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Epidemiological Investigation of *Salmonella enterica* Serovar Kedougou in Thailand

Srirat Pornruangwong,1 Rene S. Hendriksen,2 Chaiwat Pulsrikarn,1 Aroon Bangstrakulnonth,3 Matthew Mikoleit,4 Rob H. Davies,5 Frank M. Aarestrup,2 and Lourdes Garcia-Migura2

Abstract

**Objective:** Salmonella enterica serovar Kedougou is among the top 10 serovars reported in northern Thailand. The objective of this study was to identify risk factors associated with *Salmonella* Kedougou infection in Thailand and to compare the molecular types and antimicrobial resistance with *Salmonella* Kedougou isolates of human origin from United States and of animal origin from the United Kingdom.

**Methods:** Data from 13,976 *Salmonella* infections of which 253 were *Salmonella* Kedougou collected in Thailand between 2002 and 2008 were analyzed by logistic regression. Antimicrobial susceptibility testing and pulsed-field gel electrophoresis (PFGE) were performed on selected *Salmonella* Kedougou strains causing infections in Thailand (n = 66), and compared to isolates from the United States (n = 5) and the United Kingdom (n = 20).

**Results:** Logistic analysis revealed season (hot/dry; p = 0.023), region (northern Thailand; p < 0.001), and specimen (stool; p < 0.001) as significant risk factors associated with *Salmonella* Kedougou infection compared to other nontyphoid *Salmonella*. Of the *Salmonella* Kedougou isolates of human origin, 84% exhibited resistance to at least three antimicrobial classes. Three strains recovered from human stool in Thailand were resistant to third-generation cephalosporins: two harbored *blaCTX-M-63* and one *blaCMY-2*. PFGE revealed 45 unique clusters. Isolates obtained from humans in Thailand and the United States presented identical PFGE profiles suggesting a travel association, whereas the majority of the animal isolates from United Kingdom clustered separately.

**Conclusions:** This study reveals *Salmonella* Kedougou as a major cause of human infections in northern Thailand especially during the hot period and suggests a global spread probably due to travel. The clonal types causing infections in humans differed from those observed in animals in United Kingdom, which suggests the absence of an epidemiological link and could suggest differences in virulence. The high frequency of antimicrobial resistance, including emergence of resistance to fluoroquinolones and third-generation cephalosporins, might pose problems for treatment of infections.

Introduction

Although over 2500 serovars of *Salmonella enterica* have been identified (Guibourdenche et al., 2010), most human infections are caused by a limited number of serovars, with *Salmonella enterica* serovars Typhimurium and Enteritidis being the most common causes of salmonellosis worldwide (Galanis et al., 2006). In industrialized countries, most cases of salmonellosis are due to exposure to contaminated animal products. In developing countries, contaminated vegetables, water, and human-to-human transmission are also believed to contribute to a comparatively larger proportion of the human cases than those in industrialized countries (Wegener et al., 2003; Okeke et al., 2007).

There are only a few reports of human infections caused by *Salmonella* Kedougou. Outbreaks have been attributed to the consumption of products such as salami (Emberland et al., 2006), commercially grown mushrooms (Doran et al., 2005),

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turkey meat (Salmonella Kedougou and cooked meats, 1992), and infant formula (Soler et al., 2008). In addition, the first report of Salmonella Kedougou producing extended-spectrum β-lactamases (ESBL) was described from a hospital strain isolated in Algeria in 2007 (Touati et al., 2008). Although Salmonella Kedougou does not appear to be an invasive serovar, the emergence of antimicrobial resistance in Salmonella Kedougou may pose a threat to human health, especially if these strains acquire resistance to third-generation cephalosporins.

Cyprus and the United Kingdom reported Salmonella Kedougou as one of the 10 most common Salmonella serovars isolated from pig carcasses during a baseline survey performed in Europe between 2006 and 2007 (EFSA, 2009). This serovar has also been described in broiler farms in the United Kingdom mainly associated with feed mills (Snow et al., 2008) and turkey fattening holdings (www.defra.gov.uk/corporate/consult/turkey/turkey-ncp.pdf). In addition, an increase in human cases associated with the presence of Salmonella Kedougou was reported by the International Surveillance Network for the Enteric Infections—Salmonella, verotoxin-producing Escherichia coli (VTEC) (VTEC) 0157, and Campylobacter during 2006 (www.hpa.org.uk/web/HPAwebFile/HPAweb_C/1194947410 123). Out of 75 Salmonella Kedougou cases registered in six different European countries, four cases were confirmed to be travel related. Three cases confirmed travel to Thailand and the fourth case reported travel to an unspecified location. In the United States, 19 cases of Salmonella Kedougou causing infections in humans were reported between 1996 and 2006 (www.cdc.gov/ncidod/dbmd/phlsdata/salmtab/2006/ SalmonellaAnnualSummary2006.pdf).

Recent studies conducted in Thailand have identified a large number of human infections caused by Salmonella Kedougou (Hendriksen et al., 2009). In the northern region of Thailand, this serovar was described as the 10th most common serovar and accounting for 5% of the total cases of salmonellosis reported in this region between 2002 and 2008 (Hendriksen et al., 2009). Salmonella Kedougou was also detected in other regions of Thailand although at much lower rates, counting for 1.8% of the Salmonella infections reported in Thailand during the same period. Studies carried out in retail meat in Thailand have also identified Salmonella Kedougou as one of the most common serovars isolated from fresh pork for human consumption (Vindigini et al., 2007). Moreover, multidrug-resistant Salmonella Kedougou containing class 1 integrons and Salmonella genomic islands have recently been reported in Thailand (Khemtong and Chuanchuen, 2008). In a previous survey covering 1993 to 2002, this serovar was rarely detected (Bangtrakulnonth et al., 2004); therefore, the increasing proportions of Salmonella Kedougou infections appear to represent an emerging problem.

Specific risk factors for infection with Salmonella Kedougou in Thailand or any association to isolates found in Europe and the United States have not been identified. Thus, the objective of this study was to evaluate the different variables such as age, season, gender, and geographical location as possible risk factors associated with infection with this serovar compared to other nontyphoidal Salmonellae. This knowledge may facilitate the recognition and control of Salmonella Kedougou as a new and emerging pathogen in the affected areas of Thailand. Further, by identifying the risk factors associated with Salmonella Kedougou, it may be possible to establish intervention measures to reduce infection due to the presence of this serovar. In addition, the clonality and antimicrobial resistance profile of isolates from Thailand was assessed. For logistical reasons, the authors were not able to obtain Salmonella Kedougou isolates from the animal reservoir in Thailand; therefore, the clonality of Salmonella Kedougou isolates of human origin from Thailand was compared to Salmonella Kedougou isolated from humans in the United States and from various nonhuman reservoirs in the United Kingdom.

Materials and Methods

Study design

This study is a cross-sectional study carried out during a 7-year period (2002–2008). The data were collected by the different regional medical centers in Thailand through passive surveillance recording only confirmed cases of salmonellosis. All data from the different regional centers were merged at the WHO Salmonella and Shigella center in Bangkok.

Statistical analysis

SAS version 9.1.3 (SAS Institute Inc., Cary, NC) was used to perform the statistical analysis. The dataset contained a total of 13,976 Salmonella cases collected during 2002 to 2008 from blood and fecal samples (rectal swabs or stool). Due to the low number of isolates, human clinical strains isolated from other specimen types such as pus or urine were excluded from the study. Originally, age groups were given in intervals of 5 years. These data were later aggregated into five levels, 0–5, 6–20, 21–40, 41–60, and >60 years. Similarly, regions of Thailand previously labeled in the database from number 1 to 13 were also aggregated into five larger regions according to the national regions of Thailand: central, northeast, south, north, and Bangkok. Finally, a new variable named “season” was created as the outcome of grouping the months into two seasons corresponding to rainy season (from July to October) and hot/dry season (from November to June).

A preselection of independent variables, age group, season, gender, specimen, and geographical location was initiated using univariable analysis. All independent variables with a p-value of <0.05 were included in the logistic analysis. However, interactions between the independent variables were outside the scope of the analysis. Backward selection of variables was performed using p-values. The criteria for keeping variables in the model were p-value of <0.05. Possible confounding effects between variables were also assessed.

Strain collection

From Thailand, a total of 66 Salmonella Kedougou isolates of human origin were selected for further analysis. They represented isolates from all regions of Thailand obtained from stool and rectal swabs during 2008.

Isolates from the United Kingdom were obtained from an active surveillance program that monitors the prevalence of Salmonella spp. in various animal production lines. During 2007, there were a total of 2244 Salmonella submissions of which 90 were confirmed to be Salmonella Kedougou. During 2008, the number of submissions increased to 2444 and the confirmed cases of Salmonella Kedougou also increased to 124. Twenty isolates collected during this 2-year period were selected manually to represent the different production lines, as they were too many isolates from the same or related sources.
that they would have predominated in a random selection (Table 3). Five *Salmonella* Kedougou isolates from human stool (n = 4) and urine (n = 1) collected in the United States were also included in the study.

**Serotyping**

Serotyping of all the presumptive biochemical-positive *Salmonella* isolates was performed according to previously described methods (Le et al., 1990). All human *Salmonella* Kedougou isolates recovered from Thailand were serotype in the WHO International *Salmonella* and *Shigella* Centre, National Institute of Health, Bangkok, Thailand. The animal isolates were serotype at the Veterinary Laboratories Agency, Weybridge, United Kingdom. Finally, the five isolates from the United States were serotype at the Centre for Disease Control and Prevention, Atlanta, GA.

**Antimicrobial susceptibility testing**

Susceptibility to 17 antimicrobial agents was performed at the National Food Institute, Copenhagen, Denmark (DTU-FOOD), on all *Salmonella* isolates as minimum inhibitory concentration (MIC) determinations according to previously described methods (Hendriksen et al., 2008). For cefotaxime, a clinical breakpoint of >2 mg/L was used (www.eucast.org).

**Pulsed-field gel electrophoresis**

To assess the clonality of the strains and the epidemiological relatedness, *XbaI*- pulsed-field gel electrophoresis (PFGE) was conducted as described by the Centre for Disease Control and Prevention (CDC) PulseNet protocol (Ribot et al., 2006). The *Salmonella* Braenderup H9812 strain was used as molecular standard. PFGE profiles were compared using Bionumerics software version 4.6 (Applied Maths, Sint-Martens-Latem, Belgium). Isolates were considered to have a unique pattern when at least one band difference was detected. The analysis of the bands generated was performed using the Dice coefficient and unweighted pair group method with arithmetic averages (optimization of 1.00% and position tolerance 1.5%).

**Antimicrobial-resistant determinants**

The three strains exhibiting resistance to third-generation cephalosporins were further characterised by polymerase chain reaction for the presence of the *bla*TEM, *bla*CTX, *bla*CMY-1, *bla*CMY-2, *bla*SHV, and *bla*ACC genes as described previously (Hasman et al., 2005). Sequence analysis was performed using Vector NTI advance 11 (InforMax, Inc., Bethesda, MD). The amplified nucleotide sequences were compared to previously described sequences obtained from public databases (www.ncbi.nlm.nih.gov/and www.lahey.org/studies/webt.html).

**Results**

**Descriptive data**

A total of 13,976 *Salmonella* isolates were collected in Thailand between 2002 and 2008. Of those, 253 were identified as *Salmonella* Kedougou and 13,723 as other nontyphoid *Salmonella* (NTS) serovars. The authors have not received any information to suggest that these data are biased due to the occurrence of a local or regional outbreak. Information on the number of hospitals reporting to each of the regional reference centers or efficacy at reporting was not available; therefore, clustering by geographical location could not be performed.

The overall percentage of *Salmonella* Kedougou cases versus NTS during the study period is given in Table 1. The proportion of this particular serovar in the northern region fluctuated significantly during the years, with 0% cases of *Salmonella* Kedougou registered in 2002 followed by an increase to 8% in 2005 and subsequent decrease to 6% and 3.3% in 2006 and 2008, respectively (Fig. 1).

The age groups with the highest incidence of *Salmonella* Kedougou cases were children from 0 to 5 years of age (94/13,976) followed by adults of age between 21 and 40 (58/13,976). Conversely, the group age between 6 and 20 years of age presented the lowest proportion of infections (18/13,976). Same trends followed the incidence rates of other NTS cases, with similar number of infections recorded from males and females (Table 1).

The dry season was the period with the highest proportions of *Salmonella* Kedougou cases. Conversely, infections with other NTS reached the highest rates during the rainy season. The majority of *Salmonella* Kedougou infections were reported from the northern region of Thailand (103/253) followed by the central region (76/253). The northeast region of Thailand was the area with the lowest proportions of *Salmonella* infections during the 7-year period, with only 1309 (9.4%) cases of the 13,976 reported in this dataset. During the study period, isolates were primarily recovered from stool samples and very rarely from blood (Table 1).

**Table 1. Descriptive Analysis of the Proportions of *Salmonella* Kedougou and Nontyphoid *Salmonella* Found for Each One of the Individual Variables**

<table>
<thead>
<tr>
<th>Variables</th>
<th>Levels</th>
<th><em>Salmonella</em> Kedougou (%)</th>
<th>Nontyphoid <em>Salmonella</em> (%)</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall Year</td>
<td>2002</td>
<td>13 (0.7)</td>
<td>1927 (99.3)</td>
<td>1940</td>
</tr>
<tr>
<td></td>
<td>2003</td>
<td>2 (0.1)</td>
<td>1421 (99.9)</td>
<td>1423</td>
</tr>
<tr>
<td></td>
<td>2004</td>
<td>23 (1.2)</td>
<td>1870 (98.8)</td>
<td>1893</td>
</tr>
<tr>
<td></td>
<td>2005</td>
<td>47 (2.0)</td>
<td>2315 (98.0)</td>
<td>2362</td>
</tr>
<tr>
<td></td>
<td>2006</td>
<td>58 (2.8)</td>
<td>1999 (97.2)</td>
<td>2057</td>
</tr>
<tr>
<td></td>
<td>2007</td>
<td>50 (2.5)</td>
<td>1931 (97.5)</td>
<td>1981</td>
</tr>
<tr>
<td></td>
<td>2008</td>
<td>60 (2.6)</td>
<td>2260 (97.4)</td>
<td>2320</td>
</tr>
<tr>
<td>Age group</td>
<td>0–5</td>
<td>94 (2.0)</td>
<td>4525 (98.0)</td>
<td>4619</td>
</tr>
<tr>
<td></td>
<td>6–20</td>
<td>18 (1.2)</td>
<td>1434 (98.8)</td>
<td>1452</td>
</tr>
<tr>
<td></td>
<td>21–40</td>
<td>58 (1.6)</td>
<td>3501 (98.4)</td>
<td>3559</td>
</tr>
<tr>
<td></td>
<td>41–60</td>
<td>47 (2.0)</td>
<td>2302 (98.0)</td>
<td>2349</td>
</tr>
<tr>
<td></td>
<td>&gt;60</td>
<td>36 (1.8)</td>
<td>1961 (98.2)</td>
<td>1997</td>
</tr>
<tr>
<td>Season</td>
<td>Rainy</td>
<td>63 (1.4)</td>
<td>4515 (98.6)</td>
<td>4578</td>
</tr>
<tr>
<td></td>
<td>Dry</td>
<td>190 (2.0)</td>
<td>9208 (98.0)</td>
<td>9398</td>
</tr>
<tr>
<td>Zone</td>
<td>Central</td>
<td>76 (1.5)</td>
<td>4976 (98.5)</td>
<td>5052</td>
</tr>
<tr>
<td></td>
<td>Northeast</td>
<td>8 (0.6)</td>
<td>1301 (99.4)</td>
<td>1309</td>
</tr>
<tr>
<td></td>
<td>South</td>
<td>11 (0.6)</td>
<td>1685 (99.4)</td>
<td>1696</td>
</tr>
<tr>
<td></td>
<td>Northern</td>
<td>103 (4.8)</td>
<td>2029 (95.2)</td>
<td>2132</td>
</tr>
<tr>
<td></td>
<td>Bangkok</td>
<td>55 (1.4)</td>
<td>3732 (98.6)</td>
<td>3787</td>
</tr>
<tr>
<td>Gender</td>
<td>Male</td>
<td>138 (1.9)</td>
<td>7138 (98.1)</td>
<td>7276</td>
</tr>
<tr>
<td></td>
<td>Female</td>
<td>115 (1.7)</td>
<td>6585 (98.3)</td>
<td>6700</td>
</tr>
<tr>
<td>Specimen</td>
<td>Blood</td>
<td>10 (0.4)</td>
<td>2747 (99.6)</td>
<td>2757</td>
</tr>
<tr>
<td></td>
<td>Stool</td>
<td>243 (2.2)</td>
<td>10,976 (97.8)</td>
<td>11,219</td>
</tr>
</tbody>
</table>
Analysis of risk factors

The multivariable analysis (Table 2) revealed seasons and regions of Thailand as significant risk factors associated with the presence of *Salmonella* Kedougou when compared with other NTS in the final logistic analysis. The seasonal trends have a significant effect (p = 0.023) in the infection rates caused by *Salmonella* Kedougou compared with other NTS. The dry season was the period with the highest odds ratio (OR) for infection with *Salmonella* Kedougou (1.4, 95% confidence interval [CI] [1.03–1.8]). Also, the northern region of Thailand presented the highest OR of *Salmonella* Kedougou compared to other NTS. In contrast, the northeast was the region with the lowest OR (0.42, 95% CI [0.2–0.9]). In addition, the presence of *Salmonella* Kedougou in stool samples was associated with a statistically higher OR than blood samples (p < 0.01). The logistic regression analysis showed that neither the age group (p = 0.27) nor the gender of the patient (p = 0.22) posed a significant risk to Thai patients for being infected with *Salmonella* Kedougou compared to other NTS.

### Table 2. Multivariable Analysis of the Significant Single Factors Associated to the Presence of *Salmonella* Kedougou Compared to Other Nontyphoidal *Salmonella*

<table>
<thead>
<tr>
<th>Variables</th>
<th>Levels</th>
<th>OR</th>
<th>95% CI</th>
<th>p-Value (LR)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zone</td>
<td>Bangkok</td>
<td>1.13</td>
<td>0.8–1.6</td>
<td>0.5</td>
</tr>
<tr>
<td></td>
<td>Central</td>
<td>4.02</td>
<td>2.9–5.6</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>Northern</td>
<td>0.44</td>
<td>0.2–0.8</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td>South</td>
<td>0.42</td>
<td>0.2–0.9</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td>Northeast</td>
<td>1.4</td>
<td>1.0–1.8</td>
<td>0.023</td>
</tr>
<tr>
<td>Season</td>
<td>Rainy</td>
<td>0.13</td>
<td>0.07–0.25</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>Dry</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Specimen</td>
<td>Stool</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Blood</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
</tbody>
</table>

OR, odds ratio; CI, confidence interval; LR, likelihood ratio.

### Antimicrobial susceptibility testing

Multidrug resistance (resistance to at least three different classes of antimicrobials) was observed in 57/66 (86%) of the *Salmonella* Kedougou isolates from human origin isolated in Thailand, whereas 7 (11%) were pansusceptible. The two remaining isolates exhibited resistance to neomycin and ciprofloxacin together with nalidixic acid, respectively. Analysis of the resistance profiles indicated that 35 isolates (53%) were resistant to at least eight antimicrobials comprising a diverse range of phenotypes (n = 21). The most common resistance profile among the Thai isolates was amoxicillin + clavulanic acid, ampicillin, apramycin, chloramphenicol, gentamicin, spectinomycin, sulfamethoxazole, and tetracycline resistance. Two strains exhibited high level of resistance to ciprofloxacin (MIC ≥2 mg/L) and nalidixic acid (MIC ≥16 mg/L). Further, 3 out of the 66 strains (4.5%) were phenotypically resistant to cefotaxime and ceftiofur, indicating the presence of ESBL. Analysis of sequenced amplicons identified the presence of *bla*<sub>CTX-M-63</sub> in two of the isolates, whereas the remaining strain yielded a *bla*<sub>CMY-2</sub> gene. This strain exhibited resistance to cephalaxine, cefotaxime, and ceftoxitin. In addition, these three isolates also harbored the *bla*<sub>TEM-1b</sub> gene. No geographical clusters of resistance patterns were identified.

Two of the five human isolates from the United States showed a similar antimicrobial resistance pattern to the predominant resistant profile described in the Thai isolates: one was resistant to sulfamethoxazole and tetracycline, and the final two isolates were pansusceptible and were collected in 1995 and 2001.

In general, *Salmonella* Kedougou isolates of animal origin from the United Kingdom presented a lower level of resistances than the human isolates from Thailand and the United States (Table 3). Six out of 20 isolates (30%) were phenotypically resistant to four antimicrobials comprising three different phenotypes. Apart from a single isolate of duck origin exhibiting resistance to amoxicillin + clavulanic acid and ampicillin, strains generally appeared to be susceptible to these two compounds as well as apramycin, chloramphenicol, gentamicin, and spectinomycin. Isolates from turkeys showed the highest percentage of resistance, with four out of five strains resistant to spectinomycin, streptomycin, sulfamethoxazole, and tetracycline. One isolate recovered from a pig was resistant to ciprofloxacin and nalidixic acid. None of the animal isolates were resistant to third-generation cephalosporins.

### Pulsed-field gel electrophoresis

Electrophoresis of XbaI-digested genomic DNA from the 91 isolates revealed 45 different clusters (Fig. 2). XbaI profiles typically had 14 to 17 restriction fragments between 20 and 1135 kb. Indistinguishable PFGE patterns were present in isolates from Thai patients submitted in different regions of Thailand obtained at different points in time. For instance, isolates SH769, SH1133, SH3033, and SH544 were obtained from stool samples in Bangkok, south, central, and north of Thailand during April, May, December, and March 2008, respectively. All the five isolates collected from human samples in the United States shared identical XbaI patterns with isolates recovered from Thai patients. The U.S. isolates grouped in four different clusters. One of these clusters aggregated three
Table 3. Resistance Patterns Found in the Different Niches

<table>
<thead>
<tr>
<th>Country</th>
<th>N</th>
<th>AUG</th>
<th>AMP</th>
<th>ADR</th>
<th>XNL</th>
<th>CHL</th>
<th>CIP</th>
<th>COL</th>
<th>FFN</th>
<th>FOT</th>
<th>GEN</th>
<th>NAL</th>
<th>NEO</th>
<th>STR</th>
<th>SMX</th>
<th>TET</th>
<th>TMP</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>United States</strong></td>
<td>47</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><strong>United Kingdom</strong></td>
<td>0</td>
<td>0</td>
<td>0</td>
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<td>0</td>
<td>0</td>
<td>0</td>
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<td>0</td>
<td>0</td>
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<td>0</td>
</tr>
<tr>
<td><strong>Duck</strong></td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
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<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><strong>Environment</strong></td>
<td>3</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
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<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><strong>Turkey</strong></td>
<td>5</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
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AUG, amoxicillin + clavulanic acid; AMP, ampicillin; ADR, apramycin; XNL, ceftiofur; CHL, chloramphenicol; CIP, ciprofloxacin; COL, colistin; FFN, florfenicol; FOT, fosfomycin; GEN, gentamicin; NAL, nalidixic acid; NEO, neomycin; STR, spectinomycin; SMX, sulfonamides; TET, tetracycline; TMP, trimethoprim.

Discussion

To our knowledge, this is the first epidemiological study together with molecular fingerprinting and antimicrobial susceptibility testing data carried out in *Salmonella enterica* serovar Kedougou. The epidemiological data from Thailand are based on passive monitoring of samples submitted to the WHO International *Salmonella* and *Shigella* Centre and therefore should be analyzed with care. There are some limitations to the data: we have no knowledge of local laboratory skills reflecting the proficiency of *Salmonella* isolation techniques, nor do we have any information of the area in districts reporting data, or the occurrence of any outbreak that may have biased this dataset. The logistic regression analysis showed that there were significant differences in the risk factors associated with salmonellosis caused by *Salmonella* Kedougou compared with other NTS. The geographical regions and the season were identified as risk factors underlying the presence of *Salmonella* Kedougou in Thailand. Also, the presence of *Salmonella* Kedougou in stool samples was significantly higher than in blood when compared with other NTS. The authors have no biological evidence of any possible confounders that may interact in the final model.

The northern region of Thailand had a much higher prevalence of *Salmonella* Kedougou compared to the rest of the regions included Bangkok, which is normally the major urban nucleus with the highest population density and the highest incidences of NTS cases (Hendriksen et al., 2009). These geographical discrepancies in the infection rates caused by *Salmonella* Kedougou may also be a consequence of the cultural difference together with the diet regimes characteristic of each national region of Thailand. For instance, whereas the northern region has a tradition of consumption of animal meat, including raw pork (Navacharoen et al., 2009), the northeast is the poorest and least developed region in Thailand, with a diet that relies heavily on wild greens and forest foods (Chaiwat Pulsrikarn, pers. comm.). Similarly, low rates of *Salmonella* Kedougou were also observed in the southern region of Thailand. This region has a large Muslim
FIG. 2. Dendrogram analysis of the Salmonella Kedougou clones obtained from human isolates in Thailand and United States and animal isolates in United Kingdom. Blacked out antimicrobials are those for which an isolate is resistant. AUG, amoxicillin + clavulanic acid; AMP, ampicillin; APR, apramycin; XNL, ceftiofur; CHL, chloramphenicol; CIP, ciprofloxacin; COL, colestin; FFN, florfenicol; GEN, gentamicin; NAL, nalidixic acid; NEO, neomycin; SPE, spectinomycin; STR, streptomycin; SMX, sulfonamides; TET, tetracycline; TMP, trimethoprim.
population, and both pork consumption and infections with swine-associated serovars of *Salmonella* are disproportionately lower than in the other regions of Thailand (Hendriksen *et al*., 2009). In general, these cultural and dietary differences may also be the reasons for the low infection rates caused not just by *Salmonella* Kedougou but also by other NTS in the northeast and southern regions over the study period. However, there is some limitation to the data, as we do not know if there are regions of Thailand more efficient at reporting than others.

Contrary to other studies describing NTS with the infection peak over the rainy seasons (Morpeth *et al*., 2009), in Thailand the hot/dry season was the period with the highest rates of infection caused by *Salmonella* Kedougou. This observation suggests that this serovar is unlikely to be waterborne related, suggesting the animal production as the possible reservoir of infection. If open markets lack cooling systems for proper storage of retail meat, the hot temperatures reached over the hot/dry periods could promote bacterial proliferation and increase the risk of infection by consumption of contaminated meat. In the United Kingdom, although *Salmonella* Kedougou has been isolated from pig carcases, presence of this serovar is mainly associated with broiler production, in particular turkeys (www.defra.gov.uk/corporate/consult/turkey-turkey-ncp.pdf). However, the clonal types observed in United Kingdom differ from those isolated from the human cases in Thailand and the United States, suggesting an absence of any epidemiological link. We are not aware of any studies carried out in broilers in Thailand. Studies performed in retail meat in Thailand have reported *Salmonella* Kedougou as 1 of the 10 most common serovars isolated from pork (Vindigni *et al*., 2007). Shedding additional light on transmission would require performing PFGE on isolates from animals or retail meat samples in Thailand. The prevalence of *Salmonella* Kedougou isolates collected during different years. We do not have the over time and a small number of PFGE clusters aggregated isolates collected during different years. We do not have the history of the five *Salmonella* Kedougou isolates collected in the United States. However, all of them clustered together with isolates from Thailand sharing identical macrorestriction patterns. The U.S. *Salmonella* Annual Summary (2006) reported a total of 19 *Salmonella* Kedougou infections over a 10-year period (1996–2006). This is a very low incidence of *Salmonella* Kedougou cases. As described by the Enter-net Quarterly *Salmonella* Report April–June 2006/2 (www.hpa.org.uk/web/HPAweb File/HPAweb_C/1194947410123), at least 3 out of the 75 *Salmonella* Kedougou cases isolated in Europe were associated with travel to Thailand. Therefore, the results of this study strongly suggest that *Salmonella* Kedougou clones are not established in the United States, and are most likely associated to traveling. However, additional epidemiologic investigation is needed to clarify this issue. An increased number of human infections are acquired by international travel or global trade with food products. Thus, previous studies have associated the infections with certain *Salmonella* serovars such as Rissen (Hendriksen *et al*., 2008), Corvallis (Archambault *et al*., 2006), Schwarzengrund (Aarestrup *et al*., 2007), and Choleraesuis (Sirichote *et al*., 2010) with the combination of traveling to Thailand and the importation of different food products. Thus, there is a need for improving food safety to limit the sources of infections among the Thai population and travelers to the country (Aarestrup *et al*., 2007).

*Salmonella* Kedougou was isolated from several animal sources in the United Kingdom. When sub-typed by PFGE, a separate cluster was formed containing virtually all of the animal isolates. This suggests the absence of any link between isolates from animals in United Kingdom and the human cases observed in Thailand and United States. Despite the frequent occurrence of *Salmonella* Kedougou in animals in United Kingdom, this is not reflected by an increase in number of human cases. Thus, this might also suggest a difference in virulence among clones, with *Salmonella* Kedougou isolates from Thailand exhibiting more virulence factors than those present in the United Kingdom isolates.

Although one isolate of turkey origin presented identical PFGE profile to two of the human isolates, in general, animal and human isolates clustered separately, suggesting a different source of infection, probably pork. In addition, serovars associated to pork origin have been on the increased in Thailand during the last few years (Hendriksen *et al*., 2009) and *Salmonella* Kedougou was one of the 10 most common serovars isolated from pork in the country (Vindigni *et al*., 2007).

Multidrug-resistant *Salmonella* Kedougou may pose a risk to humans, especially if they are resistant to fluoroquinolones and third-generation cephalosporins. Previous studies conducted in Thailand have described the presence of multidrug-resistant *Salmonella* in a wide range of serovars (Archambault *et al*., 2006; Hendriksen *et al*., 2008; Khemtong and Chuan- chuen, 2008; Chuanchuen and Padungtod, 2009; Sirichote *et al*., 2010). Two of the *Salmonella* Kedougou strains from Thailand were resistant to fluoroquinolones, the drug of choice for treatment of severe *Salmonella* infections in humans (Archambault *et al*., 2006). In addition, three strains were identified as ESBL producers. Two of them harbored *blaCTX-M-63* and the remaining one, *blaCTX-M-63* is an uncommon variant belonging to the *blaCTX-M-8* subgroup previously described in *Escherichia coli* (Hopkins *et al*., 2006). Recent
epidemiological studies focused in ESBL-producing Enterobacteriaceae in Thailand (Hawkey, 2008; Kiratisin et al., 2008) have described bla_{CTX-M} genes, mainly bla_{CTX-M-14} and bla_{CTX-M-15}, as endemic in the country. However, to our knowledge this is the first report identifying Salmonella strains harboring bla_{CTX-M-66} and the first description of a bla_{CMY-2} in Salmonella Kedougou. Even though data on antimicrobial usage in disease prevention and as growth promoters are not accessible in Thailand, enrofloxacin and ceftiofur, a third-generation cephalosporin, are extensively used in swine production as growth promoters (Kulwicht et al., 2007). The use of these antimicrobials on farm may have contributed to the selection of certain resistant clones. Although data from this study do not suggest that Salmonella Kedougou is an invasive serovar, if these strains acquire resistance to third-generation cephalosporins, the treatment of infections with β-lactam antibiotics could be seriously compromised.

Further work should be initiated to identify the possible reservoirs and routes of transmission of Salmonella Kedougou, especially in those regions where the burden of the disease is highest. The establishment of surveillance programs for source attribution analysis would clarify the contribution of the different reservoirs to human infection. For the benefit of Thai people and international travelers, we encourage the Thai authorities to implement control measures at critical points in the food chain to improve food safety and to avoid the dissemination of potential pathogens and their resistance traits via the food chain. In addition, intervention strategies should be introduced in the animal production site to minimize the use of antimicrobial growth promoters that could compromise human treatment of infectious diseases.

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