Wide dissemination of extended-spectrum β-lactamase-producing Escherichia coli in community residents in the Indochinese peninsula

Abstract: Recent studies have reported a widespread distribution of extended-spectrum β-lactamase (ESBL)-producing bacteria, not only in the nosocomial setting, but also in the community; some local communities in Southeast Asia have been reported to show a high prevalence of ESBL-producing bacteria. However, the details regarding the quantitative/qualitative state of ESBL-producing bacterial spread in Southeast Asia are currently unclear. Thus, the aim of this study was to assess the state of ESBL-producing bacterial spread in community residents from the Indochinese peninsula, as a representative region of Southeast Asia. In order to achieve this aim, local community residents in Laos and Vietnam were examined for fecal carriage of ESBL-producing Enterobacteriaceae, and the findings were compared with data from a previous study in Thailand which was conducted in the same manner as this study. Between 47.0%–70.2% of the Laotian and Vietnamese residents carried ESBL-producing CTX-M genotype Enterobacteriaceae. The most common sub-genotypes of CTX-M were CTX-M-1 (33.0%–47.5%) and CTX-M-9 (47.5%–64.1%), and these rates were similar among all three countries. Taken together, these results confirmed that ESBL-producing Enterobacteriaceae are widely disseminated in Indochinese countries, such as Thailand, Laos, and Vietnam.

Keywords: extended-spectrum β-lactamase (ESBL), rural residents, Indochinese peninsula, fecal carriage

Introduction

Multi-drug resistant bacteria are associated with persistent disease and are continuously expanding around the world, warranting increased attention.1 Extended-spectrum β-lactamases (ESBLs) are enzymes produced by Enterobacteriaceae, including Escherichia coli and Klebsiella pneumoniae, which invalidate β-lactam antibiotics, such as penicillin and cephalosporin.2 Recent studies have indicated a high prevalence of these ESBL-producing bacteria in communities in various Asian countries,3–6 and our previous studies revealed a relatively high prevalence of ESBL-producing Enterobacteriaceae in residents of local communities in Thailand.3,4 However, whether this high prevalence of ESBL-producing bacteria in the community is similar in other communities of the Indochinese peninsula remains to be elucidated. Therefore, in this study, two other communities from Laos and Vietnam were studied to determine the quantitative and qualitative states of ESBL-producing bacterial spread in this part of the world.

Materials and methods

Specimen collection

The study was conducted at a village in the Sawanaket province, Laos, in November 2012 and a community in Bavi, Hanoi, Vietnam, in June 2013. The two communities were
selected to represent geographically different countries of the peninsula (Figure 1). A total of 57 and 198 asymptomatic resident volunteer participants were enrolled in Laos and Vietnam, respectively. One stool specimen was obtained from each participant using a transport swab with Cary-Blair transport media (Eiken Chemical, Tokyo, Japan). Written informed consent was obtained from each individual participating in the study. The study was approved by the ethics committee of Osaka University.

Detection of ESBL-producing bacteria
Stool specimens were plated on MacConkey agar (Nissui, Tokyo, Japan) supplemented with 1 and 2 mg/L cefotaxime (Tokyo Chemical Industry, Tokyo, Japan) and incubated at 37°C for 24 hours. ESBL expression was confirmed by the disc diffusion method using cefotaxime and ceftazidime with and without clavulanic acid, as recommended by the Clinical and Laboratory Standards Institute (CLSI, Wayne, PA, USA); each set of samples was tested along with the CLSI quality control strains E. coli (ATCC 700603) and K. pneumoniae (ATCC 25922 and K. pneumoniae ATCC 700603).4 Positive isolates were identified using conventional biochemical tests with the API 20E (Sysmex-bioMerieux, Tokyo, Japan) and VITEK 2 (Sysmex-bioMerieux, Tokyo, Japan) system.4

Detection of ESBL genes and genotyping
Bacterial DNA was extracted by boiling suspensions of ESBL-producing isolates. DNA samples at a concentration of 1.0 ng/µL were used as polymerase chain reaction (PCR) templates, and the blaCTX-M, blaSHV, and blaTEM genes were amplified using universal primers, as described previously.9 For genotyping of the blaCTX-M genes, four primer sets were used to amplify group-specific blaCTX-M genes, as described elsewhere,10 including the CTX-M-1, CTX-M-2, CTX-M-8, CTX-M-9, and CTX-M-25 groups. Phylogenetic grouping of identified E. coli was determined by triplex PCR using a combination of two genes (chuA and yjaA) and the DNA fragment TSPE4.C2, as described elsewhere.11 The PCR products were visualized by 2% agarose gel electrophoresis and staining with GelRed nucleic acid stain (Biotium, Hayward, CA, USA).

Statistical analysis
Statistical analysis was performed using the chi-squared test.

Results
A total of 57 Laotian and 198 Vietnamese local residents were investigated for the prevalence of ESBL-producing Enterobacteriaceae. The results showed that 71.9% of the Laotian residents and 51% of the Vietnamese residents had ESBL-producing Enterobacteriaceae in their stool specimens. All of the isolated ESBL-producing Enterobacteriaceae were identified as E. coli. The multiplex PCR detected that 70.2% of Laotian and 47% of Vietnamese residents possessed ESBL-producing bacteria with the CTX-M gene. The prevalence of ESBL-producing Enterobacteriaceae in the community assessed in Vietnam was significantly lower than that in Laos and Thailand (Table 1). However, the

Table 1 Prevalence of extended-spectrum β-lactamase-producing Enterobacteriaceae in residents of Indochinese countries

<table>
<thead>
<tr>
<th></th>
<th>Thailand¹</th>
<th>Laos</th>
<th>Vietnam</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sampling period</td>
<td>November 2010</td>
<td>November 2012</td>
<td>June 2013</td>
</tr>
<tr>
<td>No of participants</td>
<td>417</td>
<td>57</td>
<td>198</td>
</tr>
<tr>
<td>Age, mean</td>
<td>47</td>
<td>47</td>
<td>31</td>
</tr>
<tr>
<td>Age, range</td>
<td>20–85</td>
<td>17–75</td>
<td>1–90</td>
</tr>
<tr>
<td>Sex, male (%)</td>
<td>38</td>
<td>18</td>
<td>52</td>
</tr>
<tr>
<td>Phenotype</td>
<td>69.3% (289)⁴</td>
<td>71.9% (41)⁴</td>
<td>51.0% (101)</td>
</tr>
<tr>
<td>CTX-M gene</td>
<td>65.7% (274)⁴</td>
<td>70.2% (40)⁴</td>
<td>47.0% (93)⁴</td>
</tr>
<tr>
<td>CTX-M genotype groups</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M-1 group</td>
<td>38.7% (106)</td>
<td>47.5% (19)</td>
<td>33.0% (34)</td>
</tr>
<tr>
<td>M-2 group</td>
<td>0% (0)</td>
<td>0% (0)</td>
<td>1.0% (1)</td>
</tr>
<tr>
<td>M-8 group</td>
<td>0.7% (2)</td>
<td>5.0% (2)</td>
<td>1.9% (2)</td>
</tr>
<tr>
<td>M-9 group</td>
<td>60.6% (166)</td>
<td>47.5% (19)</td>
<td>64.1% (66)</td>
</tr>
</tbody>
</table>

Notes: Luvsansharav UO, Hirai I, Nakata A, et al. Prevalence of and risk factors associated with faecal carriage of CTX-M beta-lactamase-producing Enterobacteriaceae in rural Thai communities. J Antimicrob Chemother 2012;67(7):1769–1774 by permission of Oxford University Press.¹the number of CTX-M gene isolates was not matched with the total number of CTX-M due to the presence of dual CTX-M gene possession. The numbers in parentheses indicate the number of strains. P<0.0001, significantly different from the Vietnam group. P<0.01, significantly different from the Vietnam group.
majority of ESBL-producing Enterobacteriaceae isolated in both Laos and Vietnam harbored CTX-M type genes (more than the 92% of ESBL phenotype strains) assessed by the multiplex PCR.

The genotyping of CTX-M gene positive isolates showed that both the CTX-M-1 (47.5% and 33%) and -9 group (47.5% and 64.1%) were the most predominant genotype. While the CTX-M-2 (0% and 1.0%) and -8 group (5.0% and 1.9%) were the lowest rates in the Laotian and Vietnamese residents, respectively (Table 1).

The phylogenetic groups of the CTX-M ESBL-producing E. coli isolates were also determined. The most frequently detected phylogenetic group was A (50.0% and 38.6%) followed by D (25% and 25.7%) in the Laotian and Vietnamese residents, respectively. The least detected phylogenetic groups were the B1 and B2 groups (12.5%) in Laotian residents and the B2 group (13.9%) in Vietnamese residents (Table 2).

Discussion

Worldwide, the prevalence of ESBL-producing bacteria in the clinical setting has been well studied. However, the microbiological state of ESBL-producing bacteria in the community setting, particularly in Southeast Asia, is unclear, even though significant increases in the frequency of ESBL-producing bacteria in community isolates have been reported. On the other hand, the observation of CTX-M type ESBL-producing bacteria in community strains is consistent with their emergence from a widespread reservoir. In this regard, our previous study revealed the high prevalence of CTX-M type ESBL-producing Enterobacteriaceae in a local community in Thailand. However, it is still unclear whether this high prevalence of ESBL-producing bacteria can be found in other Southeast Asian countries as well. Therefore, in the present study, two local communities in Laos and Vietnam, which are geographically different, were chosen as representative communities on the Indochinese peninsula.

The study protocol and conditions of this present study were similar to those in our previous study performed in Thailand. While the age and sex of the participants differed among the three countries, this was unlikely to affect the prevalence of ESBL-producing bacteria, as the effects of age and sex on the risk of ESBL-producing bacteria carriage are reportedly limited. The concentration of cefotaxime used in the selection medium was different between the two communities. Our preliminary study with 2 mg/L cefotaxime in Vietnam showed a relatively low prevalence in comparison with that of Laos and Thailand. Therefore, we reduced the concentration of cefotaxime to 1 mg/L, which is recommended for screening ESBL-producing bacteria by CLSIF to detect more ESBL-producing bacteria than with 2 mg/L. As shown in Table 1, the prevalence of ESBL-producing Enterobacteriaceae at 1 mg/L cefotaxime was still relatively low in Vietnam (ESBL phenotype, 51%; CTX-M gene, 47%) compared with the prevalence rates in Laos and Thailand. Nevertheless, the present study revealed a high prevalence of ESBL-producing bacteria in the residents of the two communities in Laos and Vietnam.

The majority of ESBL genes were CTX-M, accounting for 97.5%, 92.1%, and 94.8% of ESBL-producing Enterobacteriaceae isolated in Laos, Vietnam, and Thailand, respectively. The widespread use of cefotaxime or ceftriaxone has been proposed as a reason for the emergence of CTX-M enzymes. The CTX-M subclasses were also investigated for characterization of the ESBL-producing bacterial dissemination in the community. Worldwide, the CTX-M-1 (CTX-M-15) group has been reported to be the dominant group of CTX-M ESBL-producing bacteria, followed by the CTX-M-9 (CTX-M-14) group, as determined by the Study for Monitoring Antimicrobial Resistance Trends using clinical isolates; the results of the present study corresponded with these previous findings.

It has been pointed out that the use of antibiotics in agriculture may contribute to the spread of antibiotic resistance in farm animals and subsequently result in transmission to humans. In fact, in the communities assessed in this study, household subsistence farming was common, indicating that the residents were likely in close contact with farm animals. In this regard, a recent study carried out in Korea reported that the most predominant CTX-M-6 type in farm animals, the farm environment, and farm workers was CTX-M-9, followed by CTX-M-1. The reasons for this observed dominance of CTX-M-1 and CTX-M-2 in the community and agriculture settings are not clear. However, clonal expansion of CTX-M ESBL-producing bacteria, such as E. coli ST131 with CTX-M-15 (CTX-M-1 group), or horizontal dissemination of certain plasmids carrying the CTX-M gene may represent potential mechanisms.

Table 2 Phylogenetic groups of the CTX-M extended-spectrum β-lactamase-producing Escherichia coli isolates

<table>
<thead>
<tr>
<th>Phylogenetic group</th>
<th>Thailand</th>
<th>Laos</th>
<th>Vietnam</th>
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<tbody>
<tr>
<td></td>
<td>No of isolates</td>
<td>%</td>
<td>No of isolates</td>
</tr>
<tr>
<td>A</td>
<td>114</td>
<td>48.7</td>
<td>20</td>
</tr>
<tr>
<td>B1</td>
<td>67</td>
<td>28.6</td>
<td>5</td>
</tr>
<tr>
<td>B2</td>
<td>16</td>
<td>6.8</td>
<td>5</td>
</tr>
<tr>
<td>D</td>
<td>37</td>
<td>15.8</td>
<td>10</td>
</tr>
<tr>
<td>Total</td>
<td>234</td>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

Assessment of the phylogenetic groups of the CTX-M ESBL-producing bacteria isolated showed dominance of A followed by D in Laotian and Vietnamese residents. Similarly, in the People’s Republic of China, phylogenetic groups A and D have been reported to be prevalent among ESBL-producing E. coli strains in asymptomatic carriers. On the other hand, a report on the population genetics of E. coli stated that commensal E. coli strains isolated from Asia mainly belonged to phylogenetic groups A and B1, whereas other studies have reported that a recent worldwide dissemination of CTX-M-14 and CTX-M-15 was likely driven by epidemic E. coli strains belonging to phylogenetic group B2 (ST131). Even though multi locus sequence typing was not conducted in this study, since the number of ESBL-producing bacteria belonging to B2 was limited in these communities, a possible clonal outbreak of CTX-M ESBL-producing E. coli B2 ST131 was considered unlikely.

Despite a limited number of participants, the results obtained in this study showed a wide dissemination of ESBL-producing bacteria among community residents in the Indochinese peninsula. These residents may be reservoirs for antibiotic resistance in the community and may contribute to the shortfalls in infection control and public health management.

Acknowledgments
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Disclosure
The authors report no conflicts of interest in this study.

References