Letter to the Editor

First report of an mcr-1-harboring Salmonella enterica subsp. enterica serotype 4,5,12:i:- strain isolated from blood of a patient in Switzerland

Sir,

In the past, colistin use has been mostly limited to veterinary medicine owing to its rather severe side effects, but given the increase in multidrug-resistant Gram-negative bacterial infections, the World Health Organization (WHO) recently re-labelled colistin as a ‘critically important antibiotic’. The first description of the plasmid-borne mobilizable colistin resistance gene mcr-1 in 2015 [1] caused great concern, as the ease of potential spread on a conjugative plasmid encoding resistance to polymyxins might change the resistance situation to colistin drastically. In line with this, mcr-1-mediated colistin resistance in Enterobacteriaceae, including Salmonella enterica, has since been reported from a wide range of geographical locations [2]. Here we report the first case of mcr-1-harboring S. enterica in Switzerland.

Salmonella enterica subsp. enterica serotype 4,5,12:i:- strain N17-0346 was isolated in 2017 from the blood of a 77-year-old male patient in Switzerland with no known travel history. The serotype was determined by the Swiss National Reference Centre for Enteropathogenic Bacteria and Listeria (NENT) according to the White-Kauffmann-Le Minor scheme. The strain was subjected to antimicrobial susceptibility testing to 16 antimicrobial agents by the disk diffusion method according to Clinical and Laboratory Standards Institute (CLSI) protocols and the results were evaluated according to CLSI criteria [3]. Determination of the colistin minimum inhibitory concentration (MIC) was performed by the broth microdilution method according to the European Committee on Antimicrobial Susceptibility Testing (EUCAST) (http://www.euCAST.org). Strain N17-0346 was phenotypically resistant to colistin (MIC = 4 mg/L), ampicillin and streptomycin but was susceptible to amoxicillin/clavulanic acid, cefazolin, cefotaxime, cefepime, gentamicin, kanamycin, nalidixic acid, ciprofloxacin, trimethoprim/sulfamethoxazole, azithromycin, fosfomycin, nitrofurantoin, tetracycline and chloramphenicol.

Strain N17-0346 was sequenced using a MiSeq platform (Illumina Inc, San Diego, CA) and a Nextera XT Library Kit (Illumina Inc.) utilising either 500 or 600 cycles of paired-end reads. De novo assembly using CLC Genomics Workbench v.9.0 (CLC Bio, Aarhus, Denmark) resulted in a genome size of 4 981 418 bp with 95 contigs and a GC content of 52.1%. The genome was annotated using the Rapid Annotation using Subsystem Technology (RAST) annotation server and 5061 coding sequences (CDS) were identified. In silico seven-gene multilocus sequence typing (MLST) using seq2mlst v.10.1 (https://github.com/lmc297/seq2mlst) identified strain N17-0346 as ST34. Core-genome MLST using the command-line implementation of SISTR v.1.0.2 classified it as ST3833327333. Genome-wide detection of antimicrobial resistance (AMR) genes using the method described and validated for Salmonella Typhimurium by Carroll et al. [4] and implemented in BTyper v.2.2.0 (https://github.com/lmc297/BTyper) in conjunction with the ARG-ANNOT AMR gene database, PlasmidFinder and PlasFlow v.1.0 produced the following hits: mcr-1 encoded on an IncX4 plasmid (pN17-0346) with no further AMR genes on the same contig; strA/strB (conferring streptomycin resistance) and sulI (encoding sulphonamide resistance) located on an IncQ1 plasmid; blβTEM-30 (coding for an inhibitor-resistant β-lactamase) on a contig classified as belonging to a plasmid; and aac(6′)-Ila (aminoglycoside acetyltransferase) encoded on the chromosome.

To generate a closed plasmid sequence of the mcr-1-coding sequence pN17-0346, the plasmid was sequenced using a Nextera DNA Flex Sample Preparation Kit on an Illumina MiniSeq Sequencer with 150-bp paired-end reads, yielding the closed 34 092-bp plasmid sequence with a per base coverage of >50× for all bases. RAST annotation determined 51 CDS and a partial insertion sequence (IS) element. The mcr-1-coding sequence was located directly upstream of an open reading frame encoding a hypothetical protein with similarities to a PAP2 superfamily protein that is frequently seen in association with mcr-1 [5]. A single copy of an incomplete version of ISAPl1 was located downstream of the mcr-1 cassette, but no ISAPl1 element was identified upstream of it.

Conjugal transfer of the mcr-1-harboring IncX4 plasmid from donor strain N17-0346 to the recipient Escherichia coli HK225 (streptomycin- and rifampicin-resistant) was tested at 25°C and 37°C in liquid medium as well as on solid agar plates. Transconjugants carrying the mcr-1 gene as confirmed by PCR were found in all of these experiments, with a conjugation efficiency of 2.2 × 10^-4 per donor cell on solid agar at 37°C. The parent E. coli HK225 had a colistin MIC of 1 mg/L; after conjugation of the IncX4 plasmid, the MIC increased to 4 mg/L. To our knowledge, this is the first report of a Salmonella strain carrying an mcr-1 colistin resistance gene in Switzerland.

The full sequence of strain N17-0346 has been deposited in GenBank under the accession no. QEA00000000 and the sequence of the mcr-1-harboring plasmid pN17-0346 under accession no. NZ_CP031291.1

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Competing interests

None declared.

Ethical approval

Not required.

References


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