Microbiology

Chemotherapy

Carriage Rates and Characteristics of Enterobacteriaceae Producing Extended-Spectrum Beta-Lactamases in Healthy Individuals: Comparison of Applicants for Long-Term Care and Individuals Screened for Employment Purposes

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Abstract

Aims: We compared the prevalence of extended-spectrum beta-lactamase (ESBL) producers in the faecal samples of 1,109 healthy individuals screened for employment purposes and in 531 asymptomatic individuals applying for long-term care (LTC). *Methods:* Eosin-methylene blue agar plates supplemented with 2 mg/l cefotaxime were used to determine which individuals were ESBL producers. ESBL phenotype was confirmed by double-disk synergy test and ESBL genes were identified by sequencing. ESBL producers were characterized by co-resistance and integron carriage. *Results:* ESBL producers were more frequent in the LTC applicants than in the employment screening individuals (7.2 vs. 2.0%; p < 0.0001), with 43 *Escherichia coli*, 18 *Klebsiella pneumoniae*, 1 *Klebsiella oxytoca* and 1 *Proteus mirabilis* being found. In the employment screening individuals, only *E. coli* was found. Most ESBL genes (79.4%, 50/63) were bla_{CTX-M} type; $bla_{CTX-M-15}$ was more frequent in the LTC applicants (p < 0.001). Regarding ESBL genes and integron diversity, *E. coli* isolates from the LTC applicants were more similar to *K. pneumoniae* than to *E. coli* from the employment screen-

ing individuals. *Conclusion:* These differences in the characteristics of ESBL producers may represent different sources of colonization. Most LTC applicants harboured *K. pneumoniae* or *E. coli* that were probably hospital-acquired whereas the *E. coli* isolates of many healthy individuals showed similarities to environmental *E. coli*.

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Introduction

Enterobacteriaceae that produce extended-spectrum beta-lactamases (ESBLs) have become a major problem worldwide, especially since the occurrence and spread of the bla_{CTX-M} family. Many of these enzymes, in contrast to the bla_{TEM} and bla_{SHV} groups, are linked to epidemic clones [1] and some of them, e.g. the $bla_{CTX-M-15}$-producing *Escherichia coli* O25b:ST131 and *Klebsiella pneumoniae* ST11, ST15 and ST147 clonal complexes, have become pandemic both in the nosocomial and community settings as well as in long-term care (LTC) facilities [1–3]. The switch from *K. pneumoniae* to *E. coli* as the major ESBL producer species is also linked to the dissemination of bla_{CTX-M} genes [4].

The gastrointestinal tracts of humans and even animals are obviously a major source of ESBL producer spe-

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cies [5, 6] and also serve as a site for the horizontal spread of resistance genes [7]. The duration of asymptomatic carriage can be very long, up to several years in some cases [8], and international travel contributes to even more efficient spreading [9]. Carriage rates, and therefore the risk of colonization, in travelers returning home are highest in Southeast Asia [6, 8], as high as 69.3%. North Africa and the eastern Mediterranean are also associated with a higher carriage risk in travelers [9].

Although a number of studies have been published on faecal carriage rates in different populations (both patients and asymptomatic individuals), colonization patterns have only started to be understood. Therefore, the aim of this study was to compare the prevalence of faecal carriage and the characteristics of ESBL producers among applicants for LTC and individuals screened for employment purposes.

Materials and Methods

Samples and Isolates

A total of 1,640 faecal samples from 2 groups of asymptomatic individuals not suffering from infection at the time of sampling were investigated between March 2009 and April 2010. Samples were collected and sent for screening for enteric pathogens from 1,109 individuals (300 males and 809 females with a median age of 34 years, range 15–68 years) being screened for employment eligibility purposes (e.g. jobs at hospitals, kindergartens and food-processing plants) and 531 individuals (218 males and 313 females with a median age of 75 years, range 0–100 years) who needed to be screened prior to admittance to LTC facilities. The vast majority of individuals were located in north-eastern Hungary from the EU regions of Northern Hungary and the Northern Great Plain.

The samples were directly inoculated onto eosin-methylene blue agar plates supplemented with 2 mg/l cefotaxime. All colonies with different morphology were further identified by means of biochemical tests; enterobacterial isolates were then processed further.

According to Clinical and Laboratory Standards Institute guidelines, antibiotic susceptibility testing was performed by the disc-diffusion method against imipenem, cefotaxime, ceftazidime, cefepime, ciprofloxacin, colistin, trimethoprim–sulfamethoxazole (co-trimoxazole), doxycycline, amikacin, gentamicin, tobramycin and tigecycline. All isolates showing a decreased susceptibility to at least 1 third-generation cephalosporin or to cefepime were tested for ESBL phenotype using the double-disc synergy test (Oxoid, Basingstoke, UK) and then re-identified using a MALDI Biotyper (Bruker, Bremen, Germany). Imipenem-resistant isolates were also tested using the Hodge test as specified in the Clinical and Laboratory Standards Institute guidelines.

In the case of isolates displaying an ESBL phenotype, DNA was extracted by heat treatment. PCR amplifications were carried out in a MyCycler PCR machine (BioRad, Hercules, Calif., USA). Species identification for *K. pneumoniae* and *E. coli* isolates was confirmed by species-specific PCRs [10, 11].

Genetic Relatedness

The epidemiological relationship was analyzed by enterobacterial repetitive intergenic consensus PCR (ERIC-PCR) with ERIC2 and ERIC1R primers (as described earlier [12]) and pulsed-field gel electrophoresis (PFGE). Plugs were prepared as described earlier [13], macrorestriction was performed using XbaI (Fermentas, Vilnius, Lithuania) in a CHEF DRIII machine (Bio-Rad) in 1% SeaKem Gold agarose (Lonza) at 14°C. Electrophoresis was performed at 6 V/cm, with a reorientation angle of 120°, and switch times were ramped between 2 and 64 s for 20 h for both species. Gels were stained with ethidium bromide and visualized under UV light. Banding patterns were analyzed with Fingerprinting II software (Bio-Rad). The threshold for probable unrelatedness was set at <85% similarity. The Simpson index of diversity (D) was calculated to assess diversity, as described earlier [14].

Identification of ESBL Genes

The bla_{TEM}, bla_{SHV}, bla_{CTX-M} genes were detected by PCR, as described previously [15–17]; bla_{TEM} and bla_{SHV} genes were identified by sequencing (Macrogen, Amsterdam, The Netherlands) using the same primers, while bla_{CTX-M} genes were identified by the sequencing of primers of group-specific PCRs [18]. Sequence alignment and analyses were performed using CLC DNA Workbench v4.0 (CLC Bio, Aarhus, Denmark).

Aminoglycoside Resistance Genes and Characterization of Integrons

The aminoglycoside resistance genes *aac(3')-IIa (aacC2)*, *aac(6')-Ib (aacA4)*, *aph(3')-Ia (aphA1)*, *ant(2')-Ia (aadB)*, *ant(3')-Ia (aadA1)*, *armA*, *rmtA* and *rmtB* were sought for as previously described [19–22]. The detection of class 1 and class 2 integrons was performed by PCR assays according to Mazel et al. [23] and the amplification and sequencing of the variable regions were performed as described by White et al. [24], using newly designed internal primers when necessary. Sequences were assembled in the CLC DNA Workbench 4.0 (CLC Bio), and gene cassettes were identified using the GenBank (http://www.ncbi.nlm.nih.gov). Identification of class 1 and class 2 integrons with variable regions of the same size was performed by restriction analysis using at least two enzymes for each cassette array, including *Eco*RI, *Hind*III, *Mse*I and *Rsa*I (Fermentas).

Phylogenetic Analysis of E. coli and Virulence Genes

Phylogenetic groups of *E. coli* were determined using the multiplex PCR method developed by Clermont et al. [25]. Phylogroups A and B1 are considered commensal, while groups B2 and D include strains responsible for extra-intestinal infections [25]. The pandemic O25b-ST131 clone was screened in the isolates of the phylogenetic group B2 with a PCR-based assay [26].

A multiplex PCR assay was used to determine the presence of virulence factor genes characteristic for enterovirulent *E. coli* pathotypes, as described previously [27].

We used PCR with previously described primers to search for genes coding for the putative extra-intestinal virulence factors of *E. coli*, including adhesins (*papC*, *fimH*, *sfaS* and *sfa/focDE*), toxins (*cnf1* and *cvaC*), factors related to iron acquisition (*iutA* and *fyuA*), the capsule system (*kpsMT II*) and miscellaneous factors (*ibeA*, PAI, *traT* and *csgA*) [28].
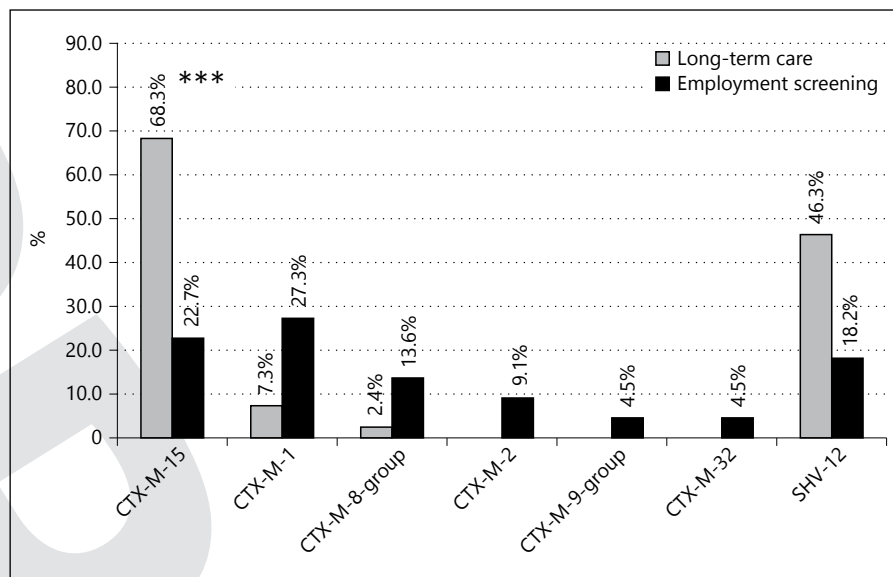
Fig. 1. Distribution of ESBL genes in the 2 groups. *** p < 0.001.

Statistical Analysis

Prevalences were compared by means of the χ^2 test or the Fisher exact test as appropriate. Age distribution in the different groups was compared by means of the Kolmogorov-Smirnov test. The distribution of genes coding for ESBLs, aminoglycoside-modifying enzyme genes and co-resistance patterns were all analyzed by χ^2 test, and the pairwise comparisons derived were adjusted with the Bonferroni correction. The association of genes with each other or with different characteristics was analyzed using the Pearson correlation. PAST v3.0 was used to perform the statistical tests [29].

Results

Epidemiology of ESBL Producers

The overall prevalence of ESBL carriers was 3.7% (60/1,640). The LTC facility applicants carried ESBL producers (7.2%; 38/531) significantly more frequently (p < 0.001) than the individuals on employment screening (2.0%; 22/1,109). The risk of carriage of ESBL producers was not affected by the region where patients lived, by their gender or by their age in any of the comparisons.

Of the 63 ESBL-producing isolates from the 60 patients, 43 *E. coli*, 18 *K. pneumoniae*, 1 *Klebsiella oxytoca* and 1 *Proteus mirabilis* were identified. Three individuals harboured multiple ESBL-producing isolates simultaneously (*K. pneumoniae* and *E. coli* in 2 individuals and *K. oxytoca* and *E. coli* in 1 individual, all from the LTC applicant group). Among the individuals on employment screening, only *E. coli* isolates could be found, but the proportion of *E. coli* and *K. pneumoniae* isolates was similar (21/41 vs. 18/41) in the LTC facility applicants.

Out of the 63 ESBL-positive isolates, 50 harboured a bla_{CTX-M} gene; the majority of these were *E. coli* (38/50). The distribution of ESBL genes in the 2 groups is shown in figure 1. All isolates carrying bla_{CTX-M} other than $bla_{CTX-M-15}$ were *E. coli*; 1 *E. coli* isolate harboured $bla_{CTX-M-15}$ and $bla_{CTX-M-2}$ simultaneously. All *K. pneumoniae* and 5 *E. coli* isolates (11.6%) carried bla_{SHV-12}; in the case of *K. pneumoniae*, 11 isolates harboured $bla_{CTX-M-15}$ and bla_{SHV-12} at the same time. The distribution of the ESBL genes was significantly different (p = 0.001) between the 2 study groups; $bla_{CTX-M-15}$ was more frequent in the LTC applicants (p < 0.001), and the frequencies of other genes were comparable. When comparing ESBL gene distribution among *K. pneumoniae* (all from the LTC applicants), *E. coli* from the LTC applicants and *E. coli* from the people on employment screening, a significant difference was found (p < 0.001). All 3 isolate groups showed significantly different distributions in pairwise comparisons (p < 0.001 to p = 0.026), mostly due to the different distribution of $bla_{CTX-M-15}$. In addition, *K. pneumoniae* carried bla_{SHV-12} more frequently than *E. coli* among the LTC applicants and in the employment screening group (p < 0.001).

Genetic Diversity

The genetic diversity among *E. coli* isolates was high on both ERIC-PCR (data not shown) and PFGE (fig. 2; D = 0.99 and 0.96, respectively). One isolate was not
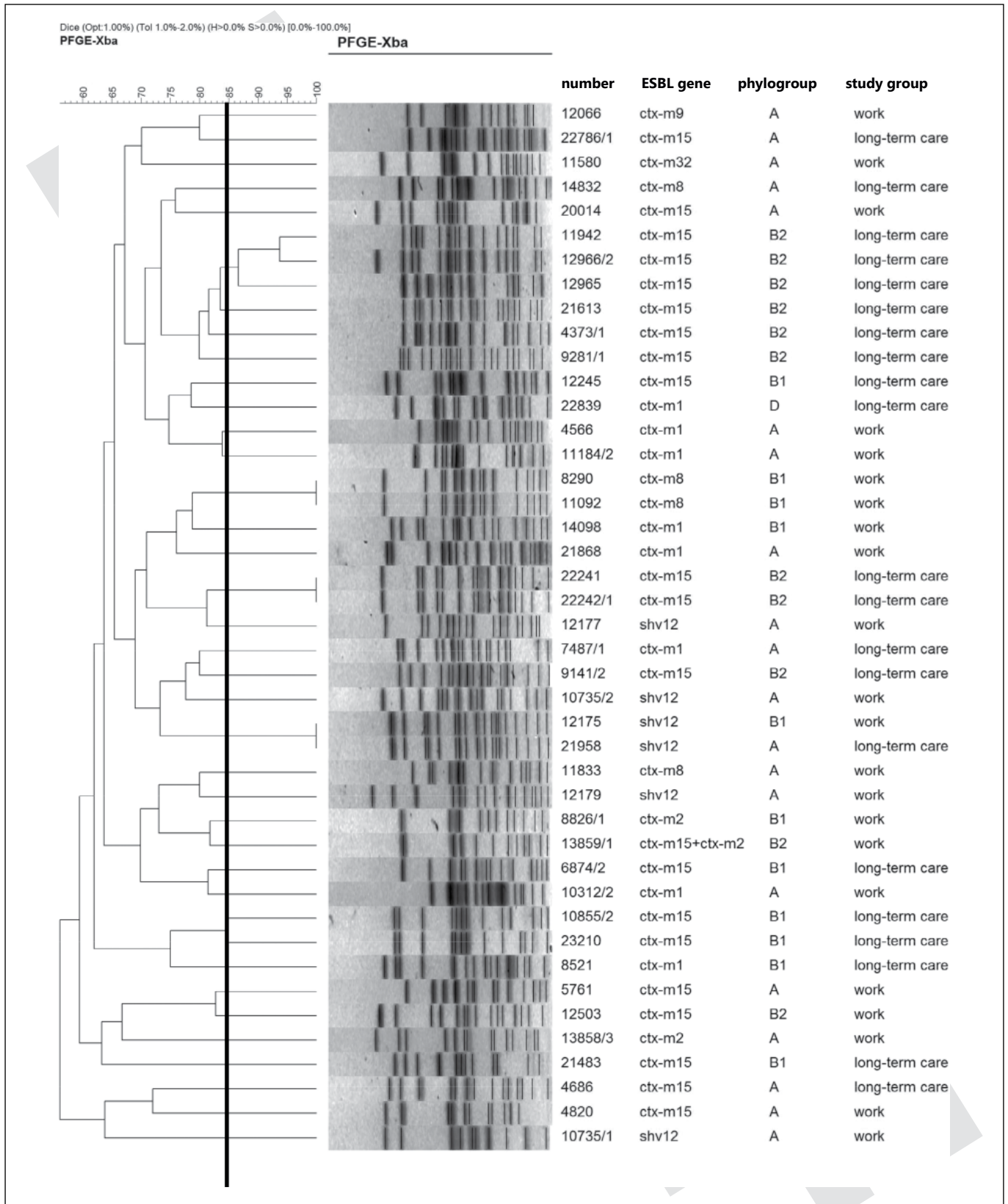
PFGE-Xba

number	ESBL gene	phylogroup	study group
12066	ctx-m9	A	work
22786/1	ctx-m15	A	long-term care
11580	ctx-m32	A	work
14832	ctx-m8	A	long-term care
20014	ctx-m15	A	work
11942	ctx-m15	B2	long-term care
12966/2	ctx-m15	B2	long-term care
12965	ctx-m15	B2	long-term care
21613	ctx-m15	B2	long-term care
4373/1	ctx-m15	B2	long-term care
9281/1	ctx-m15	B2	long-term care
12245	ctx-m15	B1	long-term care
22839	ctx-m1	D	long-term care
4566	ctx-m1	A	work
11184/2	ctx-m1	A	work
8290	ctx-m8	B1	work
11092	ctx-m8	B1	work
14098	ctx-m1	B1	work
21868	ctx-m1	A	work
22241	ctx-m15	B2	long-term care
22242/1	ctx-m15	B2	long-term care
12177	shv12	A	work
7487/1	ctx-m1	A	long-term care
9141/2	ctx-m15	B2	long-term care
10735/2	shv12	A	work
12175	shv12	B1	work
21958	shv12	A	long-term care
11833	ctx-m8	A	work
12179	shv12	A	work
8826/1	ctx-m2	B1	work
13859/1	ctx-m15+ctx-m2	B2	work
6874/2	ctx-m15	B1	long-term care
10312/2	ctx-m1	A	work
10855/2	ctx-m15	B1	long-term care
23210	ctx-m15	B1	long-term care
8521	ctx-m1	B1	long-term care
5761	ctx-m15	A	work
12503	ctx-m15	B2	work
13858/3	ctx-m2	A	work
21483	ctx-m15	B1	long-term care
4686	ctx-m15	A	long-term care
4820	ctx-m15	A	work
10735/1	shv12	A	work

Fig. 2. Dendrogram generated from macrorestriction patterns of the *E. coli* isolates.
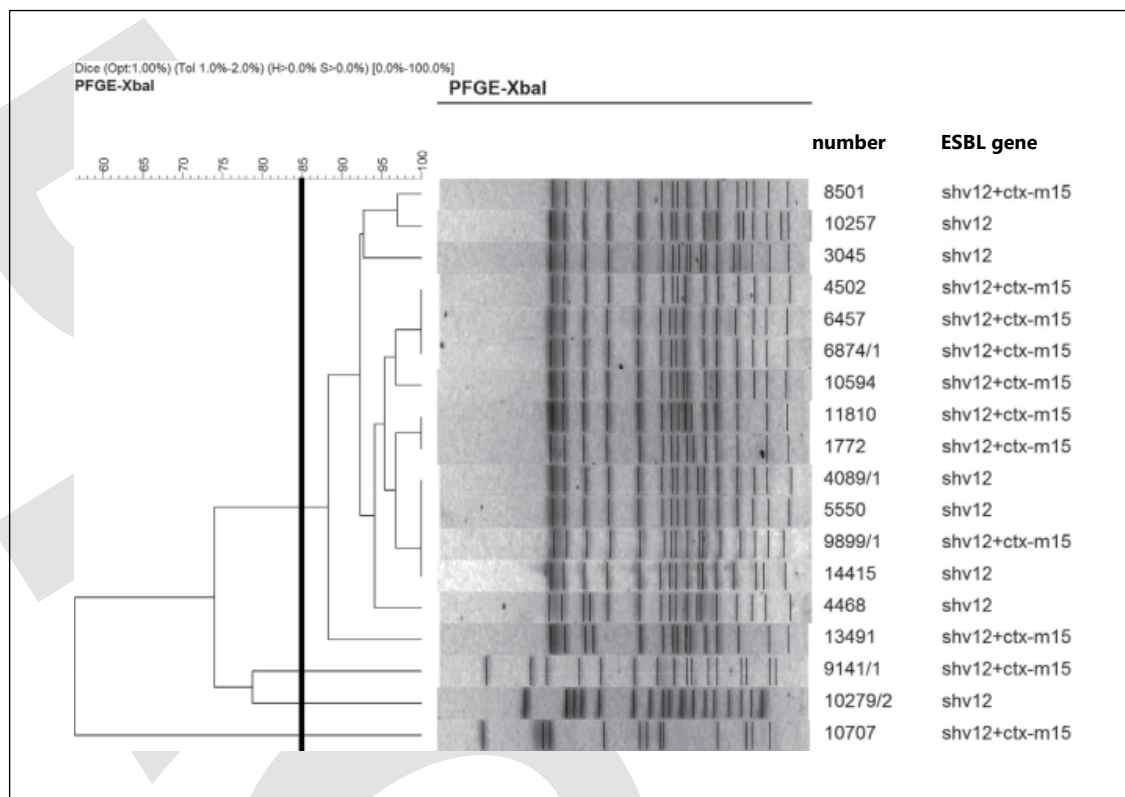
Ebrahimi/Mózes/Mészáros/Juhász/Kardos

Fig. 3. Dendrogram generated from macrorestriction patterns of *K. pneumoniae* isolates.

typeable with ERIC-PCR but its PFGE analysis was successful. The diversity of *E. coli* between the 2 groups was comparable (D = 0.99 vs. 0.99 and D = 0.94 vs. 0.95, respectively). Isolates in the same cluster were not uniform with regard to the ESBL genes and integrons carried.

In contrast, *K. pneumoniae* isolates were markedly less diverse, both with ERIC-PCR (data not shown) and PFGE (fig. 3; D = 0.58 and 0.31, respectively); a PFGE clone containing 15 of the 18 isolates was detected. Notably, 1 of the distinct isolates was the carbapenem nonsusceptible isolate.

Resistance Patterns and Aminoglycoside Resistance Genes

All ESBL-producing isolates were susceptible to colistin; carbapenem nonsusceptibility (Hodge test: negative) was detected in a single *K. pneumoniae* isolate harbouring $bla_{\text{SHV-12}}$. The *K. pneumoniae* clone detected was resistant to all the drugs tested, except for carbapenem and colistin, and it carried the *aac(6')-Ib* gene but no other tested aminoglycoside resistance genes.

As shown in figure 4, resistance rates significantly differed between *E. coli* isolates from individuals on employment screening and on screening for LTC admission in the case of ciprofloxacin, amikacin, tobramycin and trimethoprim-sulfamethoxazole (p = 0.048 to p < 0.002). When comparing commensal vs. pathogenic *E. coli* isolates as classified by phylogroups, the resistance pattern was similar; however, the significant differences in ciprofloxacin and trimethoprim-sulfamethoxazole disappeared. In contrast to the *K. pneumoniae* clone, isolates showing resistance to all non-beta-lactam antibiotic groups (except colistin) were not found in either *E. coli* group.

In the *E. coli* isolates, the presence of the $bla_{\text{CTX-M-15}}$ gene correlated positively with ciprofloxacin, amikacin and tobramycin resistance (table 1). The gene $bla_{\text{CTX-M-1}}$, on the contrary, correlated negatively with resistance to amikacin and tobramycin.

The prevalences of aminoglycoside resistance genes are shown in figure 5; all isolates were negative for *ant(2')-Ia*, *armA*, *rmtA* and *rmtB* genes. In the case of *E. coli* from the LTC applicants, aminoglycoside resistance genes were distributed unevenly (p = 0.017) with more frequent
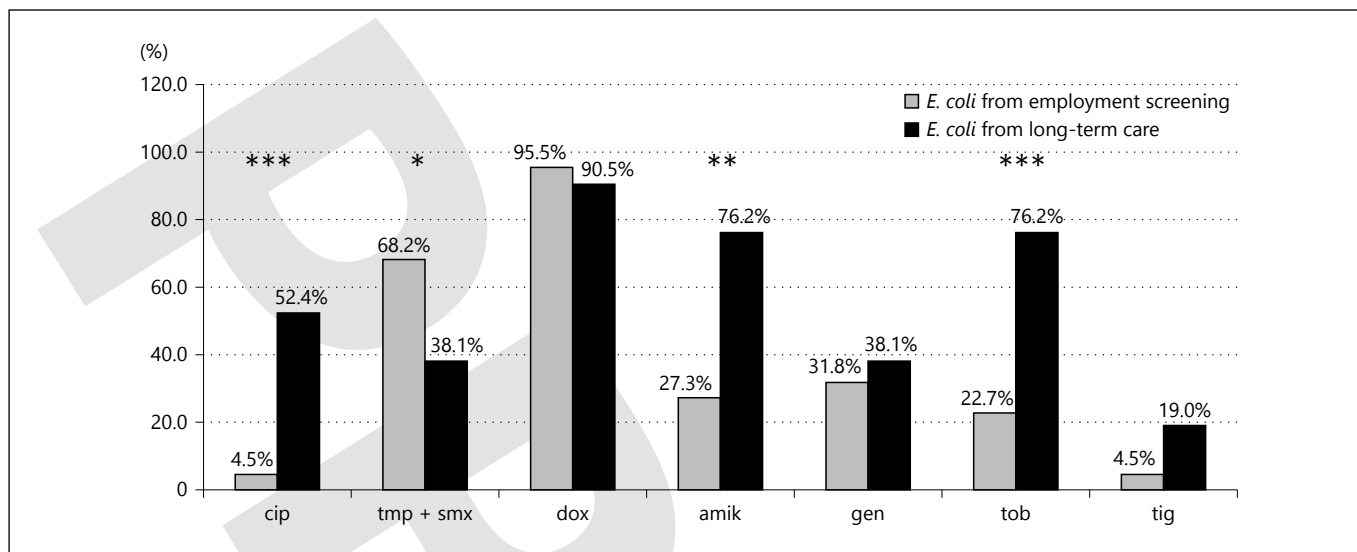
Fig. 4. Resistance rates in ESBL-producing isolates obtained from long-term care facility applicants and individuals on employment screening. amik = Amikacin; cip = ciprofloxacin; dox = doxycy- cline; gen = gentamicin; tig = tigecycline; tmp + smx = trime- thoprim-sulfamethoxazole (co-trimoxazole); tob = tobramycin. * p < 0.05; ** p < 0.01; *** p < 0.001.

Table 1. Correlation of carriage of $bla_{CTX-M-15}$ and $bla_{CTX-M-1}$ with resistance to antibiotics and aminoglycoside resistance genes in *E. coli* isolates

	$bla_{CTX-M-15}$	$bla_{CTX-M-1}$
Ciprofloxacin resistance	r = 0.43**	n.s.
Gentamicin resistance	n.s.	n.s.
Amikacin resistance	r = 0.58***	**r = −0.41****
Tobramycin resistance	r = 0.63***	**f = −0.39****
Tigecycline resistance	n.s.	n.s.
Carriage of *aac(3')-IIa*	r = 0.39*	n.s.
Carriage of *aac(6')-Ib*	r = 0.70***	**r = −0.40***
Carriage of *aph(3")-Ia*	**r = −0.35***	n.s.
Carriage of *ant(3")-Ia*	**r = −0.36***	n.s.
Carriage of class I integron	**r = −0.49*****	n.s.

n.s. = Not significant. Bold text represents negative correla- tions. * p < 0.05; ** p < 0.01; *** p < 0.001.

carriage of *aac(6')-Ib* than *aph(3')-Ia* and *ant(3')-Ia* (p = 0.048 for both comparisons), but not more frequent than the carriage of *aac(3')-IIa*. In the *E. coli* from the employ- ment screening group, the gene distribution was also un- even (p < 0.001), but *ant(3')-Ia* was more dominant than *aac(6')-Ib* and *aph(3')-Ia* (p = 0.005 and p = 0.02, respec- tively). When applying the phylogroup-based classifica- tion of *E. coli*, pathogenic isolates showed a pattern very similar to that of the isolates from the LTC applicants, but the dominance of *aac(6')-Ib* over *aph(3')-Ia* and *ant(3')-Ia* was more marked (p = 0.004 for both comparisons). The commensal isolates were similar to the isolates from the employment screening group.

The presence of the $bla_{CTX-M-15}$ gene in *E. coli* was as- sociated with the presence of *aac(3')-IIa* as well as *aac(6')-Ib*, and it correlated with the absence of *aph(3')-Ia* and *ant(3')-Ia* (table 1). The carriage of $bla_{CTX-M-1}$, in turn, was negatively correlated with *aac(6')-Ib*.

Integrons
The *K. pneumoniae* clone carried a class 1 integron with a single *ant(3')-1b* gene, which was also found in 2 of the 3 independent isolates. In *E. coli*, integron carriage correlated negatively with $bla_{CTX-M-15}$ (table 1). The car- riage rate of class 1 integrons was significantly higher in the *E. coli* from the employment screening group than from the LTC applicants (68.2%, 15/22 vs. 28.6%, 6/21; p = 0.01). Class 2 integrons were found in 6.3% (4/63) of the isolates (3 among the LTC facility applicants and 1 in an individual for employment screening); of these, 2 *E. coli* isolates harboured both integron types (1 isolate was from the LTC applicant group and the other from the employment screening group).

Seven and 2 different gene cassette arrays were identi- fied amongst the class 1 and 2 integrons, respectively (fig. 6). These arrays did not show any association with the ESBL gene carried or the study group of origin.
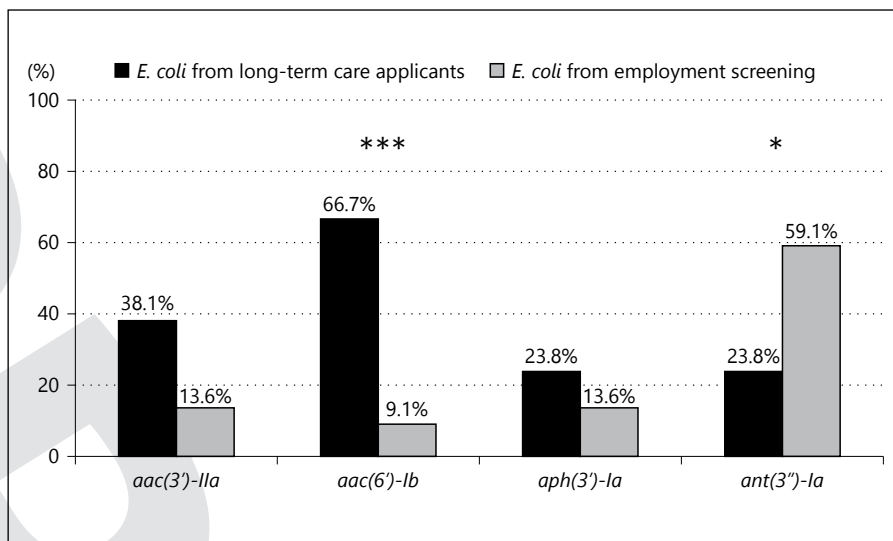
Fig. 5. The prevalence of aminoglycoside resistance genes in the 2 groups. * p < 0.05; *** p < 0.001.
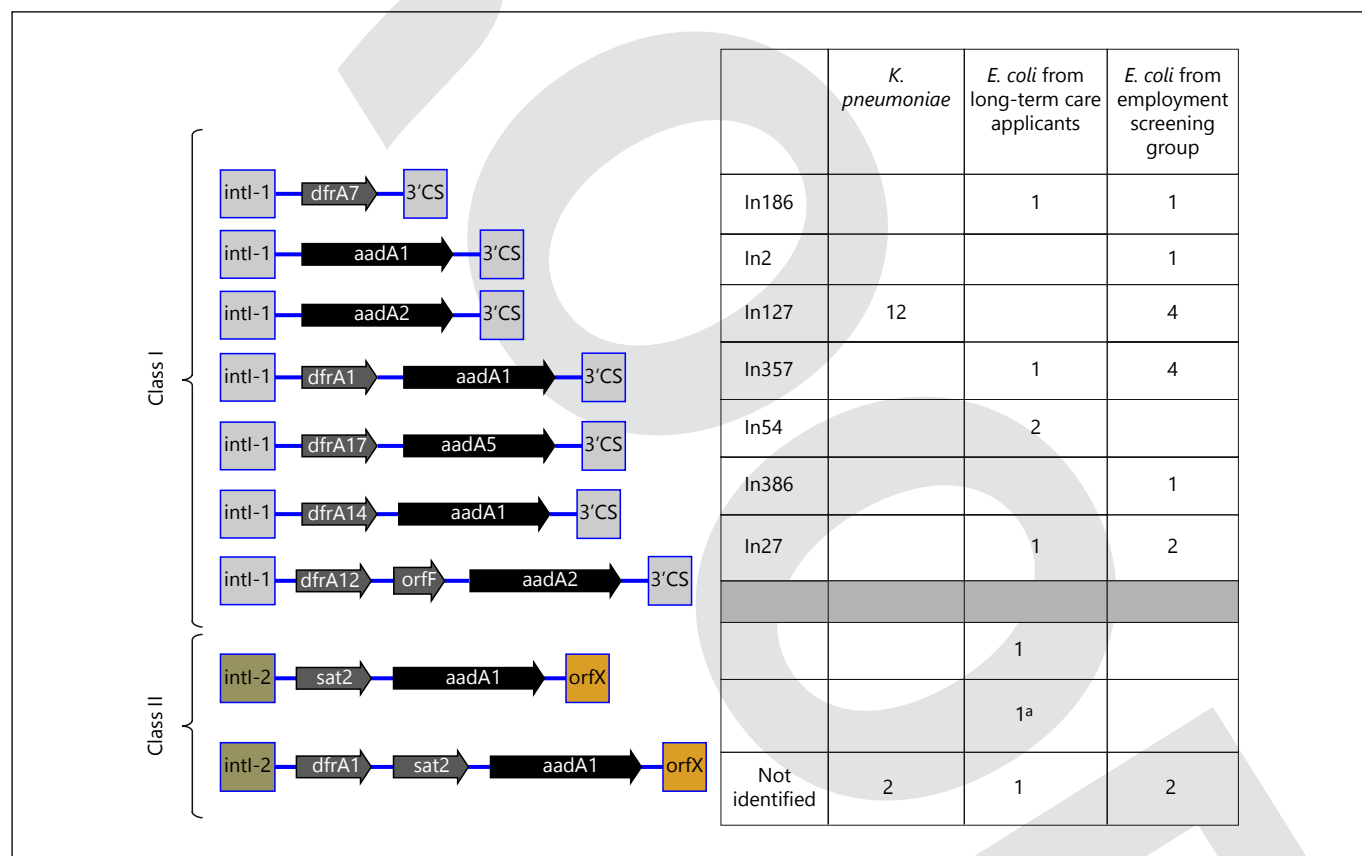

	K. pneumoniae	E. coli from long-term care applicants	E. coli from employment screening group
In186		1	1
In2			1
In127	12		4
In357		1	4
In54	2		
In386			1
In27		1	2
		1	
		1[a]	
Not identified	2	1	2

Fig. 6. Schematic representation of the various cassette arrangements found in class 1 and class 2 integrons. The arrows display different genes. The grey boxes indicate the 3′ and 5′ conservative segments. Integron numbers for class 1 integrons according to the Integrall database (http://integrall.bio.ua.pt/?nomenclature) are also shown. The number of isolates carrying each cassette array is shown on the right. [a] The *P. mirabilis* isolate also carried this array.

Table 2. Distribution of virulence genes according to *E. coli* phylogenetic groups

Gene	Number of positive isolates per phylogroup				
	A (n = 20)	B1 (n = 10)	B2 (n = 11)	D (n = 2)	Total (n = 43)
iutA	6	6	8	1	21 (48.8%)
papC	1	2	3	1	7 (16.3%)
kpsMT II	5	1	8	1	15 (34.9%)
fyuA	3	4	10	1	18 (41.9%)
sfaS	0	0	3	0	3 (7.0%)
cvaC	5	6	4	0	15 (34.9%)
traT	9	10	9	1	29 (67.4%)
cnf1	0	0	2	0	2 (4.6%)
ibeA	0	0	3	0	3 (7.0%)
sfa/focDE	0	0	3	0	3 (7.0%)
PAI	0	0	10	1	11 (25.6%)
fimH	15	10	11	2	38 (88.4%)
csgA	19	10	2	1	32 (74.4%)

Phylogrouping of E. coli and Analysis of Virulence Genes

Phylogenetic group analysis showed that group A covered 46.5% of strains, followed by groups B2 (25.6%), B1 (23.3%) and D (4.6%). The commensal *E. coli* phylogroup A was more frequent in the employment screening group (14/22 vs. 6/21; p = 0.02), but phylogroup B2 was more frequent in the LTC applicants (9/21 vs. 2/22; p = 0.01). All phylogroup B2 isolates belonged to the $bla_{CTX-M-15}$ producer O25b:ST131 pandemic clone.

None of the isolates belonged to the entero-virulent pathogroups for which tests were conducted. The frequency of the examined extra-intestinal virulence factor according to phylogroup is shown in table 2. Isolates belonging to pathogenic phylogroups showed a higher prevalence of the genes *kpsMT II* (p = 0.002) and *fyuA* (p < 0.001) than those from commensal phylogroups. The genes *sfaS*, *cnf1*, *ibeA*, *sfa/focDE* and PAI were completely absent from the commensal isolates. The comparison *E. coli* isolates according to source population yielded similar results, but only the prevalence of PAI was significantly different (p = 0.017). Virulence factor patterns were not linked to the type of ESBL gene carried.

Discussion

Asymptomatic carriage of ESBL-producing Enterobacteriaceae was documented as early as 1989 in hospital patients [30], and reports of carriage in healthy individuals soon followed [31, 32]. It became obvious that different geographical regions show different carriage rates in communities, ranging from 0.6 to 11.6% in Europe [6], with the highest prevalence being reported in LTC applicants from Belgium. Much higher rates have been found in other regions, e.g. 63.3% in Egypt [33] and 69.3% in Thailand [34]. This geographical difference is underlined by studies reporting international travel as a risk factor for the carriage of ESBL producers [8, 35, 36]. Faecal carriage rates seem to be increasing [6, 37, 38].

The prevalence rate found in this study among healthy individuals (those screened for employment purposes) is comparable or somewhat lower than rates reported by European studies [39, 40]. In contrast, the LTC applicants showed much higher prevalence, closer to that seen in hospitalized patients who were free of ESBL-related infections [41] a few years earlier and also that found in a contemporary study from Belgium targeting a similar patient group [42].

The distribution of the ESBL genes in the 2 study populations differed. In the *K. pneumoniae* and *E. coli* from the LTC applicants, it was similar to that in a European study on ESBL producers from human clinical samples [43], while higher gene diversity was found in the *E. coli* from the employment screening group.

Co-resistance patterns showed similarities with previous findings, e.g. the association of bla_{CTX-M} genes with ciprofloxacin and amikacin resistance [44, 45]. Co-resistance, as expected, was highest in *K. pneumoniae* [45], with a single, multi-resistant clone being dominant among the isolates. The co-resistance pattern of the 2 *E. coli* types was also very different. Resistance to ciprofloxacin, amikacin and tobramycin was significantly more prevalent in the LTC applicants, with the amikacin and tobramycin resistance being caused mainly by carriage of the *aac(6')-Ib* gene linked to $bla_{CTX-M-15}$ carriage. In agreement, it was also shown that ciprofloxacin resistance is associated with aminoglycoside resistance in ESBL-producer and non-ESBL producer *E. coli* [46].

In contrast, ESBL producers from the employment screening individuals were characterized by lower co-resistance rates, a low frequency of *aac(6')-Ib* and a high prevalence of *ant(3')-Ia*, a gene which is frequently present in the intestinal *E. coli* of healthy individuals [47]. This difference can be attributed to the high frequency of the ST131 carrying $bla_{CTX-M-15}$ in the LTC applicant group, while $bla_{CTX-M-1}$ carrier isolates, regardless of origin, were characterized by their low rate of co-resistance to other antibiotic classes and their lack of the *aac(6')-Ib* gene. In addition, several virulence factors were more frequently

found in the isolates from LTC applicants, also attributable to the higher frequency of ST131 in this group.

The differences in the distribution of the carried ESBL genes and other characteristics between the 2 populations in this study may be explained, in our opinion, by the different probable source of colonization. Most carriers in the group being screened for work eligibility presumably represent acquisition events independent of hospitals (rather community-related) whereas the group of LTC applicants probably represents mainly nosocomial acquisition. This serves as an explanation for the observations that *K. pneumoniae*, a typical nosocomial pathogen, was absent in the employment screening group, and that the *E. coli* isolates from the LTC applicants were more similar to *K. pneumoniae* than to *E. coli* isolates from the employment screening group with regard to a number of characteristics, including (1) a significantly lower diversity of the bla_{CTX-M} genes, (2) a significantly higher frequency of $bla_{CTX-M-15}$ and (3) the occurrence of aminoglycoside resistance genes *aac(6')-Ib* and *ant(3')-Ia*. This assumption is further supported by the lower occurrence of the commensal phylogroup A and the higher occurrence of the pathogenic phylogroup B2 in the *E. coli* from the LTC applicants and the frequent isolation of the pandemic ST131 clone from this group as well as by differences in the co-resistance patterns. Interestingly, these characteristics of *E. coli* were better explained by origin (i.e. LTC applicants vs. employment screening) than by phylogrouping (i.e. commensal vs. pathogenic).

These data, taken together, suggest that ESBL carriage in the community (employment eligibility group) is characterized by different colonization sources. The most obvious source is the exportation of hospital-derived strains either by colonized patients who have been discharged or by hospital workers; such events may be responsible for the introduction of the pathogenic ST131 clone carrying $bla_{CTX-M-15}$ which causes significant morbidity both in hospitals and in the community setting [48]. The source for colonization with isolates carrying bla_{SHV} may be horizontal gene transfer from exported *K. pneumoniae* isolates [49–51], while various potential environmental sources should also be considered for bla_{CTX-M} genes other than $bla_{CTX-M-15}$, including food [52, 53], companion animals [54, 55], wild animals [56, 57] surface water [58], drinking water [59] and food [52], either through direct transfer of the resistant bacteria or by the horizontal spread of resistance plasmids to human commensal *E. coli*.

This assumption is supported by the dominance of $bla_{CTX-M-1}$ among isolates from the work eligibility group and the parallel dominance of $bla_{CTX-M-1}$ in animal-derived Hungarian ESBL-producing *E. coli* isolates [53], in a Polish sample set from water birds [57] and in samples from food and food animals in Switzerland [52] as well as by the high frequency of $bla_{CTX-M-2}$ and $bla_{CTX-M-32}$ in samples originating in food animals [53, 60]. These *bla* genotypes were found exclusively in the individuals screened for work eligibility purposes in this study. The relatedness of animal and human isolates has also been demonstrated in some studies, pointing to a possible zoonotic spread of ESBL-producing *E. coli* [56, 61].

The presented data underline the complex epidemiology of ESBL producers in the community setting. The ESBL producers or ESBL-encoding genes may reach the human gut microbiota in the community setting via two main routes. Colonization may be the result of the exportation of nosocomial *K. pneumoniae* or *E. coli*, characterized by a dominance of $bla_{CTX-M-15}$, multiple co-resistance to other drug types and frequently by clonal spread. The other starting point may be the different environmental sources of mainly commensal *E. coli* characterized by diverse ESBL genes and a high diversity of genotypes of carrier bacteria as well as by a low rate of co-resistance. Though the extent of contribution to human infection by the latter group has still to be established, the increasing prevalence demonstrated by several authors [6, 62] is definitely cause for alarm.

Acknowledgements

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