Uropathogenic *Escherichia coli* ST131 in urinary tract infections in children

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**Purpose:** *Escherichia coli* sequence type (ST) 131, a multidrug-resistant clone causing extraintestinal infections, has rapidly become prevalent worldwide. However, the epidemiological and clinical features of pediatric infections are poorly understood. We aimed to explore the characteristics of ST131 *Escherichia coli* isolated from Korean children with urinary tract infections.

**Methods:** We examined 114 uropathogenic *E. coli* (UPEC) isolates from children hospitalized at Chung-Ang University Hospital between 2011 and 2014. Bacterial strains were classified into STs by partial sequencing of seven housekeeping genes (*adk, fumC, gyrB, icd, mdh, purA,* and *recA*). Clinical characteristics and antimicrobial susceptibility were compared between ST131 and non-ST131 UPEC isolates.

**Results:** Sixteen UPEC isolates (14.0%) were extended-spectrum β-lactamase (ESBL)-producers; 50.0% of ESBL-producers were ST131 isolates. Of all the isolates tested, 13.2% (15 of 114) were classified as ST131. There were no statistically significant associations between ST131 and age, sex, or clinical characteristics, including fever, white blood cell counts in urine and serum, C-reactive protein, radiologic abnormalities, and clinical outcome. However, ST131 isolates showed significantly lower rates of susceptibility to cefazolin (26.7%), cefotaxime (40.0%), cefepime (40.0%), and ciprofloxacin (53.3%) than non-ST131 isolates (65.7%, 91.9%, 92.9%, and 87.9%, respectively; *P* <0.001 for all).

ESBL was more frequently produced in ST131 (53.3%) than in non-ST131 (8.1%) isolates (*P* <0.01).

**Conclusion:** ST131 *E. coli* isolates were prevalent uropathogens in children at a single medical center in Korea between 2011 and 2014. Although ST131 isolates showed higher rates of antimicrobial resistance, clinical presentation and outcomes of patients were similar to those of patients infected with non-ST131 isolates.

**Key words:** Child, Escherichia coli, Multilocus sequence typing, Urinary tract infection

**Introduction**

Urinary tract infection (UTI) is one of the most common bacterial infections and *Escherichia coli* is the most common bacterial species causing UTI in children. Infants and children diagnosed with UTI usually present with fever only, and absence of urinary symptoms such as dysuria and frequency. Therefore, UTI is an important disease that should be considered in pediatric febrile illness without focus. Although febrile UTIs in children might be upper UTIs, antimicrobial therapy has mostly been performed successfully in developed countries. However, extended spectrum beta-lactamase (ESBL)-producing uropathogenic *E. coli* (UPEC) has emerged as a major global public health concern because infections due to ESBL-containing pathogens continue to be associated with significant morbidity and mortality worldwide.
ESBL can spread between bacteria in hospitals and communities via transmissible genetic elements or specific bacterial clones. Recently, the sequence type (ST) 131 E. coli strain emerged as a major clone harboring drug-resistance potential such as that provided by ESBL in various types of extraintestinal infections worldwide. Therefore, it is very important to continuously monitor and promptly identify the regional epidemiology of this virulent strain of E. coli.

In the past decade, many studies have been performed to identify the clinical, epidemiological, and microbiological characteristics of ST131 in adults. However, limited data are available regarding extraintestinal infections caused by ST131 strains in children. In this study, we aimed to determine the characteristics of ST131 UPEC strains isolated from Korean children with UTI.

1. Subjects
This was a prospective study conducted from March 2011 to December 2014. All subjects were children who were younger than 18 years, had been admitted to Chung-Ang University Hospital with diagnosis of community-acquired UTI. Informed consent was submitted by the parents/guardians of all subjects, and by the children older than 8 years when they were enrolled.

Urine samples were collected from midstream urine in toilet-trained children and by catheterization in others. UTI was defined as the presence of both positive urine culture (>10^5 colony-forming units [CFUs]/mL) and pyuria (>5 white blood cells [WBCs]/high-power field) with fever and/or urinary symptoms (i.e., urgency, frequency, and dysuria). UTI with fever or radiologic abnormality of kidney was defined as acute pyelonephritis (APN), and otherwise as cystitis. Asymptomatic bacteriuria, defined as significant growth of E. coli (>10^5 CFUs/mL) without fever or any urinary symptoms, were excluded in this study.

Clinical data were obtained from the medical records of the patients and entered into standardized case report forms. Inappropriate antimicrobial therapy was defined as the use of antimicrobials nonsusceptible to the isolated E. coli. Clinical success was defined as the resolution of fever with an improvement in urinalysis at day 7. Otherwise, patients were regarded as treatment failure. Recur was defined as 2nd episode of UTI after the completion of antimicrobial therapy for 1st episode of UTI. The present study protocol was reviewed and approved by the Institutional Review Board of Chung-Ang University College of Medicine (approval number: 10-013-02-03).

2. Isolates
All UPEC isolates were identified on the basis of typical morphology, lactose fermentation, positive spot indole test, and VITEK 2-GN cards (bioMérieux, Marcy l’Etoile, France). The isolates were stored at -80°C until further use. Extraction and purification of bacterial whole-cell DNA from UPEC colonies were performed as described in the instruction manual of the QIAamp Kit (QIAGEN GmbH, Hilden, Germany).

3. Phylogenetic grouping
Multiple polymerase chain reaction (PCR) assays for gadA, chuA, and yjaA genes and the TSPE4.C2 DNA fragment were performed. The PCR mixture composition and thermal cycling conditions have been previously described. PCR products were electrophoresed on agarose gels and photographed under ultraviolet transillumination. We could identify the 373-, 281-, 216-, and 158-base pair (bp) bands for gadA, chuA, yjaA, and TSPE4.C2, respectively. On the basis of the results of the PCR, chuA− TSPE4.C2−, chuA− TSPE4.C2+, chuA+ yjaA−, and chuA+ yjaA+ strains were classified as belonging to groups A, B1, B2, and D, respectively. The glutamate decarboxylase-alpha (gadA) gene of E. coli was used as an internal amplification control.

4. Multilocus sequence typing
The MLST scheme used the following seven housekeeping genes: adk, fumC, gyrB, icd, mdh, purA, and recA. PCR amplification for each gene was performed using the same reaction mixture, primers, and conditions as described previously. Because this study focused on ST131, we first screened the UPEC isolates on the basis of adk and fumC. If allele typing of adk and fumC identified allele 53 (adk53) and allele 40 (fumC40), respectively, further allele typing for gyrB, icd, mdh, purA, and recA was performed to confirm ST131. If the isolate had alleles other than adk53 or fumC40, the MLST procedure was stopped and the isolate was assigned to non-ST131. However, in the case of UPEC isolates with ESBL, we continued to analyze the alleles of other MLST genes to identify the full range of allele types of MLST for the isolates.

The sequencing reactions were carried out using a BigDye Terminator v3.1 Cycle Sequencing Kit (Applied Biosystems Inc., Foster City, CA, USA) with each forward primer. The reaction products were separated and detected using an Applied Biosystems 3730xl automated sequencer (Applied Biosystems Inc.), and the data were analyzed using the Sequencing analysis v3.3 software (Applied Biosystems Inc.). The allelic profile was summarized by assigning a ST via a web database (http://mlst.warwick.ac.uk/mlst/dbs/Ecoli/). The eBURST v3 software (available at http://www.mlst.net) was used to estimate the relationships between the isolates and to assign the strains to a clonal complex (CC).

5. Antimicrobial susceptibility test
Antimicrobial susceptibility testing of all isolates to ampicillin, amoxicillin-clavulanate, piperacillin-tazobactam, cefazolin, cefotaxime, cefepime, trimethoprim-sulfamethoxazole (TMP-SMX),
amikacin, gentamicin, ciprofloxacin, ertapenem, meropenem, imipenem, and tigecycline was performed using a VITEK 2 automated system (bioMérieux, Marcy l’Etoile, France). In addition, each isolate was tested using a VITEK 2 system with the ESBL test panel (bioMérieux) included in the N045 card (bioMérieux). *In vitro* antimicrobial susceptibility testing was performed by the broth microdilution method and the results were interpreted using the 2010 Clinical Laboratory Standard Institute breakpoints. Isolates resistant to antibiotics of three or more different classes were classified as multidrug resistant (MDR).

6. Statistical analyses
All statistical analyses were performed using IBM SPSS Statistics ver. 22.0 (IBM Co., Armonk, NY, USA). The prevalence of virulence genes and antibiotic resistance patterns were compared between the 2 groups using Pearson chi-square test and Fisher exact test. Continuous variables were compared with the Student *t* test and Mann-Whitney *U* test. A *P* value of <0.05 was considered statistically significant.

Results

1. Characteristics of subjects
A total of 114 subjects were enrolled in the study; the median age was 5.7 months (range, 0.5–227.6 months), and the male to female ratio was 1.53:1. Ninety-five of subjects (83.3%) presented with fever and 21 (18.4%) presented with urologic symptoms. APN and cystitis were diagnosed in 101 (88.6%) and 13 (11.4%), respectively. The mean serum WBC count was 14,942.5±5,606.2/mm³ and mean C-reactive protein (CRP) level was 48.2±49.3 mg/L. Kidney ultrasonography (US), dieremcapsuocinic acid (DMSA) scan, and voiding cystourethrogram (VCUG) were performed in 99 (86.8%), 92 (80.7%), and 14 subjects (12.3%), respectively. Radiological abnormalities were found in 40 subjects (40.4%) in US, 23 (25.0%) in DMSA scan, 5 (35.7%) in VCUG. The number of subjects with abnormality in any of the three radiologic modalities was 53 (52.4%).

Cystitis were all treated only with oral 3rd generation cephalosporins such as cefpodoxime (n=4), cefdinir (n=1), and cefixime (n=8). In all APNs, cefotaxime was injected empirically, and continued to the end of admission in most cases (n=90, 89.1%). Among the 16 UTIs caused by ESBL-producing *E.coli*, 8 cases were treated definitively with another antibiotic agent, which was carbapenem (n=6, 75.0%) or piperacillin-tazobactam (n=2, 25.0%). Treatment failure or recurrence did not occur in the study subjects.

2. Phylogenetic grouping and MLST
One UPEC isolate from each subject was analyzed for phylogenetic grouping and MLST. In the phylogenetic grouping, *gadA* was amplified in all the UPEC isolates. Group B2 was the most common one at 83% (n=95), followed by groups D (11%, n=13), A (3%, n=3), and B1 (2%, n=3) (Fig. 1). All ST131 isolates and most isolates with ESBL (87.5%) were in the B2 group.

In the MLST, a total of 15 UPEC isolates were assigned to both adk53 and fimC40. All these isolates were confirmed as ST131 with an allelic profile of 53-40-47-13-36-28-29. In the other 99 isolates, none were assigned to adk53 or fimC40. Additionally, isolates with ESBL (n=16) were all fully analyzed for MLST, with the following results: ST131 (n=8, 50.0%), CC14 (ST14 and ST1193; n=3, 18.8%), ST69 (n=2, 12.9%), ST38 (n=1, 6.3%), ST73 (n=1, 6.3%), ST95 (n=1, 6.3%).

3. Clinical characteristics of ST131
UTIs caused by ST131 isolates (n=15) were compared with those caused by non-ST131 isolates (n=99) in terms of the clinical aspects (Table 1). The mean ages of subjects were 13.61±17.15 and 22.62±40.09 months (*P*=0.409), and sex ratios were 1.8 and 1.5 in ST131 and non-ST131 groups (*P*=0.759), respectively. Fever and urologic symptoms manifested in 80.0% and 20.0%, respectively, of ST131 group, which did not differ significantly from the findings for the non-ST131 group (*P*=1.000 and *P*=0.720, respectively). The proportion of APN was also similar in ST131 (n=13, 86.7%) and non-ST131 (n=86, 86.9%) groups (*P*=1.000). In addition, laboratory findings such as urine nitrite, pyuria, leukocytosis, and CRP elevation were not significantly different between the 2 groups (*P*>0.05, for all). Finally, in ST131 group, initial empirical antimicrobials were more inappropriate than in non-ST131 group (53.3% vs. 8.1%, respectively; *P*<0.001), but definitive antimicrobials were not significantly different between the 2 groups in terms of inappropriateness (20.0% vs. 7.1%, respectively; *P*=0.223).
conventional phylogenetic group B2 was the most common group (83.3%) and the proportion of ST131 isolates was 13.2%. The proportion of isolates producing ESBL was 14.0%, and half of them were assigned to ST131. Among the ST131 isolates, 53.3% were ESBL producers and 46.7% were resistant to quinolone. Besides the inappropriate empirical antimicrobial therapy, no clinical and prognostic factor differed between ST131 and non-ST131 UPEC isolates. After a PCR strategy to assign *E. coli* isolates rapidly to 1 of the 4

![Fig. 2. Antimicrobial susceptibility, ST131 vs. non-ST131. Results of susceptibility testing with carbapenems and tigecycline were not included because all uropathogenic *Escherichia coli* isolates in the study showed 100% susceptibility, regardless of ST. ST, sequence type; AMX-CLV, amoxicillin-clavulanate; PIP-TAZ, piperacillin-tazobactam; TMP-SMX, trimethoprim-sulfamethoxazole; ESBL, extended spectrum beta-lactamase. *P<0.001.](https://example.com/fig2.png)
phylogroups A, B1, B2, or D was developed in 2000, it has been used extensively. Although the distribution of phylogenetic groups varies among countries, *E. coli* isolates causing extraintestinal infections were generally found to belong to groups B2 and D. Recently, MLST was developed as an attractive sequence-based genotyping technique, which provides reproducibility, comparability, and transferability between laboratories. MLST is thus a powerful tool for both global and long-term surveillance.

The advent of next-generation sequencing and precision typing methods has expanded the understanding of the molecular epidemiology of ESBL-producing bacteria and revealed widespread dissemination of *E. coli* ST131 bearing CTX-M-15. This clone might be the “driving force” behind the current ESBL pandemic, and circulating CTX-M strains often harbor multiple co-resistance genes such as those conferring resistance to TMP-SMX, aminoglycosides, and fluoroquinolones, thereby limiting therapeutic options.

*E. coli* ST131 has been shown to carry a broad range of virulence and resistance genes on transferable plasmids, and has been globally disseminated. Recent reports indicate that this drug-resistant *E. coli* clonal group is responsible for community-acquired UTI and bacteremia, as well as a significant number of healthcare-associated infections. Among the ESBL-producing *E. coli* isolates from various sources, 39%–57% belonged to the ST131 lineage in recent studies. In Korea, ST131 was found to be the predominant ST; it represented 31%–36% of ESBL-producing *E. coli* strains, and was distributed in both community and hospital isolates. ST131 isolates were significantly associated with ESBL, especially CTX-M-15, and the majority of them were MDR. In a study of outpatients with acute cystitis wherein *E. coli* had been isolated, treatment failure was more common among patients infected with ST131 *E. coli*. However, patients with bloodstream infection by ST131 *E. coli* isolates did not exhibit poorer treatment outcomes than those with non-ST131 isolates in the other study.

Although numerous studies of *E. coli* in adults have been performed, the clinical and molecular biological data for children remain very limited. As UTIs are the most common bacterial infection in children, it is especially important to identify the clinical significance and epidemiology of ST131 *E. coli* in pediatric UTI. UPEC strains recently isolated from Korean children belonged to phylogenetic groups B2 (61.6%–68.1%) and D (12.8%–26.8%), comprising the majority of all isolated strains. In a Taiwanese study of infants hospitalized for community-acquired UTIs caused by ESBL-producing *E. coli*, ST131 accounted for 65% of the 111 isolates. In the current study, ST131 accounted for 13.2% of total UPEC and 50.0% of ESBL-producing UPEC strains.

In addition, *E. coli* ST131 isolates also had high rates of non-susceptibility to quinolones. In contrast to ESBL, which is transmissible between UPEC strains by plasmids, quinolone resistance mainly develops in the bacterial chromosome. Therefore, a high resistance rate to quinolone in ST131 *E. coli* isolates supports the notion that this clone has highly drug-resistant characteristics. In this study, ST131 UPEC strains showed higher resistance to ciprofloxacin, that is, 46.7%, than non-ST131 strains (12.1%).

It is generally accepted that invasive infections by ESBL producers in children are associated with longer hospital stays, recurrence, frequent complications, and increased mortality. However, a Taiwanese study reported that children with UTI caused by ST131 clones had similar risk factors, clinical features and outcomes as those with UTI caused by non-ST131. Additionally, in our study, there was no significant difference between ST131 and non-ST131 UPEC cases in relation to radiologic abnormality, treatment failure, or recurrence although empirical antimicrobial therapy was more inappropriate in UTIs caused by ST131 *E. coli*, which produced ESBL more often. This finding might be due to the easy clearance of uropathogens from the infected site (urinary tract) by natural drainage (urination), despite inappropriate antimicrobial coverage. Additionally, previous studies showed that use of non-carbapenem antimicrobial therapy against ESBL-producing uropathogens did not affect the time to defervescence after the initiation of antimicrobial treatment, treatment failure, and relapse rate in pediatric UTIs.

This study has several limitations. First, we screened ST131 using two (adk and fimC) of the 7 genes used in the MLST scheme rather than by sequencing all 7 genes. Although neither adk53 nor fimC40 was detected in non-ST131 isolates in this study, it is possible that a double locus variant (DLV) of ST131 was missed by this approach. This could result in underestimation of the influence of strains related to ST131. However, among a total of 18 DLVs of ST131 in the MLST database, no DLV had different alleles both in adk and fimC from ST131. Second, the number of ST131 isolates was very small; therefore, the comparison with non-ST131 isolates was not statistically powerful. Finally, the types of fimH or ESBL that have been investigated in adult studies of ST131 were not analyzed in this study. Larger scale studies of pediatric ST131 infection should be performed based on the findings of this study.

In summary, this is the first study for molecular epidemiology and clinical characteristic of ST131 *E. coli* in Korean children. ST131 *E. coli* was a prevalent uropathogen in Korean children admitted at a single hospital between 2011 and 2014. The ST131 UPEC isolates in this study were highly drug-resistant and a high proportion produced. Although no change of clinical outcome was produced by ST131 isolates, the epidemiology and clinical implications of ST131 *E. coli* should be monitored in invasive and noninvasive pediatric infectious diseases.

**Conflict of interest**

No potential conflict of interest relevant to this article was reported.
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