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Predominance of CTX-M-15-producing ST131 strains among ESBL-producing *Escherichia coli* isolated from asylum seekers in the Netherlands

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Objectives: Numerous studies show increased prevalence of MDR bacteria amongst asylum seekers, but data on the molecular profiles of such strains are limited. We aimed to evaluate the molecular profiles of ESBL-producing *Escherichia coli* (ESBL-*E. coli*) strains isolated from asylum seekers and investigate their phylogenetic relatedness.

Methods: WGS data of ESBL-*E. coli* isolates from asylum seekers, retrieved from 1 January to 31 December 2016, were analysed to assess MLST STs, *fim* types, phylogroups and resistance genes. Fifty-two ESBL-*E. coli* isolates from the Dutch–German border region were used for genome comparison purposes as a control group.

Results: Among 112 ESBL-*E. coli* isolates from asylum seekers, originating mostly from Syria ($n=40$) and Iraq ($n=15$), the majority belonged to ST131 (21.4%) and ST10 (17.0%). The predominant gene for β -lactam resistance was *bla*_{CTX-M-15} (67.9%), followed by the often co-detected *bla*_{TEM-1B} (39.3%). No *mcr* or carbapenemase genes were detected. The majority of the strains belonged to phylogroups B2 (38.4%) and A (32.1%), carrying *fimH27* (25%) and *fimH30* (19.6%). A core genome MLST minimum spanning tree did not reveal clusters containing strains from the asylum seekers and the control group. Five clusters were formed within the asylum seeker group, by strains isolated from people originating from different countries.

Conclusions: The most frequently isolated clones in this study were isolated on a regular basis within the Dutch population before the increase in the asylum seeker population. No *mcr*- or carbapenemase-producing clones were detected among the asylum seeker population. Minor clustering was observed amongst the asylum seeker strains.

Introduction

Increased numbers of refugees and asylum seekers have entered Europe during the last decade. At the end of 2018, the United Nations High Commissioner for Refugees reported nearly 6.5 million refugees and migrants residing in Europe.¹ According to the Immigration and Naturalization Service of the Ministry of Justice and

Security, 196 519 asylum seekers entered the Netherlands from January 2013 to December 2018. The main countries of origin include the Syrian Arab Republic, Afghanistan, Iraq, Iran and Eritrea.²

This increase in displaced populations has public health implications. The health needs of refugees and asylum seekers require

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coordinated efforts from public health institutions. Medical care of refugees and asylum seekers can be challenging for public health systems and hospitals.³ Although data on the epidemiology of MDR organisms (MDROs) in most of the countries of origin are limited, studies have shown an increased MDRO prevalence amongst refugees and asylum seekers. A recently published systematic review and meta-analysis on antimicrobial resistance among migrants in Europe showed a pooled prevalence of any detected antimicrobial resistance carriage or infection of 25.4% (95% CI 19.1%–31.8%). The pooled prevalence of MDR Gram-negative bacteria was 27.2% (95% CI 17.6%–36.8%).⁴

The need for adequate and timely MDRO detection and, subsequently, the implementation of infection prevention measures regarding refugee inpatients is imperative. WGS has proved to be a valuable tool for microbial analysis on a molecular level and for MDRO outbreak investigation. During the last decade, WGS has been increasingly integrated in microbiology laboratories as part of the daily routine diagnostics, as it became easier, faster and cheaper to use.⁵ This technique combined with bioinformatics tools can provide, in less than 48 h, an abundance of valuable information regarding MDROs, including detection, identification, genetic resistance profile, genotype and epidemiological typing. Further analysis of the data can also determine genetic and phylogenetic relatedness amongst strains, revealing clustering and aiding outbreak investigation.^{6,7}

The main mechanism of resistance to β -lactams in *Escherichia coli* strains is production of ESBLs, a group of enzymes mainly encoded by CTX-M, TEM and SHV variants.⁸ During the past decade, *bla*_{CTX-M} genes have been increasingly detected in Gram-negative bacteria, including *E. coli*, worldwide, leading to a 'CTX-M pandemic' situation.⁹ This rapid spread of certain CTX-M-producing *E. coli* lineages carries with it difficulties in typing. Conventional typing methods, such as PFGE and MLST, do not have the discriminatory power to identify clusters of dissemination. Even next-generation sequencing ('NGS')-based typing, which has higher discriminatory power, does not always provide conclusive proof of dissemination and should always be interpreted in combination with epidemiological data. Additional methods, such as typing of *fimH* genes, are useful to subtype certain lineages.¹⁰ In addition, sequences of epidemiologically unrelated isolates, so-called context isolates, should be added to the analysis to provide insight into the genetic background of the bacterial population.¹¹

In this study, we evaluated the molecular profiles, including STs, *fim* types, phylogroups and resistomes, of ESBL-producing *E. coli* (ESBL-*E. coli*) strains isolated from hospitalized asylum seekers in 2016. We compared the molecular epidemiology of the isolates from the refugees with that of a collection of ESBL-*E. coli* strains from the Dutch-German border region, from hospitalized patients and a community population in 2012, before the number of refugees started to increase.

Materials and methods

Study design

When entering the Netherlands, all asylum seekers are appointed to an asylum seeker centre (ASC) and are registered under the ASC's address. Asylum seekers included in the study were identified by the ASC address at which they resided. Data on patient characteristics were retrospectively collected from the Certe laboratory system. Study material included screening

samples for MDRO carriage before admission (throat, rectum and nose) and clinical samples (e.g. blood, wounds and urogenital) from asylum seekers. All of these samples were obtained as part of standard care.

Study population

We included asylum seekers, hospitalized in the northern part of the Netherlands from 1 January to 31 December 2016, who tested positive for ESBL-*E. coli* strains. Demographic data, such as age, sex and country of origin, were collected from the laboratory system and the healthcare system for asylum seekers. All ESBL-*E. coli* strains isolated from the study population were included in the group of asylum seeker strains. Duplicate strains with the same molecular, phenotypic and genotypic profile that were from the same asylum seeker were excluded.

Bacterial identification and antimicrobial resistance mechanism detection

ESBL-*E. coli* strains from asylum seekers were obtained in the Certe laboratory. This laboratory performs routine microbiological analyses for primary and secondary medical care in the north-east of the Netherlands, including the ASC population in this part of the country. Screening and clinical samples were cultured in a variety of selective (solid) media used for MDRO detection, including MacConkey agar with 0.5 mg/L ciprofloxacin and 2 mg/L gentamicin (Mediaproduits BV, Groningen, The Netherlands), ChromID ESBL agar and ChromID Carbapenemase agar (both from bioMérieux, Marcy-l'Étoile, France). The presence of ESBL was confirmed with cefotaxime/clavulanate, ceftazidime/clavulanate and cefepime/clavulanate Etests (bioMérieux). Possible carbapenemase-producing Enterobacterales (CPE) were confirmed by CIM test and PCR (Check-Direct CPE assay, Check-Points, Wageningen, The Netherlands) and typed by the national reference network for CPE at the RIVM (National Institute for Public Health), as part of standard care.

Antimicrobial phenotype detection

Susceptibility to amikacin, amoxicillin/clavulanic acid, ampicillin/sulbactam, cefepime, cefotaxime, ceftazidime, ciprofloxacin, colistin, ertapenem, fosfomicin, gentamicin, imipenem, levofloxacin, meropenem, nitrofurantoin, piperacillin/tazobactam, tigecycline and trimethoprim/sulfamethoxazole was determined using Vitek 2 (bioMérieux). EUCAST guidelines were used for interpretation of MICs.

Control group isolates

The control isolate collection consisted of 41 ESBL-*E. coli* from hospitalized patients and people in the community in the Netherlands and 11 ESBL-*E. coli* strains from hospitalized patients in Germany.¹² The isolates were collected in 2012 and were used as context isolates in the genomic comparisons. All strains included in the control group were analysed using the same workflow as the asylum seeker group strains.

DNA isolation

A total of 112 frozen ESBL-*E. coli* strains isolated from unique asylum seekers were recultured and incubated for 24 h at 37°C. DNA was extracted using the DNeasy UltraClean Microbial Kit (MoBio Laboratories, Carlsbad, CA, USA), according to the manufacturer's instructions. A 5 μ L aliquot of each isolate was suspended with 300 μ L of PowerBead solution. DNA purity was measured using a NanoDrop 2000C spectrophotometer (Thermo Fisher, Waltham, MA, USA). DNA concentration was measured with a Qubit 2.0 fluorometer, using the double-stranded DNA BR Assay Kit (Life Technologies, Carlsbad, CA, USA).

WGS

Prior to library preparation, isolated DNA was diluted to a concentration of 0.2 ng/μL. DNA library preparation was performed with the Nextera XT v.01 (Illumina Inc., San Diego, CA, USA) kit using 5 μL of diluted DNA according to the manufacturer's instructions. Libraries were sequenced on an MiSeq sequencer (Illumina Inc.) aiming to generate 250 bp paired-end reads.

Quality check and WGS data analysis

Trimming and *de novo* assembly was performed using CLC Genomics Workbench v.10.1.2 (QIAGEN, Hilden, Germany). A minimum Phred score (Qscore) of 30 was used. Six parameters were checked for assembly quality: number of contigs <1000, N50 > 15 000, maximum contig length >50 000, percentage of reads used for the assembly >90%, coverage >30× and percentage of the expected genome size >90% to <115%.

The assembled genomes were uploaded to SeqSphere v.5.5.1 (Ridom GmbH, Münster, Germany) for investigation of phylogenetic relatedness. A minimum spanning tree based on allelic mismatch between the isolates was designed. A maximum of 10 allelic differences was considered as clonal clustering.

SNP-based Neighbor-Joining (NJ) trees were constructed based on the genome sequences using Ridom SeqSphere+ version 5.1.0 (Münster, Germany) with default settings. The genomes were analysed using an *ad hoc E. coli* scheme based on 2764 targets, including 242 851 bp.

Assembled genomes were uploaded to the web tools ResFinder 3.1 to identify acquired resistance genes¹³ and FimTyper 1.0 to determine *fim* type.¹⁰ Phylogroups were determined via the EzClermont web app and command-line tool.^{14,15}

Sequences are publicly available at the ENA database (study accession number PRJEB36686).

Statistical analysis

Data were collected in and analysed with SPSS (version 2.23). Descriptive statistics were used for the general characteristics of the study population.

Ethics

This study was evaluated by the Ethics Committee and approval was waived in accordance with Dutch legislation owing to its retrospective nature (University Medical Centre Groningen, METc number 2016/516). No written informed consent was obtained from patients for the use of retrospective data. Patient information was anonymized and de-identified prior to analysis.

Results

General characteristics of the study population

We evaluated single ESBL-*E. coli* isolates from 112 asylum seekers. General characteristics of the study population and the included samples are described in Table 1.

Routinely measured resistance

Antimicrobial susceptibility of the strains to different antibiotic agents, routinely tested in the Certe laboratory, is described in Table S1 (available as [Supplementary data](#) at JAC Online).

All of the isolates were resistant to penicillins, cephalosporins and combinations of penicillins and β-lactamase inhibitors. Also, 56.3% of the isolates were resistant to trimethoprim/sulfamethoxazole. Resistance to ciprofloxacin was observed in 31.3% of the isolates. Regarding aminoglycosides, 27.7% and 33.0% of the

Table 1. General characteristics of the study population and the included samples; *N* = 112

Female, <i>n</i> (%)	75 (67)
Age (years), median (IQR)	28.0 (20.4–36.1)
Number of days in the Netherlands, median (IQR)	192 (77–347)
Country of origin, <i>n</i> (%)	
Syria	40 (35.7)
Iraq	15 (13.4)
Iran	12 (10.7)
Afghanistan	9 (8.0)
Eritrea	6 (5.4)
other from Europe	6 (5.4)
other from Eastern Europe/Russia	4 (3.6)
other from Asia	8 (7.1)
other from Africa	9 (8.0)
Samples, <i>n</i> (%)	
rectal	101 (90.2)
urine	6 (5.4)
skin	2 (1.8)
sputum	1 (0.9)
nasal	1 (0.9)
stool	1 (0.9)

isolates were resistant to gentamicin and tobramycin, respectively. No isolate was resistant to meropenem or imipenem. All isolates were susceptible to fosfomycin and colistin.

ST and genotypic profile

The most frequent ST of the asylum seeker isolates was ST131 (21.4%), followed by ST10 (17.0%), ST38 (8.0%) and ST69 (8.9%). Among the control group isolates, the most frequent ST was ST38 (15.4%), followed by ST10 and ST131 (both 11.5%) and ST58 (7.7%). Table 2 shows the STs of the asylum seeker isolates and the *bla*_{CTX-M} resistance genes carried by the isolates for each ST.

The distribution of *bla*_{CTX-M} resistance genes harboured by the isolates from asylum seekers and the control group can be seen in Figure 1.

The most frequently observed CTX-M gene for β-lactam resistance was *bla*_{CTX-M-15} for both groups, followed by *bla*_{CTX-M-27} for the asylum seekers and *bla*_{CTX-M-1} for the control group. Other, non-CTX-M β-lactam resistance genes detected in the asylum seeker isolates were *bla*_{TEM-1B} (*n* = 44, 39.3%), *bla*_{OXA-1} (*n* = 11, 9.8%), *bla*_{SHV-12} (*n* = 2), *bla*_{TEM-33} (*n* = 2), *bla*_{DHA-1} (*n* = 2), *bla*_{TEM-1C} (*n* = 1) and *bla*_{CMY-60} (*n* = 1).

All of the asylum seeker isolates carried resistance genes related to more than one antibiotic group, including aminoglycosides, fluoroquinolones, sulphonamides and trimethoprim. Regarding aminoglycoside resistance, 44 isolates harboured *strA*, 43 harboured *strB*, 42 harboured *addA5* and 21 harboured *acc(3)-IId*. The main genes carried by the isolates that encoded quinolone resistance were *qnrS1* (*n* = 26) and *aac(6')Ib-cr* (*n* = 10). For sulphonamide resistance, 48 isolates carried *sul1* and 40 isolates carried *sul2*. Lastly, the main trimethoprim resistance genes detected

were *dfrA17* ($n=44$) and *dfrA14* ($n=14$). Of note, no *mcr* and carbapenemase genes were detected.

Phylogroup and *fim* type

Asylum seeker isolates belonged primarily to phylogroups B2 ($n=43$, 38.4%) and A ($n=36$, 32.1%). The remaining isolates belonged to phylogroups D ($n=24$, 21.4%), B1 ($n=6$, 5.4%) and F ($n=3$, 2.7%).

Subtyping of *fimH* alleles of isolates from asylum seekers showed that *fimH27* was the most frequent type ($n=28$, 25%), followed by *fimH30* ($n=22$, 19.6%) and *fimH5* ($n=6$, 5.4%). Thirteen isolates (11.6%) did not carry any *fim* gene. The remaining isolates carried a wide variety of different *fim* genes. Of note, 19 out of the 24 isolates that belonged to ST131 carried *fimH30* and 8 out of the 19 isolates that belonged to ST10 carried *fimH27*.

Table 2. STs of the asylum seeker isolates and the *bla*_{CTX-M} resistance genes carried by the isolates for each ST

ST	Total, N (%)	<i>bla</i> _{CTX-M} resistance gene, n (%)				
		<i>bla</i> _{CTX-M-15}	<i>bla</i> _{CTX-M-27}	<i>bla</i> _{CTX-M-3}	<i>bla</i> _{CTX-M-14}	<i>bla</i> _{CTX-M-14b}
ST131	24 (21.4)	12 (50.0)	7 (29.2)	1 (4.2)	0 (0)	0 (0)
ST10	19 (17.0)	16 (84.2)	0 (0)	2 (10.5)	1 (5.3)	0 (0)
ST69	10 (8.9)	8 (80)	0 (0)	1 (10)	0 (0)	0 (0)
ST38	9 (8.0)	1 (11.1)	1 (11.1)	0 (0)	1 (11.1)	3 (33.3)
ST12	7 (6.3)	7 (100.0)	0 (0)	0 (0)	0 (0)	0 (0)
ST120	4 (3.6)	3 (75.0)	0 (0)	0 (0)	0 (0)	0 (0)
ST93	4 (3.6)	4 (100.0)	0 (0)	0 (0)	0 (0)	0 (0)
ST1193	3 (2.7)	2 (66.7)	0 (0)	0 (0)	0 (0)	0 (0)
ST73	3 (2.7)	2 (66.7)	0 (0)	0 (0)	0 (0)	0 (0)
ST648	2 (1.8)	2 (100.0)	0 (0)	0 (0)	0 (0)	0 (0)
ST3877	2 (1.8)	2 (100.0)	0 (0)	0 (0)	0 (0)	0 (0)
ST58	2 (1.8)	2 (100.0)	0 (0)	0 (0)	0 (0)	0 (0)
Other	23 (20.5)	14 (60.9)	1 (4.3)	1 (4.3)	2 (8.7)	0 (0)

Analysis of the core genome MLST (cgMLST) NJ tree, including asylum seeker and control group isolates, is shown in Figure 2. Furthermore, the cgMLST NJ tree is shown in Figure S1 in rectangular form, including metadata, such as country of origin of the asylum seekers, isolate ST and isolate phylogroup (given in the columns next to the tree).

Phylogenetic relatedness and cluster analysis

A minimum spanning tree of the asylum seeker and control group isolates can be seen in Figure S2. The observed allelic distance ranged from 0 to 2371 alleles.

Cluster analysis revealed five clusters within the asylum seeker group isolates (Figure S2). Cluster 1 consisted of six isolates belonging to ST69, phylogroup D, subtype *fimH27* and carrying *bla*_{CTX-M-15} (Figure 2). Two of the isolates were from asylum seekers originating from Syria and the remaining four were from asylum seekers originating from Palestine, Afghanistan, Iraq and Yugoslavia. The isolates from the Syrian and Iraqi asylum seekers were from rectal and skin samples, cultured a day apart, respectively, and the isolates from the Palestinian and Afghan asylum seekers came from sputum and urine samples, respectively, cultured 17 days apart. Cluster 2 was formed by five isolates belonging to ST10, phylogroup A, subtype *fimH27* and carrying *bla*_{CTX-M-15} (Figure 2). All isolates were obtained from asylum seekers originating from different countries, namely Syria, Iraq, Eritrea, Turkey and Benin. The isolates from the Syrian and Iraqi asylum seekers were from rectal samples, cultured 4 days apart. Cluster 3 consisted of four isolates belonging to ST12, phylogroup B2, no *fim* subtype and carrying *bla*_{CTX-M-15} (Figure 2). Two of the strains were isolated from Syrian asylum seekers, one from a Mongolian asylum seeker and one from an Afghan asylum seeker. One of the isolates that was from a Syrian asylum seeker and the isolate that was from an Afghan asylum seeker were from rectal samples, cultured a week apart. Cluster 4 included two isolates belonging to ST1193, phylogroup B2, subtype *fimH64* and carrying *bla*_{CTX-M-15} (Figure 2). One was isolated from a Syrian asylum seeker and the other from an Iranian;

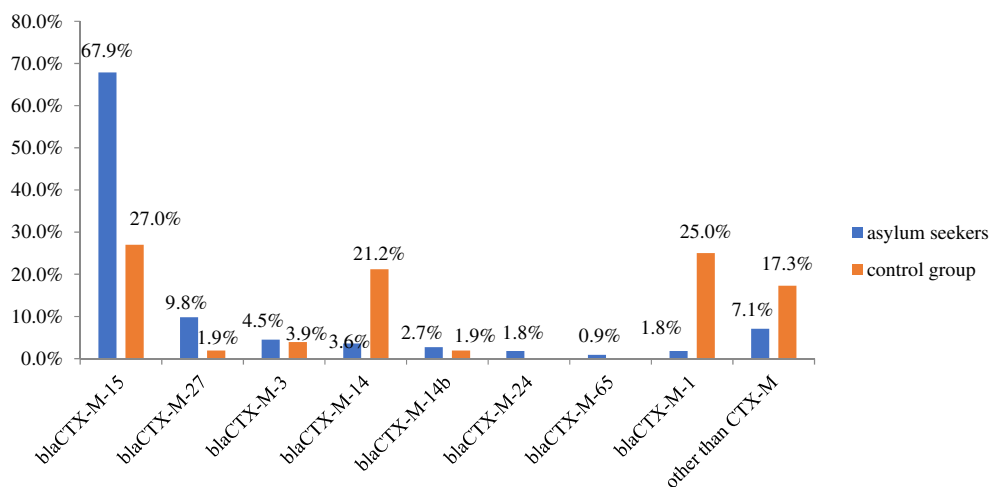


Figure 1. Distribution of the most frequently detected *bla*_{CTX-M} genes among the asylum seeker and control group isolates. This figure appears in colour in the online version of JAC and in black and white in the print version of JAC.

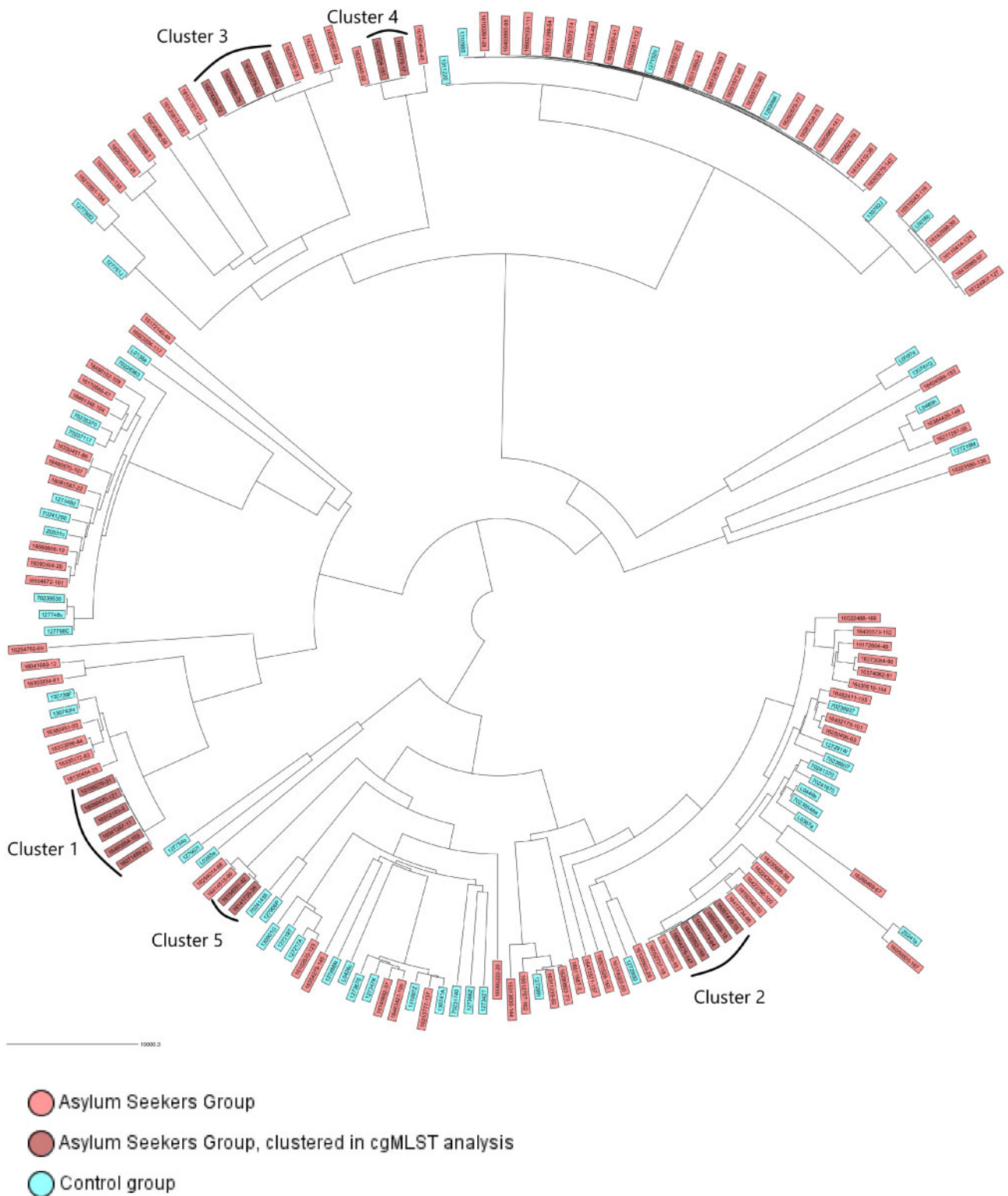


Figure 2. cgMLST Neighbor-Joining tree, including asylum seeker and control group isolates. Genomes were analysed using an *ad hoc* *E. coli* scheme based on 2764 targets, including 24 2851 bp. Control group isolates were from health institutions near the Dutch–German border region. Isolates that formed clusters in further phylogenetic analysis are indicated in dark pink. This figure appears in colour in the online version of JAC and in black and white in the print version of JAC.

they were isolated from urine and rectal samples, respectively, cultured 7 months apart. Cluster 5 consisted of two isolates belonging to ST120, phylogroup A, subtype *fimH237* and carrying *bla*_{CTX-M-15} (Figure 2). The two isolates were from a Syrian asylum seeker and an Eritrean asylum seeker; they were isolated from rectal samples, cultured 10 days apart. No clusters contained isolates belonging to both asylum seekers and control group isolates.

Discussion

A total of 112 ESBL-*E. coli* strains isolated from asylum seekers were analysed using WGS in order to investigate their phylogenetic relatedness and possible transmission within the asylum seeker population in the northern part of the Netherlands. The asylum seeker study population originated mainly from Syria and had been residing in the Netherlands for a median of 192 days.

All isolates were phenotypically resistant to β -lactams and exhibited various resistance profiles, with all of them being resistant to at least one more antibiotic group, such as aminoglycosides, quinolones and sulphonamides. The genes that encode resistance to these antibiotic agents are often located on plasmids that can co-harbour different resistance genes and can be horizontally transferred amongst Enterobacteriales, such as *E. coli*, rendering the strains MDR.¹⁶

The majority of the isolates belonged to ST131 and ST10, and harboured a *bla*_{CTX-M-15} gene. This is in accordance with the epidemiological profile of the high-risk ST131 *bla*_{CTX-M-15} clone. ST131 *bla*_{CTX-M-15} is currently globally disseminated and is identified as the most widespread CTX-M ESBL enzyme worldwide.^{17,18} The Netherlands has also been affected by the ST131 *bla*_{CTX-M-15} clone. In a recently published study conducted in Dutch hospitals, between 2014 and 2016, the dominant clone found among ESBL-*E. coli* blood isolates was ST131 carrying *bla*_{CTX-M-15}.¹⁹ Furthermore, in a study conducted in the Netherlands in 2016, the clone was isolated among community-associated and hospitalized patients,²⁰ indicating that the clone existed in both the community and hospitals in the Netherlands before the number of refugees started to increase in 2015 and 2016.

The majority of the study isolates harboured *bla*_{CTX-M-15}, regardless of the ST to which they belonged. Even though strains carrying *bla*_{CTX-M-15} have been reported all over Europe, strains carrying this gene are isolated at a higher rate in Middle Eastern, Asian and African regions.²¹ Furthermore, epidemiological data on the distribution of such strains indicate that African and Asian regions could serve as a reservoir and facilitate global dissemination.²² A German study that investigated the antibiotic resistomes of refugees reported high prevalence rates for β -lactamase genes: mainly *bla*_{TEM}, *bla*_{CTX-M} group 1 and *bla*_{SHV}.²³ Another German study reported high detection of *bla*_{CTX-M} group 1 genes, followed by *bla*_{TEM} and *bla*_{SHV}, among ESBL-producing Enterobacteriaceae isolates from Libyan and Syrian patients.²⁴ Furthermore, a study performed in Saudi Arabia showed a prevalence of *bla*_{CTX-M-15} or *bla*_{CTX-M-14} of 60% among ST131 uropathogenic *E. coli* strains.²⁵ In addition, an Iranian study, published in 2017, demonstrated a high prevalence of ST131 *bla*_{CTX-M-15} amongst clinical *E. coli* strains.²⁶ A high prevalence of strains carrying these genes amongst asylum seekers from Iran, Syria and Afghanistan was also documented in our study.

Despite the fact that some clustering among the isolates from the asylum seekers was observed, no clear pattern of transmission was documented. Isolates that exhibited close phylogenetic relatedness formed five clusters. As expected, isolates within each cluster exhibited identical genetic characteristics, such as ST, phylogroup and *fim* type. However, isolates included within each cluster did not show a clear epidemiological link, since they were isolated from asylum seekers mostly originating from different countries. Furthermore, even though certain isolates in clusters 1, 2, 3 and 5 were isolated within 10 days or less, clear epidemiological links cannot be hypothesized without additional information, such as department and institution of hospitalization, ASC of residence and countries they have travelled through before entering the Netherlands. Due to limited clustering and wide dispersion of the origin of the asylum seekers carrying the isolates within each cluster, no conclusion can be drawn regarding the geographical epidemiology and origin of the isolates.

A limited number of studies have previously sequenced MDROs in a refugee/asylum seeker patient population.^{27,28} To our knowledge, this is the first study to investigate ESBL-*E. coli* strains isolated from asylum seekers using WGS on a large scale documenting various genetic characteristics, such as STs, genotypic resistance profiles and phylogenetic relatedness. This information is still scarce in the related literature and can help to optimize treatment, hospital hygiene strategies and infection control measures. Furthermore, our study population exhibited a large variation in age, number of days in the Netherlands and country of origin, reflecting the main countries from which migrants originate, namely Syria, Afghanistan and Iraq.¹

Due to the retrospective nature of this study, we did not have access to important information, such as travelling and antibiotic consumption history. In addition, information regarding asylum seeker hospitalization, such as reason for admission, department of admission, duration of hospitalization and treatment given, was not available. Furthermore, we had no access to data regarding the specific ASCs where our study population resided after their arrival in the Netherlands. Close contact within a facility can lead to transmission of MDROs. In our study, no clear pattern of transmission was observed.

Conclusions

The most frequently isolated clones in the study are already detected on a regular basis within the Dutch population. No *mcr*- or carbapenemase-producing clones were detected among the asylum seeker population. No clustering between asylum seekers and control group strains was observed.

Even though no assumptions can be made on whether transmission within the asylum seeker population occurs or not, small clustering within the asylum seeker strains could be an indication of this. Based on the results of this study, there is no clear evidence as to whether asylum seekers obtained their MDROs in their country of origin, during their journey to the Netherlands or after their arrival in the Netherlands. Asylum seekers originating from the same country showed a large variability in resistance and phylogenetic relatedness.

Further research on the genetic characteristics of MDRO isolates carried by asylum seekers could reveal important information on transmission and cluster formation.

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Transparency declarations

J.W.R. is currently an employee of IDbyDNA. IDbyDNA did not have any influence on the interpretation of the reviewed data and conclusions drawn, or on the drafting of the manuscript, and did not (financially) support the study. All other authors: none to declare.

Supplementary data

Table S1 and Figures S1 and S2 are available as [Supplementary data](#) at JAC Online.

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