Ciprofloxacin-resistant *Escherichia coli* in Central Greece: mechanisms of resistance and molecular identification

Angeliki Mavroidi, Vivi Miriagou, Apostolos Liakopoulos, Evi Tzelepi, Angelos Stefos, George N Dalekos, and Efthymia Petinaki

**Abstract**

**Background:** Fluoroquinolone resistant *E. coli* isolates, that are also resistant to other classes of antibiotics, is a significant challenge to antibiotic treatment and infection control policies. In Central Greece a significant increase of ciprofloxacin-resistant *Escherichia coli* has occurred during 2011, indicating the need for further analysis.

**Methods:** A total of 106 ciprofloxacin-resistant out of 505 *E. coli* isolates consecutively collected during an eight months period in a tertiary Greek hospital of Central Greece were studied. Antimicrobial susceptibility patterns and mechanisms of resistance to quinolones were assessed, whereas selected isolates were further characterized by multilocus sequence typing and β-lactamase content.

**Results:** Sequence analysis of the quinolone-resistance determining region of the *gyrA* and *parC* genes has revealed that 63% of the ciprofloxacin-resistant *E. coli* harbored a distinct amino acid substitution pattern (GyrA:S83L + D87N; ParC:S80I + E84V), while 34% and 3% carried the patterns GyrA:S83L + D87N; ParC:S80I and GyrA:S83L + D87N; ParC:S80I + E84G respectively. The *aac (6 \text{'})-Ib-cr* plasmid-mediated quinolone resistance determinant was also detected; none of the isolates was found to carry the *qnrA*, *qnrB* and *qnrS*.

Genotyping of a subset of 35 selected ciprofloxacin-resistant *E. coli* by multilocus sequence typing has revealed the presence of nine sequence types; ST131 and ST410 were the most prevalent and were exclusively correlated with hospital and health care associated infections, while strains belonging to STs 393, 361 and 162 were associated with community acquired infections. The GyrA:S83L + D87N; ParC:S80I + E84V substitution pattern was found exclusively among ST131 ciprofloxacin-resistant *E. coli*. Extended-spectrum β-lactamase-positive ST131 ciprofloxacin-resistant isolates produced CTX-M-type enzymes; eight the CTX-M-15 and one the CTX-M-3 variant. CTX-M-1 like and KPC-2 enzymes were detected in five and four ST410 ciprofloxacin-resistant *E. coli* isolates, respectively.

**Conclusions:** Our findings suggest that, ST131 and ST410 predominate in the ciprofloxacin resistant *E. coli* population.

**Keywords:** *Escherichia coli*, Quinolones, MLST, Beta lactamases

**Background**

*Escherichia coli* is among the major pathogens in both community and hospital settings [1]. The prevalence of multidrug-resistant *E. coli*, (i.e., *E. coli* isolates resistant to more than three classes of antimicrobial agents) has been increased worldwide in the past decades. The emergence and worldwide dissemination of fluoroquinolone resistant *E. coli* isolates, that are also resistant to newer β-lactams due to the production of extended-spectrum β-lactamases (ESBLs) particularly CTX-M-type enzymes, is a significant challenge to antibiotic treatment and infection control policies [1]. The application of multilocus sequence typing (MLST) to isolates producing CTX-M-15 ESBL led to the recognition of an internationally disseminated clone, ST131, which is a virulent phylgroup B2, uropathogenic *E. coli* lineage. ST131 *E. coli* is associated with resistance to fluoroquinolones and aminoglycosides.

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Quinolones are widely used antimicrobials for the treatment of bacterial infections [2]. Their wide use has triggered increased bacterial resistance worldwide. Mutations in gyrA and parC genes are the most common mechanism involved in high-level quinolone resistance, yet the spread of plasmid-mediated quinolone resistance genes and efflux-pump mutants have also been described. In Greece, according to the recent data of WHONET the rate of ciprofloxacin resistant (CIP-R) E. coli varied from hospital to hospital and ranged from 5.6 to 49.5%. In the University Hospital of Larissa (UHL), that is the main tertiary hospital of Central Greece and serves a region of 1,000,000 habitants, an increase of ciprofloxacin-resistant E. coli from 16.3% in 2010 to 21% in 2011 was recorded. The aim of this study was to assess the epidemiological features of bacterial infections [2]. Their wide use has facilitated the spread of these resistant strains, which are being identified at an alarming rate in many parts of the world. In the case of Greece, the increase in the rate of CIP-R E. coli from 16.3% in 2010 to 21% in 2011 highlights the need for more effective surveillance and control measures to prevent further spread of these resistant strains. 

Methods

Bacterial isolates, susceptibility testing and clinical data

From May to December 2011, a total of 505 consecutive E. coli isolates were recovered from clinical samples taken as part of standard care of an equal number of individual patients admitted to the University Hospital of Larissa, Central Greece. Identification and antimicrobial susceptibility testing of bacterial isolates was performed by the Vitek-2 system (bioMérieux, Marcy l’Étoile, France). The interpretive criteria of the European Committee on Antimicrobial Susceptibility Testing were used (http://eucast.org/clinical_breakpoints).

Determination of MICs to ciprofloxacin norfloxacin, imipenem and meropenem was performed by the Etest method (bioMérieux). The double disk synergy test (DDST) was used to determine the ESBL production, as described previously [3]. The modified Hodge test was used for phenotypic detection of carbapenemase production [3]. All E. coli, that were classified as resistant to ciprofloxacin (MICs > =4 mg/L), were further analyzed for the underlying mechanisms of quinolone resistance, molecular typing and β-lactamase content.

The medical records of patients diagnosed with CIP-R E. coli infection were reviewed regarding their current and previous hospitalizations. Before obtaining the clinical information of the patients, approval was received by the Ethics Committee of the UHL, that is represented by the Infection Control Committee (number of permission 1234). Infections developed 48 h after hospital admission were characterized as UHL hospital-acquired infections. Infections due to CIP-R E. coli diagnosed within 48 h of hospital admission were characterized as community acquired infections. Finally, infections in patients who had been hospitalized in the preceding 6 months for more than 48 h in hospital facilities or nursing homes such as infections in patients transmitted to UHL from other hospitals were considered as health care associated infections.

PCR and sequencing of the Quinolone Resistance-Determining Region (QRDR) of the gyrA and parC genes

All Cip-R E. coli were screened for the presence of mutations in the QRDR of the gyrA and parC genes. After DNA extraction by using the Quick-gDNA™ MiniPrep kit (ZYM0 RESEARCH Corp., USA), the QRDRs of both gyrA and parC genes were amplified by polymerase chain reaction (PCR), as described previously [4] and the amplicons were sequenced on both DNA strands using an ABI3730 DNA sequencer (Applied Biosystems, Warrington, United Kingdom). For each isolate, the sequences of the gyrA and parC gene fragments were concatenated, maintaining the +1 reading frame, aligned and a neighbor-joining tree was constructed from the aligned sequences using the MEGA software [5].

Detection of plasmid-mediated quinolone resistance genes qnrA, qnrB, qnrS and aac(6′)-Ib-cr variant

Primers and conditions for PCR amplification of the qnrA, qnrB and qnrS genes, which encode three target-protecting proteins and the aac(6′)-Ib-cr variant, which encodes a bifunctional aminoglycoside-fluoroquinolone modified enzyme, were used as described previously [6], and the amplified PCR products obtained were sequenced on both DNA strands as described previously.

Detection of β-lactamases

Detection of the bla genes was performed by PCR using a panel of specific primers for blaTEM-1, blaCTX-M, blaSHV, blaOXA-1, blaQPC, blaCMY, blaVIM and blaKPO as described previously [7]. PCR products were purified by using the PureLink™ PCR Purification Kit (Invitrogen, USA) kit and sequenced. Nucleotide and deduced protein sequences were identified by comparing the sequences of the database of G. Jacoby and K. Bush (http://www.lahey.org/Studies).

Molecular typing of isolates

The major phylogenetic groups (A, B1, B2, D) were determined by PCR amplification of the three gene fragments of the scheme (chuA, yjaA and TSPE4.C2). Phylogroups were determined as described previously [8]. MLST was performed by PCR amplification and sequencing of seven housekeeping genes: adk (adenylate kinase), fumC (fumarate hydratase), gyrB (DNA gyrase), icd (isocitrate/isopropylmalate dehydrogenase), mdh (malate dehydrogenase), purA (adenyllosuccinate dehydrogenase), recA (ATP/GTP binding motif) by using primers and conditions as described at the...
whereas, only 25% to gentamicin. Nineteen percent of methoxazole, 70% to tetracycline, 60% to tobramycin, bapenem, 72% were resistant to trimethoprim- sulfa- while, 4.7% exhibited also resistance to at least one car-
an ESBL phenotype, exhibiting resistance to penicillins, to quinolones.
tance to four other classes of antimicrobial agents, (58 isolates) from health care associated infections.
recovered from community acquired infections, 15% (16 sions and sputum. Among them, 30% (32 isolates) were
11%), but also from bronchial secretions, cutaneous les-
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E. coli
One hundred six out 505 (21%) E. coli isolates were
Results
One hundred six out 505 (21%) E. coli isolates were found to be resistant to quinolones. These isolates were obtained from various clinical specimens; mainly from urine (76 out of 106; 72%) and blood (12 out of 106; 11%), but also from bronchial secretions, cutaneous lesions and sputum. Among them, 30% (32 isolates) were recovered from community acquired infections, 15% (16 isolates) from UHL hospital acquired infections and 55% (58 isolates) from health care associated infections.

Among 106 CIP-R E. coli isolates, 57% showed an ESBL phenotype, exhibiting resistance to penicillins, expanded- spectrum cephalosporins and aztreonam, while, 4.7% exhibited also resistance to at least one car-
papenem, 72% were resistant to trimethoprim- sulfa-
ketoxazole, 70% to tetracycline, 60% to tobramycin, whereas, only 25% to gentamicin. Nineteen percent of the CIP-R E. coli isolates exhibited concurrent resis-
tance to four other classes of antimicrobial agents, 58.5% to three classes, 13.2% to two classes and 3.8% to one class; the remaining 5.5% showed resistance only to quinolones.

Sequencing of the QRDR regions of the gyrA and parC genes of CIP-R isolates has revealed three different amino acid substitution patterns: GyrA:S83L + D87N; ParC:S80I + E84V (n = 36, 34%) and GyrA:S83L + D87N; ParC:S80I + E84G (n = 3, 3%). All but one of the 67 CIP-R E. coli isolates, that possessed the GyrA:S83L + D87N; ParC:S80I + E84V pattern, showed identical nucleotide sequences in the gyrA and parC, with the exception showing only a synonymous nucleotide substitution in the parC gene. In addition, the three isolates with the GyrA:S83L + D87N; ParC:S80I + E84G pattern had identical nucleotide sequences. On the contrary, the nucleotide sequences of gyrA/parC gene fragments of the 36 CIP-R isolates with the GyrA:S83L + D87N; ParC:S80I substitution pattern were found to be more polymorphic.

Out of the 106 CIP-R coli isolates, 35 were selected for further investigation including 21 with the pattern GyrA: S83L + D87N; ParC:S80I + E84V, 13 with the pattern GyrA:S83L + D87N; ParC:S80I, and one with the pattern GyrA:S83L + D87N; ParC:S80I + E84G. The selection was designed so as to include proportionally all the variations in the amino and nucleotide acid substitution patterns observed in the GyrA and ParC, the antimicrobial resistance patterns, the origin and the distribution of the isolates over the study period (Table 1).

The 35 representative CIP-R E. coli isolates were distributed into phylogroups B2 (30 isolates), D (two iso-
lates), A (two isolates) and B1 (one isolate) [Table 1]. Genotyping by MLST has revealed the presence of nine STs. ST131 (21 out 35 isolates), and ST410 (six out 35 isolates) were the most prevalent; ST44, ST90, ST162, ST361, ST1140, ST2509 included one isolate, while, ST393 included two isolates. According to clinical data, all ST131 isolates were associated with UHL hospital acquired infections and health care associated infections, ST410 CIP-R E. coli were exclusively associated with health care associated infections, while STs 393, 361 and 162 from community acquired infections. Strains that belonged to ST90, 44, 1140 and 2509 were isolated from patients with UHL hospital acquired infections.

Although, only three amino acid substitution patterns in the GyrA/ParC were identified among the CIP-R E. coli, we have sought to investigate any possible correla-
tions of the nucleotide polymorphisms and the STs of the isolates. For this purpose, a neighbor-joining tree was constructed from the concatenated sequences of the gyrA/parC gene fragments (Figure 1). The pattern GyrA:S83L + D87N; ParC:S80I + E84V was associated with ST131 strains possessing identical nucleotide sequences, while, the GyrA:S83L + D87N; ParC:S80I was found to various STs, including ST410. Except from one isolate (ID 362), the rest isolates with this substitution pattern differed in 12 polymorphic sites of the 643 bp nucleotide sequence of the gyrA/parC fragments. The third pattern, GyrA:S83L + D87N; ParC:S80I + E84G, was associated with ST393 (ID 296). The isolate ID 362, that belonged also to ST393, showed similarity in the gyrA/parC nucleotide sequences with the isolate ID 296 and they differed at only a single nucleotide site.

The presence of plasmid- mediated quinolone resist-
ance genes was also investigated in the 35 CIP-R E.coli isolates (Table 1, Figure 1). The aac (6’)-1b-cr variant was detected in 23 out of 35 CIP-R E.coli; of these 14 were ST131 and five ST410. The remaining four isolates belonged to ST44, ST90 and ST162 and ST393. None isolate was found to carry any of the qnrA, qnrB and qnrS gene.

The β-lactamase content of the 35 Cip-R E. coli was also determined (Table 1). ESBL- positive CIP-R isolates were found to produce enzymes of the CTX-M-1 family (n = 15); CTX-M-15 (n = 12) and CTX-M-3 (n = 3). Among them, nine belonged to ST131, five to ST410 and one to ST44 (Table 1). One out of the twelve ESBL-negative ST131 was a VIM-1 producer. Three ST410 CTX-M-producing strains co-produced the KPC-2 carbapenemase.
Table 1 Microbiological characteristics of the 35 representative CIP-R *E. coli*; phylogroups, MLST STs (clonal complexes; Cplx), presence/absence of the *aac (6')*-Ib-cr variant and beta-lactamase content

| Strain ID | Isolation date | Antimicrobial resistant pattern | ESBL | Phylogroup/MLST ST (CC) | *aac (6')*-Ib-cr | beta-lactamase content |
|-----------|----------------|---------------------------------|------|-------------------------|-----------------|------------------------|
| GlycylS83L + D87N/ ParC: S80I + E84V (n = 21) | | | | | | |
| 347 | 13/6/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (-) | CTX-M-15 |
| 161 | 27/5/2011 | AM, AMC, CAZ, CTX, ATMP, NN, SXT, CIP NOR positive | | B2/ ST131 | (+) | CTX-M-15 + OXA-1 |
| 391 | 30/5/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP NOR positive | | B2/ ST131 | (+) | CTX-M-15 + OXA-1 |
| 392 | 9/6/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | CTX-M-15 + OXA-1 |
| 393 | 18/7/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | CTX-M-15 + OXA-1 |
| 399 | 2/8/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | CTX-M-15 + OXA-1 |
| 160 | 3/5/2011 | AM, AMC, CAZ, CTX, ATMP, SXT, CIP, NOR positive | | B2/ ST131 | (+) | CTX-M-15 + TEM-1 |
| 505 | 4/9/2011 | AM, AMC, CAZ, CTX, AM, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | CTX-M-15 + TEM-1 + OXA-1 |
| 397 | 4/8/2011 | AM, AMC, CAZ, CTX, ATMP, T, SXT, CIP, NOR positive | | B2/ ST131 | (-) | CTX-M-3 |
| 301 | 1/6/2011 | AM, AMC, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | OXA-1 |
| 307 | 6/6/2011 | AM, AMC, GM, NN, TE, CIP, NOR negative | | B2/ ST131 | (+) | OXA-1 |
| 252 | 1/5/2011 | AM, SXT, CIP, NOR negative | | B2/ ST131 | (-) | TEM-1 |
| 206 | 16/5/2011 | AM, AMC, GM, TE, SXT, CIP, NOR positive | | B2/ ST131 | (+) | TEM-1 + OXA-1 |
| 320 | 25/5/2011 | AM, AMC, CIP, NOR negative | | B2/ ST131 | (+) | TEM-1 + OXA-1 |
| 346 | 3/6/2011 | AM, AMG, G, NN, SXT, TE, CIP, NOR negative | | B2/ ST131 | (+) | TEM-1 + OXA-1 |
| 803 | 4/12/2011 | AM, AMG, C, CTX, IMP, MEM, CIP, NOR negative | | B2/ ST131 | (+) | VIM-1-like + TEM-1 + OXA-1 |
| 252 | 10/6/2011 | AM, AMC, CAZ, CTX, ATMP, GM, NN, SXT, CIP, NOR positive | | B2/ ST131 | (+) | KPC-2 + CTXM-15 + TEM-1 + OXA-1 (ST23 Cplx) |
| 648 | 31/12/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, SXT, CIP, NOR positive | | B2/ ST410 | (+) | KPC-2 + CTXM-15 + TEM-1 + OXA-1 |
| 182 | 17/5/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, TE, CIP, NOR positive | | B2/ ST410 | (+) | CTXM-3 + TEM-1 + OXA-1 (ST23 Cplx) |
| 252 | 10/6/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, TE, CIP, NOR positive | | B2/ ST410 | (+) | CTXM-3 + TEM-1 + OXA-1 (ST23 Cplx) |
| 252 | 15/8/2011 | TE, SXT, CIP, NOR negative | | B2/ ST131 | (+) | none |
| 321 | 18/7/2011 | TE, CIP, NOR negative | | B2/ ST131 | (+) | none |
| 354 | 1/7/2011 | TE, CIP, NOR negative | | B2/ ST131 | (+) | none |
| 355 | 15/8/2011 | TE, SXT, CIP, NOR negative | | B2/ ST131 | (+) | none |
| GlycylS83L + D87N/ ParC: S80I + E84V (n = 1) | | | | | | |
| 296 | 6/8/2011 | AM, TE, CIP, NOR negative | | D/ ST393 (ST1 Cplx) | (+) | TEM-1 |
| GlycylS83L + D87N/ ParC: S80I (n = 13) | | | | | | |
| 148 | 9/6/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, SXT, CIP, NOR positive | | B2/ ST410 | (+) | KPC-2 + CTXM-15 + TEM-1 + OXA-1 (ST23 Cplx) |
| 182 | 17/5/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, AN, NN, TE, CIP, NOR positive | | B2/ ST410 | (+) | KPC-2 + CTXM-3 + TEM-1 + OXA-1 (ST23 Cplx) |
| 252 | 10/6/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, TE, CIP, NOR positive | | B2/ ST410 | (+) | KPC-2 + CTXM-3 + TEM-1 + OXA-1 (ST23 Cplx) |
| 648 | 31/12/2011 | AM, AMC, CAZ, CTX, ATMP, IMP, MEM, NN, TE, CIP, NOR negative | | B2/ ST410 | (+) | KPC-2 + TEM-1 (ST23 Cplx) |
| 132 | 24/5/2011 | AM, AMC, CAZ, CTX, AM, AN, NN, SXT, CIP, NOR positive | | B2/ ST410 | (+) | CTXM-15 + TEM-1 + OXA-1 (ST23 Cplx) |
| 281 | 15/8/2011 | AM, AMC, CAZ, CTX, AM, AN, GM, NN, SXT, CIP, NOR positive | | B2/ ST410 | (+) | CTXM-15 + OXA-1 (ST23 Cplx) |
| 384 | 30/11/2011 | AM, AMC, CAZ, CTX, AM, AN, SXT, CIP, NOR positive | | B2/ ST44 | (+) | CTXM-15 (ST10 Cplx) |
| 383 | 22/9/2011 | AM, TE, SXT, CIP, NOR negative | | A/ ST90 | (+) | TEM-1 (ST23 Cplx) |
| 259 | 14/7/2011 | AM, TE, SXT, CIP, NOR negative | | D/ ST162 | (+) | TEM-1 (ST469 Cplx) |
Discussion

During an eight months period of 2011, CIP-R E. coli accounted for 21% of the total E. coli isolates recovered from various clinical specimens from outpatients and inpatients of the UHL. The majority of the CIP-R E. coli isolates were multidrug-resistant, posing a challenge for therapeutic options. Only 5.5% were resistant to fluoroquinolones, but susceptible to various antimicrobial classes. The latter strains were recovered from both community and hospital acquired infections.

The majority of CIP-R E. coli belonged to ST131 and ST410, which were recovered from hospital and healthcare associated infections, whereas other studies have shown that such strains were disseminated in the community [11-16]. We have also shown previously that ST410 was linked with an outbreak of KPC-producing E. coli in a long-term care facility unit of Thessalia [17]; this clone, apart from carbapenemase producers, includes also isolates with CTX-M-15, as it was firstly reported in Spain [16]. All ST131 and ST410 CIP-R E. coli belonged to the virulent phylogroup B2, which is associated epidemiologically and experimentally with extraintestinal virulence [12,18]. The remaining isolates, that were isolated in a sporadic fashion in our study, were distributed equally to various STs, which have been previously reported among Cip-R E. coli isolates, and originated from community and hospital acquired infections [15].

Table 1 Microbiological characteristics of the 35 representative CIP-R E. coli; phylogroups, MLST STs (clonal complexes; Cplx), presence/absence of the aac (6')-1b-cr variant and beta-lactamase content (Continued)

| Isolate | Date | Antibiotics | Resistance Pattern | MLST ST | Notes |
|---------|------|-------------|--------------------|---------|-------|
| 328     | 10/9/2011 | AM,SXT,CIP,NOR | negative | B2/ST361 | TEM-1 |
| 362     | 21/10/2011 | AM,TE,SXT,CIP,NOR | negative | D/ST393 | TEM-1 |
|         |       |             |                   | ST31 Cplx |       |
| 129     | 25/5/2011 | AM,TE,SXT,CIP,NOR | negative | B2/ST1140 | TEM-1 |
| 327     | 9/6/2011  | CIP, NOR    | negative | A/ST2509 | none |

Figure 1 Neighbor-joining tree of the 35 representative CIP-R E. coli; for each isolate, the GyrA/ParC amino acid substitution pattern, the presence/absence of the aac (6')-1b-cr variant and the MLST ST is indicated.
In Greece, there are few studies on the mechanisms of resistance to fluoroquinolones [19-21], but to our knowledge this is the first report on the association of gyrA and parC mutations and the phylogenetic lineages of ciprofloxacin-resistant *E. coli* in Greece. According to our results, ST131 and ST410 carried specific patterns of GyrA/ParC amino substitutions. In more details, ST131 Cip-R *E. coli* in our hospital possessed the same amino acid substitution pattern GyrA:S83L + D87N; ParC:S80I + E84V, which has been previously identified among ST131 CIP-R *E. coli* isolated from humans and companion animals in the United States, United Kingdom, Australia and Korea [13-15]. On the other hand, the six ST 410 *E. coli* of our study carried the GyrA: S83L + D87N; ParC:S80I pattern, which has been identified in previous studies [22,23]. Nevertheless, the sequence types of the latter strains have not been determined.

The presence of plasmid-mediated resistance genes (*qnrA*, *qnrB* and *qnrS*) has also been reported in previous studies in Greece [19-21], but these genes were not detected in our collection of CIP-R *E. coli* isolates. On the other hand, the aac (6’)-Ib-cr variant was detected mainly in strains belonging to both ST131 and ST410, a finding that was also previously described [13,16].

Since isolates possessing the GyrA:S83L + D87N; ParC:S80I + E84V comprised a high percentage (67%) among CIP-R *E. coli*, and the representative isolates of this group have been assigned to ST131, we assume that the increase of fluoroquinolone-resistance during 2011 can be attributed to the dissemination of ST131 strains. The source and the time of importation of these isolates are unknown. As mentioned previously, since ST131 was first described in 2008 [24], it has disseminated worldwide [11-15,23-26]. Several factors such as host-to-host or foodborne transmission or environmental contamination have been suggested to contribute in their dissemination; reservoirs of ST131 have been identified in food and water sources, in nursing home residents, companion animals and food sources [27-29].

Central Greece is a rural area, where fluoroquinolones are widely used in veterinary and poultry, but no data exist about the incidence of resistant *E. coli* of animal source. It would be interesting to investigate the mechanisms of resistance and genetic relatedness of such strains in future studies. These studies would elucidate the relationships between STs of isolates recovered from human and animals and the roots of transmission.

Conclusions
In the present study we have characterized the mechanisms of resistance and explored the genetic relatedness of CIP-R *E. coli* recovered from community, hospital and health care associated infections in a tertiary care hospital in Central Greece. Our findings suggest that ST131 and ST410 predominate in the CIP-R *E. coli* population in our institution. The increase of resistance to fluoroquinolones observed during 2011 is attributed mainly to the wide dissemination of ST131 CIP-R *E. coli*.

Abbreviations
CIP: Ciprofloxacin; MLST: Multilocus sequence typing; ST: Sequence type; CC: Clonal complex; ESBL: Extended-spectrum β-lactamases; KPC: *Klebsiella pneumoniae* carbapenemase; QRDR: Quinolone Resistance-Determining Region; UHL: University Hospital of Larissa; DDST: Double disk synergy test; AM: Amoxicillin; AMC: Amoxicillin + clavulanic acid; CAZ: Ceftazidime; CTX: Cefotaxime; ATM: Aztreonam; GM: Gentamicin; NN: Tobramycin; AN: Amikacin; TE: Tetracycline; SXT: Trimethoprim-sulfamethoxazole; CIP: Ciprofloxacin; NOR: Norfloxacin; IMP: Imipenem; MEM: Meropenem.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
EP, AM, and VM conceived and designed the study. AM wrote the first draft of the paper and other co-authors contributed to the final draft. EP and GND were responsible for conducting the study and managing the data. AS conducted the interpretation of data. Others participated in data analysis and data interpretation. All authors read and approved the final manuscript.

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