Molecular Characterization of AmpC β-lactamases in Enterobacteriaceae

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Abstract

AmpC β-lactamases are enzymes that are resistant to β-lactams, such as penicillin and cephalosporin, but not cefoxitin and cefotetan. This study was conducted to characterize AmpC β-lactamases in Enterobacteriaceae. This study included 200 cephalosporin-resistant Gram-negative isolates recovered from different samples between January 2015 and December 2016. The isolates were subjected to phenotypic tests, and those that tested positive were further analyzed by PCR for six AmpC genotypes: ACC, DHA, FOX, CIT, MOX, and EBC. Among the 200 strains, 32% (64) were positive for AmpC β-lactamases by different phenotypic methods. The target genotypes were detected in 20 (10%) of the isolates. Pus was the predominant source of AmpC isolates. Klebsiella pneumoniae (55%) was the most common producer of AmpC β-lactamase. CIT-FOX was the predominant gene type. As there is variation in the prevalence of AmpC β-lactamases in different geographic regions, periodic surveillance and measures to control infection can prevent the spread of these genes.

Keywords: AmpC β-lactamases, AmpC Disk Test, Antibiotic Resistance

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INTRODUCTION

Resistance to antibiotics such as cephalosporin and carbapenem, is increasingly reported in Enterobacteriaceae and is associated with the activity of several types of enzymes, notably Extended spectrum β-lactamases (ESBL), AmpC β-lactamases, and carbapenemases.

AmpC beta-lactamases are enzymes that confer resistance to penicillin, monobactam and 1st, 2nd, and 3rd generation cephalosporin as well as cephamycins. These enzymes are not inhibited by beta-lactam inhibitors. Sometimes also confer resistance to carbapenems along with reduced membrane permeability. AmpC β-lactamases are of two types: chromosomal- and plasmid-mediated. AmpC β-lactamases are chromosomally encoded in a few Gram-negative bacteria, such as Citrobacter freundii, Enterobacter spp., Proteus spp., and Serratia marcescens. Plasmid-encoded AmpC β-lactamases were first discovered in 1988. These were found in Klebsiella pneumoniae (K. pneumoniae), Salmonella spp., and Proteus spp., through the mobility of AmpC genes in horizontal transfer between Enterobacterales.

There are no standardized guidelines for identifying AmpC β-lactamase-producing Enterobacteriaceae strains. The worldwide prevalence of plasmid-mediated AmpC ranges from 2% to 47%. In this study, we report the prevalence of AmpC β-lactamase-producing Enterobacteriaceae using phenotypic and molecular methods.

MATERIALS AND METHODS

We carried out a prospective, lab-based cross-sectional study that included 200 randomly collected non-replicate cephalosporin-resistant Enterobacteriaceae isolates from various sample materials, including pus (n = 29), wound swab (n = 28), urine (n = 107), blood (n = 11), stool (n = 2), endotracheal aspirate (ETA) secretion (n = 7), IV tip (n = 2), IV line (n = 1), and sputum (n = 13) from various departments of tertiary care (medicine, surgery, ICU, NICU, urology, nephrology, orthopedics, gynecology, pediatrics, chest and TB, special wards, and casualty) during the period from January 2015 to December 2016.

Identification of the organisms were performed using standard biochemical tests. Antibiotic susceptibility testing was performed using the Kirby–Bauer disc diffusion method. The isolates were then subjected to phenotypic tests, namely the AmpC disc antagonism test, AmpC disc test, modified Hodge test (MHT) and phenylboronic acid test. Subsequently, phenotypic test-positive isolates were subjected to molecular characterization via multiplex PCR for the common AmpC genotypes: DHA, MOX, CIT, ACC, FOX, and EBC.

Antibiotic Susceptibility Testing

A single colony taken from nutrient agar was inoculated into peptone water and placed in an incubator at 37°C for 2 h, and the opacity of the broth was checked with 0.5 MacFarland. Lawn cultures of the test isolates were inoculated on Muller Hinton agar (MHA) plates. Antibiotics were placed on the plates and incubated for 18-24 h at 37°C. The results were interpreted as per CLSI guidelines 2015.

Cefoxitin Screening Test

The test isolate was taken from nutrient agar, inoculated in peptone water, and incubated for 2 h at 37°C, after which the broth turbidity was checked with 0.5 MacFarland standard. The test isolates were seeded on MHA plates. Cefoxitin discs (Himedia) were placed on the surface of the MHA plates and incubated for 18-24 h. The zone of inhibition was interpreted as per CLSI guidelines 2015.

AmpC Disc Antagonism Test (DAT)

The test isolate was inoculated in peptone water for 2 h. The opacity of the broth was checked with 0.5 MacFarland standard spread over MHA. Cefotaxime (30μg) and ceftazidime (30μg) discs were placed 20 mm apart from cefoxitin disc on MHA plates and incubated for 18-24 h. Blunting of cefotaxime or ceftazidime inhibition zone near cefoxitin was considered a positive isolate for the AmpC test.

Modified (Cefoxitin) Hodge Test (MHT)

E. coli ATCC 25922 was seeded on MHA. Cefoxitin (30μg) discs were placed on the MHA. The
isolates was inoculated from the edge of cefoxitin disc until the periphery and incubated at 37°C overnight. Isolates with cloverleaf patterns were positive for AmpC production.

**AmpC Disc Test**

*E. coli* ATCC 25922 lawn culture was inoculated on an MHA plate. Cefoxitin disc was placed on MHA. Saline solution (20µL) was added to a sterile plain disc and placed near the cefoxitin disc. Three to four colonies of the strain were placed on the plain disc. The plate was incubated at 35°C for 18 h. The plates were read as follows: indentation (heart-shaped)/flattening of the zone around cefoxitin disc was positive for ampC production and no distortion around cefoxitin disc indicated a negative result.

**Phenylboronic Acid Test**

Phenylboronic acid (Himedia) (120 g) was dissolved in three mL of dimethylsulfoxide. Sterile water (3 mL) was added to this solution. Twenty microliters of the prepared phenylboronic acid were pipetted onto cefoxitin discs. The discs were then dried for 30 min. The test strains were seeded on MHA, air-dried and cefoxitin (30 μg) and discs containing cefoxitin/phenylboronic acid (30/400 μg) were placed on the plate. The seeded plates were then incubated for 18 h at 35°C. An organism that produces an enhanced zone of inhibition of 5 mm or more around the boronic acid/cefoxitin disc compared to the zone of inhibition around the cefoxitin disc was considered an AmpC-producing organism.

**Multiplex PCR for AmpC β-lactamases**

**Preparation of Template DNA**

All 64 isolates that were positive for the ampC disc test and phenylboronic acid were subjected to multiplex PCR for amplification.11

The isolates were grown on nutrient agar and the colonies were suspended in

<table>
<thead>
<tr>
<th>No.</th>
<th>Organism</th>
<th>No. of isolates</th>
<th>AmpC producers positive in AmpC disk &amp; MHT test</th>
<th>AmpC producers by phenyl boronic acid test</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td><em>Escherichia coli</em></td>
<td>87</td>
<td>24 (27.5%)</td>
<td>25 (28.7%)</td>
</tr>
<tr>
<td>2.</td>
<td><em>Klebsiella pneumoniae</em></td>
<td>84</td>
<td>31 (36.9%)</td>
<td>32 (38.0%)</td>
</tr>
<tr>
<td>3.</td>
<td><em>Klebsiella oxytoca</em></td>
<td>13</td>
<td>1 (7.6%)</td>
<td>1 (7.6%)</td>
</tr>
<tr>
<td>4.</td>
<td><em>Proteus mirabilis</em></td>
<td>7</td>
<td>2 (28.5%)</td>
<td>2 (28.5%)</td>
</tr>
<tr>
<td>5.</td>
<td><em>Proteus vulgaris</em></td>
<td>3</td>
<td>2 (66.7%)</td>
<td>2 (66.7%)</td>
</tr>
<tr>
<td>6.</td>
<td><em>Citrobacter</em></td>
<td>1</td>
<td>1 (100%)</td>
<td>1 (100%)</td>
</tr>
<tr>
<td>7.</td>
<td><em>Enterobacter</em></td>
<td>5</td>
<td>1 (20%)</td>
<td>1 (20%)</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td>200</td>
<td>62 (31%)</td>
<td>64 (32%)</td>
</tr>
</tbody>
</table>

**Figure 1.** Modified Hodge test (Cefoxitin Test)  
**Figure 2.** AmpC disc Test
Statistical Package for the Social Sciences (SPSS) Version 20, using simple descriptive statistics, percentiles and qualitative analysis.

RESULTS

Of the 200 isolates, 149 were found to be resistant to cefoxitin (i.e., results were positive for cefoxitin screening). Of these 149 isolates, 64 (42.9%) were *Escherichia coli* (*E. coli*), 65 (43.5%) *K. pneumoniae*, 9 (6.0%) *Klebsiella oxytoca*, 4 (2.6%) *Proteus mirabilis* (*P. mirabilis*), 2 (1.3%) *Proteus vulgaris* (*P. vulgaris*), 1 (0.7%) *Citrobacter* spp., and 4 (2%) *Enterobacter* spp. The resistogram of the isolates indicated: 94.5% resistant to ceftriaxone, 69% to cefepime, 28% to cefepime-tazobactam, 29.5% to piperacillin-tazobactam, 32.5% to imipenem, 31.5% to meropenem, 41.5% to norfloxacin, 79% to ciprofloxacin, 41.5% to nitrofurantoin, 4% to fosfomycin, 65.5% to gentamycin, 48% to amikacin, 37% to chloramphenicol and 70% to cotrimoxazole.

Among the 200 isolates, 62 (31%) were found to be positive for AmpC β-lactamases by the AmpC disc test (Figure 1) and MHT test (Figure 2), and 64 (32%) isolates were positive by the phenylboronic acid method. The resistogram of the isolates indicated: 94.5% resistant to ceftriaxone, 69% to cefepime, 28% to cefepime-tazobactam, 29.5% to piperacillin-tazobactam, 32.5% to imipenem, 31.5% to meropenem, 41.5% to norfloxacin, 79% to ciprofloxacin, 41.5% to nitrofurantoin, 4% to fosfomycin, 65.5% to gentamycin, 48% to amikacin, 37% to chloramphenicol and 70% to cotrimoxazole.

The positive isolates from the phenotypic
AmpC test were subjected to multiplex PCR using primers for six families. In Table 2 we present the AmpC producers identified by multiplex PCR. Twenty isolates were assigned the genotypes DHA, FOX, CIT, and EBC (Figure 3). The majority of organisms were found to be positive for FOX-CIT. The distribution of AmpC producers in the various samples is shown in Table 3.

Pus and wound samples were the predominant sources of AmpC. *Klebsiella pneumoniae* is the predominant AmpC producer. (Table 4)

**DISCUSSION**

Treatment failures due to broad-spectrum antibiotics may be due to over expression of AmpC β-lactamases. AmpC β-lactamases encoded by plasmids were detected more frequently in *E. coli* and *Klebsiella* species. Several studies have revealed discrepancies in the types of AmpC β-lactamases among different geographical regions. AmpC beta-lactamase prevalence in different parts of India ranges from 8% to 47%. The AmpC producer rate was 32% in our study, which is similar to the results of other studies. In our study, all isolates were negative for the disc antagonism test (DAT), thus revealing the presence of only plasmid-mediated resistance in our isolates.

Cefepime is generally sensitive to AmpC producers. However, in our study, a few AmpC isolates were resistant to cefepime and cefepime-tazobactam. This may be due to a point mutation in the active site R2 loop, which results in variation that may act on cefepime in AmpC β-lactamase isolates and is known as extended-spectrum cephalosporinases. Many of the isolates in the present study were multidrug-resistant (MDR) because the genes are encoded on very large plasmids that are responsible for the multiresistant phenotype.

Presently, there are no recommended methods (CLSI guidelines) for screening and identification of AmpC producers in *Enterobacteriaceae*; however, several detection methods have been developed, such as the AmpC disc test, modified three-dimensional test, modified hodge test and inhibitor-based methods such as cloxacillin and boronic acid tests.
A reduction in susceptibility to cefoxitin antibiotic is considered a screening method for the detection of AmpC enzyme. However, ACC types are the only AmpC enzymes that can be missed when using the cefoxitin screening test. In this study, all 200 Enterobacteriaceae isolates that were not susceptible to oxyiminocephalosporin were screened for cefoxitin susceptibility. Of the 200 isolates, 149 (74.5%) were resistant to cefoxitin. However, only 62 (31%) of 200 isolates were positive in the AmpC disc test and MHT, and 64 (32%) were positive in the phenylboronic acid test. The cefoxitin-resistant isolates that were AmpC-negative in the phenotypic test may be due to ESBL or MBL production, or it may be due to decreased porin channels or increased efflux pump action, as demonstrated by porin Omp K 35 and Omp K36 loss in E. coli and K. pneumoniae isolates. This may also be due to the over expression of chromosomal AmpC genes caused by mutations. In several other studies conducted in India, 19-27% of cefoxitin-resistant isolates were found to be non-producers of AmpC β-lactamases. Porin loss is found to enhance the resistance of ESBL and AmpC β-lactamases and also leads to resistance to carbapenems.

The percentage of AmpC-producing isolates in the clinical samples was higher in samples from inpatients, indicating its nosocomial importance.

Among the AmpC disc and MHT tests, 24 isolates (38.7%) were E. coli, 31 (36.9%) were K. pneumoniae. There was one isolate each of K. oxytoca, Citrobacter spp., and Enterobacter spp. As well, two isolates each were from P. mirabilis and P. vulgaris. In other studies, 24% and 20.5% of E. coli were positive in the AmpC disc test. Among the 64 AmpC-positive isolates tested, AmpC genes were detected by PCR in 20 isolates (10%). The FOX-CIT family member was detected in 16 isolates (80%). FOX-CIT was detected in nine isolates of K. pneumoniae and seven of E. coli. The genotypes detected included DHA, EBC, CIT and FOX, respectively. MOX and ACC were not detected in the present study.

AmpC β-lactamases mediated by plasmids were detected more frequently in E. coli and Klebsiella species. Several studies have revealed discrepancies in the types of AmpC β-lactamases among different geographical regions. The discrepancy between phenotypic and genotypic tests may be due to the presence of other AmpC β-lactamase genes that continue to expand apart from the six gene families detected by PCR, or may be due to hyper producers due to over expression of the chromosomal gene. Of the 20 isolates identified as possessing plasmid-mediated AmpC, the sources were from urine, pus, and ETA samples.

Among the 64 isolates of AmpC producers from the phenotypic test, 20 could be assigned a genotype, which included CIT-FOX (16), EBC (2), FOX (1), and DHA (1).

AmpC beta-lactamases were identified in the urine (n=9), pus (n=6), wound (n=4), and ETA (n=1). Of these samples, 80% were from inpatients. Overall, the prevalence of AmpC genotypes among the tested Enterobacteriaceae was low (10%). FOX-CIT (80%) was the predominant genotype detected in this study. FOX-CIT was more common in K. pneumoniae (45%) than in E. coli, and EBC was detected in K. pneumoniae and E. coli, one DHA from P. mirabilis, and one FOX in K. pneumoniae.

In our study, the FOX group (n=17) was predominant (85%), followed by the CIT (n=16) (80%), EBC (n=2 (10%), and DHA (n=1(5%) groups. In a 2012 study by Mohamudha et al., the DHA gene was predominant, whereas, in the study by Manoharan et al., FOX-CIT was predominant, as was the case with the current study.

In our study, 40% of E. coli and 55% of Klebsiella pneumoniae were AmpC producers among the 64 phenotypic type positive clinical isolates, which is in accordance with the results of previous studies. However, 25.5% of E. coli and 39.1% of K. pneumoniae isolates have been reported in one study.

Plasmid-mediated AmpC β-lactamases were obtained from nine urine samples, six pus samples, four wound samples, and one ETA sample. Of these, 80% were from inpatients. The overall prevalence of common plasmid-mediated AmpC genotypes among the tested Enterobacteriaceae was low (10%). In the present study, CIT-FOX (80%) was the predominant genotype. In our
study, the FOX group (n=17) was predominant (85%), followed by the CIT (n=16) (80%), EBC (n=2 (10%), and DHA (n=1(5%)) groups. In a study by Mohamudha et al.15 the DHA-CIT genotype was predominant, whereas, in Manoharan et al.,18 the FOX-CIT genotype was predominant, which is in concordance with our study.

Many studies have shown that pAmpC β-lactamase-producing organisms are detected in prolonged hospitalized patients, and very recent studies suggest that it is found in tertiary care centers and outpatients, indicating its presence in the community. In this study, pAmpc was isolated from both inpatients and outpatients, indicating its presence both in hospitals and in the community.

CONCLUSION

In this study, *E. coli*, *K. pneumoniae* and *P. mirabilis* showed the presence of plasmid-mediated AmpC β-lactamases by PCR. In this study, many cefoxitin-resistant isolates were negative for AmpC, indicating the presence of other resistance mechanisms. FOX-CIT was the predominant AmpC β-lactamase gene, followed by EBC. The broad use of cephalosporins in empirical therapy in *Enterobacteriaceae* has increased the selective pressure among AmpC lactamase-generating isolates, increasing its prevalence. Hence, the identification of these enzyme-producing isolates will help in understanding the epidemiology of resistance mechanisms.

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CONFLICT OF INTEREST

The authors declare that there is no conflict of interest.

AUTHORS’ CONTRIBUTION

All authors listed have made a substantial, direct, and intellectual contribution to the work, and approved it for publication.

FUNDING

None.

DATA AVAILABILITY

All datasets generated or analysed during this study are included in the manuscript.

ETHICS STATEMENT

The study was conducted according to the guidelines of Institutional Human ethical committee, SBMCH, India with reference number 002/SBMC/IHEC/2013-99

REFERENCES


