mcr-1–Positive Colistin-Resistant Escherichia coli in Traveler Returning to Canada from China

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DOI: http://dx.doi.org/10.3201/eid2209.160177

To the Editor: A 61-year-old man underwent transurethral prostate resection in Vancouver, British Columbia, in January 2016. On postoperative day 1, he was febrile (39.1°C) and had leukocytosis (12.7 × 10⁹ cells/L). Blood and urine cultures were ordered on postoperative day 2, and ceftriaxone was started. On postoperative day 3, urine culture grew Escherichia coli (>100 million CFU/L). Susceptibility testing (VITEK2, bioMérieux, Quebec, Canada) indicated a possible extended-spectrum β-lactamase producer and showed resistance to ampicillin, cefazolin, ceftriaxone, gentamicin, ciprofloxacin, and trimethoprim/sulfamethoxazole; intermediate resistance to tobramycin; and susceptibility to amoxicillin/clavulanate, piperacillin/tazobactam, ertapenem, meropenem, and nitrofurantoin. Treatment was switched to amoxicillin/clavulanate. The urinary catheter was removed 48 hours later. The patient was discharged on postoperative day 5 and completed 14 days of oral amoxicillin/clavulanate. Blood cultures were negative after 7 days’ incubation.

The E. coli cultured from the patient underwent further testing and grew in equal amounts on Columbia Colistin-Nalidixic acid Agar (CNA) with 5% sheep blood and Columbia agar with 5% sheep blood (OXOID, Ontario, Canada). This result was brought to the attention of the hospital’s medical microbiologist. A colistin Etest (bioMérieux, Quebec, Canada) showed a MIC of 3 µg/mL; EUCAST defines colistin resistance as >2 µg/mL for Enterobacteriaceae (I).

A real-time PCR to detect the mobile colistin resistance (mcr-1) gene was developed at the Provincial Public Health Laboratory by using primers MCR-1F (5’-CATC-GCTCAAGTATCCAGTGG-3’), MCR-1R (5’-CCATGCTCAAGTATCCAGTGG-3’), and probe MCR-1P (5’-Cy5-TGCAGACGCACAGCAATGCCTATGAT-TAGATAGACACCGTTCTCAC-3’), and probe MCR-1P (5’-Cy5-TGCAGACGCACAGCAATGCCTATGAT-TAGATAGACACCGTTCTCAC-3’) with TaqMan Fast Advanced Master Mix (Life Technologies, Burlington, Ontario, Canada), on an ABI 7500 FAST thermocycler (Applied Biosystems, Foster City, CA) by using manufacturer’s’ recommended conditions. The mcr-1 gene was confirmed by Sanger sequencing by using previously described oligonucleotides (2). The isolate was also PCR-positive for a blaCTX-M gene. The strain was sequenced by using MiSeq (Illumina, Victoria, British Columbia, Canada), and predicted to be sequence type 3944 based on multilocus sequence typing databases (http://github.com/tseemann/mlst; http://mlst.warwick.ac.uk/mlst/dbs/Ecoli) and serotype O159:H4 (3). Sequence type 3944 does not belong to any clonal groups; 1 isolate from Asia is in the MLST database. Abricate (http://github.com/tseemann/abricate) and PlasmidFinder (4) were used to query the SPAdes-assembled genome (5). Results showed that this isolate carries 3 plasmids that have IncR, IncFIA/HI, and IncI2 replicons. The mcr-1 gene was found on a 60,599-nt contig with the IncI2 replicon; this contig is ≈87% identical to pHNSHP45 (Figure) (2). The blaCTX-M-27 gene was found on the same contig as the IncFIA/HI replicon, and no resistance genes were found with the IncR replicon.

At the time of mcr-1 detection, the patient and all other patients who shared a hospital room with this patient,
all for <24 h, had been discharged. According to a public health representative, and on the basis of the limited exposure duration of roommates, discharged roommates were not screened. The patient improved clinically, so no changes to therapy were indicated.

The patient had traveled to China in November 2015 for 2 weeks, where he required catheterization in a hospital emergency department in Zhejiang Province for acute urinary retention. He experienced acute urinary retention and fever 6 days after catheter removal, requiring another catheter insertion and 3 days of intravenous antimicrobial drugs in Guangdong Province. He denied contact with farm animals, live poultry markets, or undercooked meat. On return to Canada, obstructive urinary tract symptoms persisted, requiring 5 emergency department visits before prostate resection.

Colistin is a last-resort antibacterial drug because of its toxic effects and is increasingly used for treating carbapenem-resistant Enterobacteriaceae (CRE). Plasmid-mediated resistance genes have been described in agricultural animals and meat, as well as in humans. Initial reports described the mcr-1 gene in China and Southeast Asia (2,7). Retrospective reviews have detected mcr-1 in Enterobacteriaceae from Europe, South America, Africa, and Japan (7).

Unlike laboratory detection of CRE, where screening media and automated susceptibility panels were available, no commercial screening media exist for mcr-1. MIC testing is only recommended for Enterobacteriaceae resistant to all other antimicrobial classes, and molecular testing may not be accessible. Furthermore, some Enterobacteriaceae are intrinsically resistant to colistin. Only Etest (http://etest.net/) could be performed in our laboratory, which is a limitation that may underestimate the actual MIC (8). However, the reference broth microdilution method is unavailable to most clinical laboratories. This isolate was identified by a technologist who recognized heavy growth of E. coli on a CNA plate, an unusual occurrence because CNA plates are used for the isolation of gram-positive bacteria while inhibiting gram-negative bacteria. Despite serendipitously identifying mcr-1 on the CNA, this method is an inadequate for detection of mcr-1.

Retrospective screening has identified mcr-1 isolates in Canada (9). However, we describe a prospectively identified patient in Canada with E. coli harboring the mcr-1 gene. The patient’s travel history suggested that acquisition occurred in China, although only 1% of inpatients with infection in Guangdong/Zhejiang Provinces harbor mcr-1 (2).

Limited laboratory screening procedures have implications for laboratories and public health. Routine colistin testing for Enterobacteriaceae would be costly and low-yield; however, without such testing, the real prevalence of mcr-1 will be underestimated. A coordinated approach to

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**Figure.** The mcr-1–containing contig from Escherichia coli isolated from a traveler returning to Canada from China compared to pHNSHP45 plasmid. BRIG (6) was used to generate a visual representation of the 60,599-nt contig assembled from the sequencing of the mcr-1–positive isolate to pHNSHP45 (2). Coding regions are represented in the inner ring; the mcr-1–containing contig is represented in the outer ring. The mcr-1 gene is indicated in a black segment in the inner ring.
the prevention of mcr-1 dissemination is needed, particularly to prevent the proliferation of an organism harboring a plasmid with mcr-1 and a carbapenemase (10).

Acknowledgments
We thank Michael Mulvey and Laura Mataseje for genome sequencing, which was supported by the Public Health Agency of Canada Genome Research and Development Initiative grant awarded to Dr. Mulvey. We also thank the laboratory technologists at Providence Health Care and the BC Centre for Disease Control Public Health Laboratory for their work on this study.

We acknowledge the National Microbiology Laboratory of Canada for contributions to the mcr-1 PCR testing.

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Carbapenem-Resistant Enterobacter spp. in Retail Seafood Imported from Southeast Asia to Canada

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DOI: http://dx.doi.org/10.3201/eid2209.160305

To the Editor: Carbapenems, antimicrobial drugs of last resort, are recommended only for severe community- and healthcare-associated multidrug-resistant bacterial infections. In Canada, carbapenem-resistant infection rates in hospitals remained low (≤0.25 cases/1,000 patient admissions) over 5 years’ (2009–2014) surveillance (1). Carbapenemase-producing bacteria have rarely been detected in the food chain in industrialized countries. However, carbapenemase genes were detected in bacteria isolated from produce in Switzerland (2) and seafood in Canada (3); implicated food items originated from Southeast Asia. We conducted targeted sampling to assess, using selective media, the occurrence of carbapenem-resistant Enterobacteriaceae in imported seafood products sold in Canada.

For testing, we selected 1,328 retail seafood samples: 928 were imported fresh and frozen raw shrimp collected during 2011–2015 by CIPARS (the Canadian Integrated Program for Antimicrobial Resistance Surveillance), and 400 comprised an assortment of imported niche-market fresh and frozen raw seafood collected specifically for this study during January–April 2015. Product information and origin country were recorded for each sample. We used chromoID CARBA agar (bioMérieux, St. Laurent, QC, Canada) to select putative colonies. To determine carbapenemase production on nonsusceptible (zone of inhibition <25 mm) isolates, we used disk diffusion susceptibility to ertapenem and meropenem (10 μg each) and the Carba NP test as previously described (4). Isolates were identified to species using matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (Bruker Daltonics Ltd, Milton, ON, Canada) and tested for susceptibility using the Sensititre Complete Automated System with the Sensititre NARMS Gram Negative Plate (CMV3AGNF)