cobalt-zinc-cadmium-resistance protein (bases 118,576 to 119,544); operon copS/copE (bases 165,340 to 167,337); the efflux system rcnR/rcnA (bases 167,595 to 169,105); the arsenic resistance genes arsCBRH (bases 181,666–184,550 bp); and 1 complete mercury resistance operon, merRTCADE (base 134,519–138,533).

We identified a similar organization in the other bla_{VIM-4} isolates (Figure 4, panel B). However, in C309, the VIM-4–encoding plasmid designated pC309-VIM4 (254,277-bp) differed by 2 deletion sites (≈24 kb and ≈21 kb) flanked by mobile elements (IS26 and ISPa21). These deletions resulted in the loss of resistance genes aac(6′)-Ib, tetA, bla_{TEM-1}, ant(3′), and ant(2′)-Ia. These results suggest the horizontal transfer of the same bla_{VIM-4}–encoding plasmid in several lineages of ECC.

Three related IncHI2 plasmids encoding bla_{VIM-1} were identified in GenBank (Figure 4, panel C). Except in the region encoding resistance genes, pC45-VIM4 shared 94% of its sequence and most of gene synteny with pMRVIM0813; pRH-R27 and pRH-R178 reported from Germany are more distantly related (85% and 60% of overlap).

A New Cluster in the Phylogenomic Tree of the ECC

Because specific subgroups within the ECC are more prone to cause nosocomial infections or outbreaks, we characterized the isolates at the species and subspecies levels as described by Hoffmann and Roggenkamp (14) and Chavda et al. (15). In the hsp60–based neighbor-joining tree comprising 52 representative reference and type strains (online Technical Appendix Figure 1), the sequences of C47 and C308 co-localized with hsp60 cluster III and that of C309 localized with hsp60 cluster VIII; both had ≥99.3% identity within the clusters. The hsp60 sequences of the ST873 isolates formed a new cluster that shared only 96.7%–97.1% identity with the closest related sequences.

To confirm that the ST873 isolates formed a new subgroup, we performed a phylogenomic analysis with 398 ECC genomes downloaded from GenBank. In the resulting
phylogenomic tree (Figure 5), the genomes were distributed in 2 major branches corresponding to the *E. hormaechei* and *E. cloacae* metaclusters as previously reported (14,15). The *E. hormaechei* metacluster comprised 6 branches corresponding to Chavda’s phylogenomic groups A–E, and a new phylogenomic group, designated S, comprising solely the ST873 isolates. As expected, the C47, C308, and C309 isolates clustered in Chavda’s phylogenomic groups D and B, which correspond to Hoffmann’s *hsp60* clusters III and VIII (Figure 5). These findings fortify the hypothesis that the ST873 isolates could be a new species or subspecies in the *E. hormaechei* metacluster.

**A New Species in the *E. hormaechei* Metacluster**

Average nucleotide identity (ANI) and percentage of conserved DNA (PCD) can accurately replace DNA–DNA hybridization values for species delineation by using 0.95 and 0.69 as ANI and PCD thresholds, respectively (25,26). We therefore calculated ANIs and PCDs for the ST873 isolates against 398 ECC genomes by using BLAST (http://blast.ncbi.nlm.nih.gov/Blast.cgi). The PCD values were high enough (>0.69) within the *E. hormaechei* and *E. cloacae* metaclusters for the delineation of species and subspecies by ANI calculations (online Technical Appendix Figure 2). Genomes within the same phylogenomic group shared ANI mean values ≥98% (Figure 6). In the *E. cloacae* metacluster, the ANI values supported the designation of the phylogenomic groups as different species (ANI, 0.87–0.94), except for *E. cloacae* subsp. *cloacae* and *E. cloacae* subsp. *dissolvens* (ANI, 0.95). In the *E. hormaechei* group, most phylogenomic groups shared ANI values 0.96–0.98, supporting the split of the *E. hormaechei* metacluster into different subspecies. As expected, isolates C47 and C308 clustered in the *E. hormaechei*
phylogenomic group D and isolate C309 in the phylogenomic group B (also designated *E. hormaechei* subsp. *steigerwaltii*). The new phylogenomic group S, comprising the ST873 isolates, had ANI values below the species cutoff (<0.95) against all groups, including those in the *E. hormaechei* metacluster. We obtained similar results by using the MUMmer-based approach (online Technical Appendix Figures 3, 4), suggesting that the ST-873 isolates are a new species of the *E. hormaechei* metacluster.

The 3 subspecies *E. hormaechei* subsp. *hormaechei*, *E. hormaechei* subsp. *oharae*, and *E. hormaechei* subsp. *steigerwaltii* can be differentiated by using D-adonitol, D-arabitol, D-sorbitol, and D-melibiose fermentation tests (bioMérieux). By using the API 50CH system, we found that the biochemical characterization of our isolates yielded results compatible with *E. hormaechei* subsp. *steigerwaltii* (*hsp60* cluster VIII and phylogenomic group B), as expected (online Technical Appendix Tables 1, 2). Of interest, the ST873 isolates produced a distinguishable biochemical phenotype in the *E. hormaechei* metacluster by growing on only D-melibiose as the sole carbon source (online Technical Appendix Table 2). Overall, our molecular and biochemical data agree with those of previous studies in the field and suggest that the ST873 isolates are a new species of the *E. hormaechei* metacluster.

**Functional Genomics in the *E. hormaechei* Metacluster**

To investigate the functional features of the ST873 isolates and other phylogenomic subgroups in the *E. hormaechei* metacluster, we constructed a pangenome by using 245 strains including our isolates. The pangenome was divided into 3 sections: 1) the core genome (the set of genes shared by 99% of strains), 2) the accessory genome (the set of genes present in some but not all representatives), and 3) the unique genome (genes unique to individual strains). The 245 strains examined yielded a pangenome of 25,221 genes. On the basis of this dataset, the core genome is composed of 2,575 genes, the accessory genome of 14,849 genes, and the unique genome of 7,797 genes (Figure 7, panel A). Functional annotation of the pangenome on the COG database showed an overrepresentation of genes belonging to the groups with housekeeping functions (COG categories C, G, E, F, I, P, H, J, O, D, and T) in the core genome (Figure 7, panel B). The accessory genome and the unique genes had a similar distribution of functional annotations with an overrepresentation of genes involved in DNA recombination (genes encoding integrases, transposases, or resolvases) and defense mechanisms, as well as those belonging to the mobilome (COG categories L, U, V, and X), as expected.

The distribution of variable genome annotations among the phylogenomic groups showed closely related distributions of functional annotations (Figure 7, panel C), except for cell mobility annotations (COG category N, comprising the flagellar- and adhesion-related functions), which were overrepresented in phylogenomic group B and the ST873 isolates (phylogenomic group S), and mobilome annotations (COG category X), which were overrepresented in phylogenomic group A. The differences in functional distribution were enhanced by the analysis of variable genes specific to phylogenomic groups (Figure 7, panel D). Phylogenomic group B (*hsp60* group VIII), the most prevalent subgroup in human infections, and the new epidemic phylogenomic group S (ST873) had an overrepresentation of genes involved in cell motility (COG category N), as previously observed among variable genes, showing that these functions are overrepresented and based on specific genes in...
these ECC lineages. The ST873 group also exhibited enrichment in specific genes related to signal transduction (COG category T), and phylogenomic group A exhibited enrichment in genes related to recombination and the mobilome (COG categories L and X). We also observed other differences in genes related to metabolic functions. Phylogenomic group A demonstrated an enrichment in specific genes involved in lipid metabolism (COG category I), and phylogenomic group D has specific genes involved in coenzyme, nucleotide, and secondary metabolism (COG categories F, H, and Q); phylogenomic group B accumulated specific genes involved in energy production and carbohydrate, amino acid, and ion metabolism (COG categories C, G, E, and P). Overall, these data suggest that there exist distinct lifestyles in ECCs explaining varied abilities to colonize the hospital environment and to induce nosocomial infections.

Adhesion to Abiotic Surface and Epithelial Cells
To investigate the overrepresentation of mobility/adhesion functions in the ST873 isolates in the context of their prolonged persistence in hospital, we compared the ability of our isolates to initiate biofilm formation on an abiotic surface and their adhesion to intestinal epithelial cells. The ST873 isolates had greater ability to initiate biofilm on PVC than did isolates of ST118 and ST110 (online Technical Appendix Figure 5, panel A). We also observed slight differences of adhesion to HT29 intestinal epithelial cells (online Technical Appendix Figure 5, panel B). These results suggest that ST873 isolates have original adhesion features, as suggested by the pan genomic analysis.

Discussion
One major issue regarding CPEs is whether the main driver of the spread of carbapenemases is the transmission of successful clonal lineages or the horizontal transfer of carbapenemase genes by mobile genetic elements such as plasmids. Our study provides evidence for the spread of an epidemic VIM-4–encoding IncHI2 plasmid in distinct lineages of the ECC and the 1-year persistence of an epidemic strain ST-873, suggesting a double-string diffusion mechanism involving the emergence of both a VIM-4–encoding plasmid and a persistent ECC phylogenetic lineage.

ECC accounted for 9.4% and VIM-4–producing ECC for 3.9% of CPEs during 2015 in France (27). Only 16 VIM-4–producing Enterobacteriaceae cases were diagnosed in France during the same period; the 7 ECC cases included in this study, 6 cases of E. coli, and 1 additional case of a VIM-4–producing ECC, in the Paris area, and 2 cases of Citrobacter freundii in northern France. During this period, 6 other cases of CPE were identified in the same hospital: 5 produced oxacillinase 48 carbapenemases, and 1 produced New Delhi metallo-β-lactamase 1; these cases, in line with common observations in France and other countries in Europe, were imported (4). Our investigation was therefore prompted by the contrast between the low incidence of VIM-4–producing ECC cases in France and our case series, which suggested a new carbapenemase-spreading factor.

All the VIM-4 isolates we found harbored a similar blaVIM-4–encoding, self-conjugative plasmid. The blaVIM-4 gene has previously been reported only in IncA/C plasmids in a neighboring country (Italy) (28). The epidemic plasmid we identified belongs to the ST1-IncHI2 incompatibility...
group. IncHII plasmids are frequent among the ECCs and are often associated with the dissemination of genes encoding extended-spectrum β-lactamases and, at least to some extent, bla\textsubscript{VIM-1} (29). Although bla\textsubscript{VIM-4} differs from bla\textsubscript{VIM-1} by a point mutation, the IncHII bla\textsubscript{VIM-4} plasmids we identified substantially diverge from previously reported bla\textsubscript{VIM-1} plasmids by the ≈50kb region encoding the associated resistance genes.

The VIM-4 epidemic strain ST873 persisted in the hospital for ≈1 year despite the application of specific isolation precautions for patients colonized or infected. During the investigation of another nosocomial outbreak that occurred in 2016 in Caen University Hospital (692 km from Lyon), we identified 2 CTX-M-15–encoding ECC strains belonging to ST873, which suggests that the spread of ST873 is not geographically limited. These strains had traits of speciation and specific genes related to signal transduction, cell motility, and adhesion. These functions have a crucial role in the initiation of biofilm formation (30), which was enhanced in the ST873 isolates compared with the other VIM-4 isolates in our study. Biofilm formation, a key function for host–pathogen interactions and environmental survival, may explain the successful persistence of ST873 isolates in the hospital settings of this study.

Nosocomial infections mediated by Enterobacteriaceae can be transmitted to patients in medical settings by HCWs, patient-to-patient spread, or environmental sources (31). Our findings emphasize the need to consider clinical circumstances such as bed contamination, as previously observed (32). In addition, the VIM-4–encoding plasmid pECC-VIM4 transferred to a bacterial recipient at room temperature, which could explain why successful horizontal transfer into ECC multiclonal isolates occurred in the hospital environment and provided support for the crucial importance of environmental reservoirs in the transmission of nosocomial pathogens. Hence, the threat of outbreaks can be limited by high-quality cleaning and disinfection of patient-care areas and the regular replacement of equipment such as mattress covers.

In conclusion, we report a nosocomial outbreak of multiclonal VIM-4–producing ECC that originated from contamination in the hospital environment. The
predominant clone belongs to a new lineage in the ECC and should be closely monitored in the context of nosocomial infections caused by its apparent ability to maintain and spread in a hospital setting. Our study also delineated the multifactorial spread of the VIM-4 carbapenemase and emphasizes the usefulness of ECC phylogenomic typing in the investigation of outbreaks.

Acknowledgments
We thank Laurent Argaud, Thomas Rimmle, Jacques Ninet, Laurent Juillard, Michel Chuzaville, Olivier Monnave, Patrick Lemrusiaux, Lionel Badet, Bernard Flocard, and their staff for patients’ care and Laurent Guillouard and Alexis Pontviande for technical assistance.

This work was supported by INSERM, INRA, the Université Clermont Auvergne, and the Centre Hospitalier Universitaire de Clermont-Ferrand.

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References


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Novel Enterobacter Lineage as Leading Cause of Nosocomial Outbreak Involving Carbapenemase-Producing Strains

Technical Appendix

Material and Methods

Contact Patients and Environment Study

In accordance with French recommendations (http://www.hcsp.fr), contact patients were defined as patients hospitalized in the same unit as a carbapenemase-producing Enterobacteriaceae (CPE) carrier and/or managed by the same medical or nursing staff. These patients were screened weekly for CPE by rectal swab. All CPE carriers were placed on strict contact precautions with dedicated staff as soon as they were identified. In addition, three rooms for CPE patients including the beds, mattress covers, and shared equipment were sampled after cleaning, either with swabs (Copan, Brescia, Italy) or with sterile wipes (bioMérieux, Marcy l’Etoile, France). Each epidemiologic investigation included on-site visits and the collection of patient demographic characteristics, hospitalization details, and clinical and bacteriological data. The attack rate over the 13-month period was calculated for 10,000 hospital stays and compared with that of the previous 13-month period (11/01/2013 to 11/31/2014) using Fisher exact test. p < 0.05 was considered significant.

wgMLST and Core Genome SNP-based Typing

The wgMLST approach combined the analysis of core genome loci and the presence/absence of accessory genes using BWA read alignments as previously described (1). Core and accessory genes were extracted from 60 genomes belonging to the hormaechei metacluster, yielding 2315 core genome loci (2,115,786 bases) and 2091 accessory genes (2,109,600 bases). SNP-based typing was based on the best practices guide produced by the Broad Institute (2). After quality trimming with sickle, the reads were aligned against the core
genome with BWA-mem. PCR duplicate removal and read realignment around indels were then performed with Picard tools (http://broadinstitute.github.io/picard/). Following this alignment processing, variant calling and variant filtering were performed with VarScan (3). As recombination events mask the true phylogenetic signal by generating tight clusters of SNPs, SNPs were filtered out in each mapping if they were called within the vicinity of 25 bp of another SNP, as previously performed (4). Core genomes with all variants instantiated were then generated from filtered SNPs to obtain the phylogenetic tree using RAxML (5) with 100 bootstrap replicates.

**Gene Detection from NGS Data**

Multilocus sequence typing (MLST) and plasmid double locus sequence typing (pDLST) were performed from online resources (http://pubmlst.org) with the ARIBA package (6). Replicons were detected by PlasmidFinder (http://www.genomicepidemiology.org/). The antibiotic resistance genes were assembled with the ARIBA package by mapping short reads against a manually curated and updated database of resistance-associated genes (2835 genes and 1308 SNPs from 130 genes) derived from CARD and Resfinder (7,8). The assembled genes were filtered and identified by alignment against the database with Exonerate (9) using a 95% minimum threshold for coverage and identity percentages. The distance presence/absence matrix derived from resistance gene detection and the corresponding UPGMA tree were computed in R (https://cran.r-project.org/) with ade4 (https://cran.r-project.org/web/packages/ade4) and fastcluster (https://cran.r-project.org/web/packages/fastcluster) packages.

**SNP and hsp60 Enterobacter cloacae Complex Trees**

The 85,257 SNPs were identified by Parsnp (10) using the recombination filtering option. These SNPs were used to infer an approximately-maximum-likelihood tree with FastTree (11). The reliability of the nodes was assessed by the Shimodaira-Hasegawa test. The phylogenetic tree was visualized with FigTree (http://tree.bio.ed.ac.uk/software/figtree). hsp60 typing was performed as previously described (12).

**ANI and PCD Calculation**

The average nucleotide identity (ANI) and the percentage of conserved DNA (PCD) were calculated as previously described (13) with the pyani module (https://github.com/widdowquinn/pyani).
Pangenome Construction

To determine the pangenome of the hormaechei metacluster, 244 genomes were annotated by Prokka (14) and the corresponding dataset was analyzed with the Roary package (15). The homologous genes were functionally annotated with the Clusters of Orthologous Groups (COG) database (16) using EggNOG (17). Functional enrichment analysis was performed with the stats package (https://github.com/SurajGupta/r-source/tree/master/src/library/stats/R) in R to calculate odds ratios and Fisher exact tests with p-values corrected for multiple testing by the Bonferroni adjustment. A p value less than 0.05 and an odds ratio higher than 1.5 were considered statistically significant.

List of ECC Genomes Included in this Study

The Genbank accession numbers (hsp60 cluster, phylogenomic group): AFHR01 (E,VII), AMGJ01 (B,VIII), ANIA01 (B,VIII), ANIC01 (B,VIII), ANID01 (B,VIII), AXLJ01 (D,III), AXLK01 (A,VI), AYIE01 (B,VIII), AYIG01 (D,III), AYIK01 (B,VIII), AYIM01 (D,III), AYIR01 (A,VI), AYIY01 (A,VI), AYJE01 (D,III), CP008823 (B,VIII), CP008897 (D,III), CP008905 (D,III), CP009854 (D,III), CP010377 (E,VII), CP010384 (A,VI), CP011572 (B,VIII), CP011581 (B,VIII), CP011584 (B,III), CP012165 (A,VI), CP012167 (B,III), CP017179 (B,III), CP017180 (C,VI), CP017183 (A,VI), CP017186 (D,III), FKHB00 (D,III), FKHE00 (D,III), FKHL00 (D,III), FKHW00 (D,III), FP929040 (B,VIII), JCKK01 (D,III), JCKR01 (A,VI), JCKS01 (A,VI), JCKT01 (B,III), JCKX01 (B,III), JCLC01 (B,III), JCLD01 (B,III), JCLF01 (A,VI), JCLG01 (D,III), JCLH01 (A,VI), JCLJ01 (B,III), JCLK01 (B,III), JCLL01 (D,III), JCLM01 (D,III), JCLN01 (D,III), JCL001 (B,III), JCLP01 (D,III), JCLQ01 (A,VI), JCLR01 (B,III), JCLS01 (D,III), JCLT01 (D,III), JCLU01 (A,VI), JCLV01 (D,III), JCLW01 (B,III), JJNL01 (D,III), JMUP01 (A,VI), JMUT01 (B,III), JQGO01 (B,III), JRFQ01 (B,III), JSBO01 (A,VI), JTBZ02 (A,VI), JTC02 (A,VI), JTCB02 (A,VI), JTCP01 (A,VI), JTD02 (A,VI), JTCE02 (A,VI), JTC02 (A,VI), JTCG02 (A,VI), JI02 (A,VI), JIPE01 (A,VI), JUI01 (A,VI), JUH01 (A,VI), JUH01 (A,VI), JUHY01 (B,III), JUIA01 (A,VI), JUIB01 (A,VI), JUIC01 (A,VI), JUID01 (A,VI), JUIE01 (A,VI), JULZ01 (B,III), JUN01 (C,VI), JUX01 (B,III), JVBS01 (B,III), JVGX01 (B,III), JVKL01 (B,III), JVN01 (D,III), JVOV01 (B,III), JYD01 (B,III), JYAZ01 (C,VI), JVZ01 (B,III), JWBJ01 (A,VI), JWRO01 (A,VI), JWR01 (A,VI), JYGA02 (A,VI), JYLQ01 (A,VI), JYL01 (A,VI), JYL01 (A,VI), JYLW01 (A,VI), JYLY01 (A,VI), JYLZ01 (A,VI), JYMA01 (A,VI),
Biofilm Formation and Epithelial Cell Adhesion Assays

The initiation of biofilm formation was assayed by the ability of cells to adhere to the wells of 96-well microtiter dishes made of polyvinylchloride plastic as previously described (18). Biofilm formation was detected after 3h of incubation at room temperature by determining the extent of crystal violet-stained cells attached to a surface at 595 nm. The commensal E. coli K-12 strain TG1 carrying a F-conjugative plasmid that promotes biofilm formation (19) was used as a positive control strain for biofilm formation.
HT-29 intestinal epithelial cells were purchased from ATCC and maintained in an atmosphere containing 5% CO\textsubscript{2} at 37°C in Dulbecco’s modified Eagle’s medium. They were seeded at a density of 2×10\textsuperscript{5} cells/cm\textsuperscript{2} in culture plates (Falcon) for 48 H. Cells were infected at a multiplicity of infection of 10 bacteria per cell. Infected cells were centrifuged at 900 g for 10 min at 25°C, maintained at 37°C for 3 hours and then washed three times in phosphate-buffered saline (PBS; pH 7.2). The epithelial cells were then lysed with 1% Triton X-100 (Sigma) in deionized water. Samples were diluted and plated onto Luria-Bertani (LB) agar plates to determine the number of CFU corresponding to the total number of cell-associated bacteria. The adherent-invasive \textit{E. coli} strain LF82 (20) was used as positive control. The adhesion index was expressed as the mean number of associated bacteria per epithelial cell.

The data were compared by the Kruskal-Wallis test with Dunn's post hoc tests. The p-values <0.05 were rated as significant.

Results

Clinical Data

The index case (patient 1, P1) was a 67-year-old woman hospitalized for a kidney transplant. Following surgery, a CPE designated C45 was isolated from a urine sample. However, P1 did not develop an infection and was only colonized. Five months after she was discharged, a second CPE strain designated C46 was isolated from the urine of a 72-year-old man (patient 2, P2) admitted to the intensive care unit (ICU) for endarterectomy and tracheotomy. During his 106 days of hospitalization, the patient developed septic shock. Patient 3 (P3) was a 52-year-old woman admitted to the ICU for an undetermined recurrent septic shock following bowel resection and laparotomy. Two weeks after the isolation of strain C46 from P2, a CPE strain designated C47 was identified from the urine of P3. Patient 4 (P4) was a 69-year-old man hospitalized for a kidney transplant. Two weeks after the isolation of C47 from P3, a CPE strain designated C48 was isolated from the urine of P4. Patient 5 (P5) was an 87-year-old man admitted to the medical unit and then to the ICU for a consciousness disorder. A CPE strain designated C308 was isolated from a urine sample of P5 3 months after the isolation of strain C48. Two months after the isolation of C308, CPE strain C310 was isolated from a skin sample taken from an 84-year-old woman admitted to the medical unit for the treatment of a necrotic...
ulcer (patient 6, P6). Patient 7 (P7) was an 82-year-old man admitted to the surgical unit to undergo lithotrity. One month after the isolation of C310, CPE strain C309 was isolated from the operative peritoneal fluid of P7.

References


### Technical Appendix Table 1. Fermentation tests of *Enterobacter cloacae* complex isolates performed by using API 50CH

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<th>Isolates</th>
<th>3-hydroxybutyrate</th>
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### Technical Appendix Table 2. Biochemical differentiation of the isolates ST-873 among *Enterobacter hormaechei* subspecies and other relevant species of the *E. cloacae* complex

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* +, absence of fermentation; -, fermentation; V, variable among isolates.*
Technical Appendix Figure 1: Neighbor-joining tree based on the hsp60 gene of representative Enterobacter cloacae complex strains including type and reference strains. The isolates C45, C46, C48, C310, E14, E16, CNR1568 and CNR1569 form a new hsp60 genetic cluster. The scale gives the Jukes-Cantor distance along the branches. The leaves are labeled by species name, hsp60 genetic group and the corresponding GenBank accession number.
Technical Appendix Figure 2: Percentages of conserved DNA (PCD) calculated from Blast software (http://blast.ncbi.nlm.nih.gov/Blast.cgi) pairwise comparisons of ST-873 genomes and 401 representative genomes of *E. cloacae complex*. (NA, non-attributed)

Technical Appendix Figure 3: Percentages of conserved DNA (PCD) calculated from MUMer (22) pairwise comparisons of ST-873 genomes and 401 genomes of *E. cloacae complex*. (NA, non-attributed)
**Technical Appendix Figure 4:** Average nucleotide identities (ANI) calculated from MUMer (22) pairwise comparisons of ST-873 genomes and 401 genomes of *E. cloacae complex*. (NA, non-attributed)

| Meta-Cluster | Name                      | hgap60 cluster | Genomic cluster | A | B | C | D | E | F | G | H | I | J | K | L | M | N | O | P | Q | R | NA |
| E. hormaechi | E. hormaechi subsp. stleyeae | VI             | A (n=10)        | 0.19 | | | | | | | | | | | | | | | | | | | |
|              | E. hormaechi subsp. stleyeae | VIII          | B (n=39)        | 0.17 | 0.30 | | | | | | | | | | | | | | | | | | |
|              | E. hormaechi subsp. stleyeae | VIII          | C (n=42)        | 0.17 | 0.30 | 1.00 | | | | | | | | | | | | | | | | | |
|              | E. hormaechi              | III            | D (n=38)        | 0.16 | 0.36 | 0.96 | 0.99 | | | | | | | | | | | | | | | | | |
|              | E. hormaechi subsp. hormaechi | VII          | E (n=48)        | 0.15 | 0.35 | 0.95 | 0.96 | 1.00 | | | | | | | | | | | | | | | | |
| ST-873       | NA                        | S (n=48)       | 0.34 | 0.54 | 0.94 | 0.94 | 0.91 | 1.00 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | I (n=13)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | J (n=21)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | K (n=6)        | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | L (n=16)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | M (n=30)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | N (n=3)        | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | O (n=1)        | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | P (n=4)        | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | Q (n=34)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | R (n=14)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |
| E. coli      | NA                        | S (n=10)       | 0.19 | 0.30 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | | | | | | | | | | | | | | | |

**Technical Appendix Figure 5:** Ability of A) ECC isolates to initiate biofilm formation on PCV and B) to adhere to HT29 intestinal epithelial cells. *E. coli* K12 TG1 was used as positive control for biofilm formation and the *E. coli* reference strain LF82 for adhesion to intestine epithelial cells (Kruskal-Wallis Test; ns, not significant; *, p<0.001; **, p<0.05).