Abstract

Background: Extended spectrum β-lactamase (ESBL) positive organisms are now a global health concern including in Bangladesh. These are associated with treatment failure, increased morbidity and mortality and increased health care costs. In this study, frequency of ESBL positive organisms in some health care centres in Dhaka city has been observed and their current status of antibiogram has also been observed. Objective: To observe the current status of antibiogram of ESBL positive organisms. Materials and Methods: This cross-sectional study was done in the Department of Microbiology, Bangladesh Institute of Health Sciences (BIHS) General Hospital, Dhaka, Bangladesh from March, 2012 to February, 2013. Only E. coli and Klebsiella spp. from pus and urine specimens were included in this study. Isolation, identification and antibiotic sensitivity of the organisms were done by standard procedures. Results: Organisms (Escherichia coli and Klebsiella spp.) isolated from urine and pus collected from different sites of 472 subjects were studied. Predominant organisms were Escherichia coli (82.8%) and remaining 17.2% were Klebsiella spp. ESBL positive organisms were higher in Escherichia coli (54.5%) than in Klebsiella spp. (44.4%) and higher in pus (77.0%) than in urine (49.1%) isolates. Imipenem is the most effective drug for treating ESBL positive organisms followed by colistin, tigecycline and piperacillin/tazobactam. Conclusion: Imipenem, colistin, tigecycline and piperacillin/tazobactam drugs should be kept reserved and used only when other effective drugs are not available so that emergence of resistance against these drugs is deferred. While reporting the culture and sensitivity tests, the ESBL positive organisms should be pointed out with comment like this – “The organisms are ESBL positive and resistant to penicillins, cephalosporins and monobactams”.

Key words: ESBL; Drug resistance; Cephalosporins

Introduction

Extended spectrum β-lactamase (ESBL) producing organisms are now a global threat and its treatment has become difficult because of their drug resistance over a wide spectrum of β-lactam and non-β-lactam drugs. These are emerging worldwide in community as well as in health care settings specially in Intensive Care Units (ICU). The ESBL isolates were first reported from Germany and England in 1983 and subsequently in...
USA in late 1988. In Bangladesh, it was reported first in 2001. ESBLs are enzymes that mediate resistance to β-lactam antibiotics, viz., penicillins, cephalosporins and monobactams, but do not affect cephamycins or carbapenems. Because of greatly extended substrate range, these enzymes are called extended spectrum β-lactamases. ESBL enzymes are most commonly produced by Escherichia coli (E. coli) and Klebsiella spp. and to some extent by other members of Enterobacteriaceae and Pseudomonadaceae.

ESBL organisms have continued to increase in varieties (more than 400 variants detected) and prevalence and are now a global health concern. The ESBL organisms have implications for clinicians and patients because these are associated with treatment failure, increased morbidity and mortality, poor outcomes, increased length of stay (LOS) in hospital and healthcare costs. So, it is important to treat it properly as soon as it is diagnosed. Aim of this study was to observe the current status of antibiogram of ESBL positive organisms and their prevalence in and around Dhaka city.

Materials and Methods

This cross-sectional study was done in the Department of Microbiology, Bangladesh Institute of Health Sciences (BIHS) General Hospital, Dhaka, Bangladesh from March, 2012 to February, 2013. Specimens were collected from inpatient and outpatient departments of BIHS General Hospital and nine more hospitals/centres located at different sites of Dhaka city.

Organisms isolated from urine and pus of 472 subjects were studied. There were 411 urine and 61 pus specimens. Pus was collected from diabetic foot lesions, post surgical infected wounds and traumatic wounds. Only E. coli and Klebsiella spp. were included in this study. Isolation, identification and antibiotic sensitivity of the organisms were done by standard procedures.

For detection of ESBL positive organisms, screening test was done along with routine sensitivity test. Positive screening tests were confirmed later by Phenotypic Confirmatory Test (PCT).

Screening test

Disks of ceftaxime (CTX), ceftazidime (CAZ) and amoxycillin/clavulanic acid (AMC) were placed in a line, placing AMC in the centre and other two on either side about 30 mm apart from centre to centre. Widening of cephalosporin’s inhibition zone adjacent to the disk containing AMC was regarded as ESBL positive.

Phenotypic confirmatory test (PCT)

In this method, two sets were used. One set consisted of ceftazidime (30 µg) disk and ceftazidime/clavulanic acid (30/10 µg) disk. Another set consisted of ceftaxime (30 µg) disk and ceftaxime/clavulanic acid (30/10 µg). The disks among the sets were placed at least 30 mm apart (centre to centre). When the zone of inhibition by combination disks (ceftazidime/clavulanic acid or ceftaxime/clavulanic acid) was >5 mm than the zone of inhibition produced by ceftazidime/ceftaxime alone, the test was taken as PCT positive.

Media for sensitivity

Muller Hinton agar (Oxoid)

Antibiotic disks used

All the isolates were tested for sensitivity to amoxycillin/clavulanic acid (20/10 µg), ceftaxime (30 µg), ceftazidime (30 µg), cefradine (30 µg), cefoxitin (30 µg), ampicillin (10 µg), amikacin (30 µg), imipenem (10 µg), gentamicin (10 µg), ciprofloxacin (5 µg), nalidixic acid (30 µg), nitrofurantoin (300 µg), doxycycline (30 µg), cotrimoxazole (1.25/23.75 µg), colistin (10 µg), tigecycline (15 µg) and piperacillin/tazobactam (100 µg/10 µg).

The above mentioned antibiotic discs were placed on Muller Hinton agar according to standard procedure.

Results

Isolation of ESBL positive organisms in BIHS General Hospital and different hospitals/centres located at different sites of Dhaka city has been shown in Table I. The highest isolation rate was in Wari Hospital (91.3%) followed by BIHS General Hospital IPD, Mirpur (81.4%) and Foot Care Hospital at Rampura (74.4%).

81
Frequency of organisms in different specimens is shown in Table II. E. coli was found more prevalent than Klebsiella spp. In case of E. coli, 54.5% were ESBL positive and in case of Klebsiella, 44.4% were ESBL positive. Frequency of ESBL positive organisms was more in case of pus (77.0%) than in case of urine (49.1%).

In Table III sensitivity pattern of ESBL positive and ESBL negative E. coli and Klebsiella spp. to different antibiotics has been shown. In case of E. coli, all (100%) of ESBL positive and ESBL negative strains were sensitive to imipenem and tigecycline. All (100%) ESBL negative strains were sensitive to colistin and piperacillin/tazobactam; but 93.9% and 90.6% of ESBL positive strains were sensitive to colistin and piperacillin/tazobactam respectively. For cefoxitin sensitivity of ESBL positive and negative E. coli was 85.9% and 100% respectively. Sensitivity to amikacin was about 76.5% and 77% against ESBL positive and ESBL negative strains respectively. Nitrofurantoin was used in case of urine isolates only where it was almost equally sensitive (85.2% vs 84.2%) to both of ESBL positive and ESBL negative strains. Sensitivity of ESBL positive organisms to other antibiotics was less; but ESBL negative strains were more sensitive than ESBL positive ones. All ESBL positive organisms were resistant to cephalosporins (1st to 3rd generation) while all ESBL negative organisms were sensitive to cephalosporins.

The scenario of antibiotic sensitivity was similar in case of Klebsiella spp. except that all (100%) were sensitive to colistin; 5.6%, 19.4% and 19.4% of ESBL positive strains were resistant to imipenem, tigecycline and piperacillin/tazobactam respectively. ESBL negative urine isolates showed higher (65.8%) sensitivity to nitrofurantoin than ESBL positive ones (31.6%).

Table I: Distribution of ESBL positive organisms in different sites of Dhaka city

<table>
<thead>
<tr>
<th>Centre names</th>
<th>Esch. coli</th>
<th>Klebsiella</th>
</tr>
</thead>
<tbody>
<tr>
<td>Isolated ESBL +ve Number (%)</td>
<td>Isolated ESBL +ve Number (%)</td>
<td></td>
</tr>
<tr>
<td>BIHSGH OPD, Mirpur</td>
<td>148 (77.0)</td>
<td>23 (17.4)</td>
</tr>
<tr>
<td>BIHSGH IPD, Mirpur</td>
<td>33 (26.8)</td>
<td>10 (90.0)</td>
</tr>
<tr>
<td>Savar</td>
<td>41 (18.4)</td>
<td>04 (14.3)</td>
</tr>
<tr>
<td>Uttara</td>
<td>32 (15.4)</td>
<td>07 (21.4)</td>
</tr>
<tr>
<td>Banani</td>
<td>27 (08.2)</td>
<td>07 (32.9)</td>
</tr>
<tr>
<td>Bashaboo</td>
<td>25 (10.0)</td>
<td>04 (16.0)</td>
</tr>
<tr>
<td>Jurain</td>
<td>18 (09.0)</td>
<td>01 (00.0)</td>
</tr>
<tr>
<td>Dhanmondi Hospital</td>
<td>20 (10.0)</td>
<td>06 (16.0)</td>
</tr>
<tr>
<td>Wari Hospital</td>
<td>23 (11.3)</td>
<td>00 (00.0)</td>
</tr>
<tr>
<td>Foot Care Hospital, Rampura</td>
<td>24 (19.2)</td>
<td>19 (32.9)</td>
</tr>
<tr>
<td>Total</td>
<td>391 (213)</td>
<td>81 (36)</td>
</tr>
</tbody>
</table>

BIHSGH, Bangladesh Institute of Health Sciences General Hospital

Frequency of organisms in different specimens is shown in Table II. E. coli was found more prevalent than Klebsiella spp. In case of E. coli 54.5% were ESBL positive and in case of Klebsiella 44.4% were ESBL positive. Frequency of ESBL positive organisms was more in case of pus (77.0%) than in case of urine (49.1%).

Table II: Frequency of ESBL positive Esch. coli and Klebsiella spp. in different specimens (N= 472)

<table>
<thead>
<tr>
<th>Organisms</th>
<th>Urine</th>
<th>Pus</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Isolates</td>
<td>ESBL+ve Number (%)</td>
<td>Isolates</td>
</tr>
<tr>
<td>Esch. coli</td>
<td>354 (51.7)</td>
<td>37 (81.1)</td>
<td>391</td>
</tr>
<tr>
<td>Klebsiella spp.</td>
<td>57 (33.3)</td>
<td>24 (70.8)</td>
<td>81</td>
</tr>
<tr>
<td>Total</td>
<td>411 (49.1)</td>
<td>61 (77.0)</td>
<td>472</td>
</tr>
</tbody>
</table>
Discussion

Reporting of ESBL positive organisms with their antibiogram is essential for proper treatment of the patients. In this study the latest scenario of antibiogram and frequency in some health care centres in and around Dhaka city have been presented.

From this study, it became evident that ESBL positive organisms are prevalent throughout the Dhaka city. The overall frequency of ESBL positive organisms was 52.8% (n=472). The highest frequency of ESBL positive organisms was in Wari Hospital (91.3%) followed by BIHS Gnereal Hospital IPD, Mirpur (81.4%) and Foot Care Hospital at Rampura (74.4%). The lowest frequency of ESBL positive organisms in Banani (32.4%), the area where most affluent people live, apparently gives a good relationship of economic status, improved environmental sanitation and infection control.

In this study, 472 specimens were tested, of which 249 (52.8%) were found positive for ESBL organisms. This is consistent with findings of Ahmed et al (53.4%, n=221) and Sashirekha (48.9%, n=225). Findings of Dalela (61.6%, n=219), Oliveira et al (61.1%, n=90) and Yasmin (71.3%, n=300) were slightly higher; the findings of Rahman et al (2.4%, n=14674) was very low. It is to be noted that in countries where there is good infection control practices and antimicrobial stewardship, the prevalence of ESBL +ve organisms is lower than other countries as was evidenced in a study in 2009 where global prevalence rate was found as follows: Latin America–44%, Asia/Pacific–22.4%, Europe–13.3% and North America–7.5%. However, prevalence of ESBL positive organisms is increasing throughout the world.

Predominating organism in our study was E. coli (82.8%, n=472) and the rest (17.2%) was Klebsiella spp.; 54.5% of E. coli (n=391) and 44.4% of Klebsiella spp. (n=81) were ESBL positive. These findings are consistent with the study of Rahman et al (54.4%, n=241) of which 43.2% were ESBL positive and 45.6% were K. pneumoniae of which 39.5% were ESBL positive. In the study of Sashirekha (28%, n=325) of which 52.8% were ESBL positive and 25.2% were K. pneumoniae of which 45.1% were ESBL positive. In the study of Dalela (73.5% of E. coli and 58.1% of K. pneumoniae were ESBL positive. In contrast, study of Oliveira et al showed K. pneumoniae (71.1%, n=90) as predominating organism following by E. coli (24.4%); 71.9% of K. pneumoniae and 36.4% of E. coli were ESBL positive. Study of Yasmin in Bangladesh also showed different scenario where predominating organism was E. coli (52%, n=300) followed by Proteus spp. (18.3%) and Klebsiella spp. (15%). In her study, Klebsiella spp. (80%) topped the list as ESBL producers followed by Proteus spp. (72%) and E. coli (67%).
In our study, organisms were isolated from two types of specimens – urine and pus. Urine isolates were 87.1% and pus 12.9%. Out of 411 urine isolates, the predominant organism was *E. coli* (86.1%) of which 51.7% were ESBL positive. *Klebsiella spp.* was 13.9% of which 33.3% were ESBL positive. Our findings are in consistence with that of Yasmin19 and Dalela17 except that the rate of ESBL positive organisms in urine isolates was higher (~70–80%).

In the present study, organisms were isolated from 61 pus specimens – 60.7% were *E. coli* and 39.3% were *Klebsiella spp.* Rate of ESBL positive *E. coli* and *Klebsiella spp.* was 81.1% and 70.8% respectively. This is consistent with the study of Yasmin19 and Dalela17 except that the rate of ESBL positive *E. coli* in the study of Yasmin19 (50%, n=42) and rate of ESBL positive *Klebsiella spp.* in the study of Dalela (50%, n=20)17 were lower.

ESBL producing organism is now a global threat1 and its treatment has become difficult because of drug resistance over a wide spectrum of β-lactam and non β-lactam drugs. In this study, in case of most antibiotics the difference of sensitivity between ESBL positive and ESBL negative organisms was statistically significant.

All ESBL positive organisms (100%) were found resistant to cephalosporins (except cephaplatin as discussed below) and monobactams while all (100%) ESBL negative organisms were sensitive to these drugs. Other authors showed varying degrees of sensitivity of ESBL producers to cephalosporins.1,16,17 It is to be noted that ESBL producers should be taken as resistant to all cephalosporins and monobactams even if these show sensitivity against some of the members. Cefepime is considered as a therapeutic option by some authors, but clinical data show high failure rate.20 Some patients with ESBL positive organisms were treated successfully with cephalosporins where organisms appeared sensitive in disc diffusion methods and had very low MIC.1,16,17,21,22 However, MIC is not done routinely in our department except in cases of multidrug resistance when higher than MIC level of antibiotic is targeted.

Currently, carbapenems are the most effective antibiotic for treatment of infections due to ESBL producing organisms as the outcome is better than treatment with other antibiotics.4,18,22,23 This has been true in this study also where all (100%, n=391) *E. coli* strains, both ESBL positive and negative, were sensitive to imipenem. This is consistent with the findings of Sasirekha (n=225)16, Yasmin (n=201)19 and Paterson23 who also found 100% of isolates sensitive to imipenem. Dalela (n=215)17 found 98.5% (n=135) of ESBL producer and 94% (n=219) of non-ESBL producers sensitive to imipenem and Ahmed et al1 found 98.3% of ESBL producer *E. coli* sensitive to imipenem. However, 5.6% of ESBL positive *Klebsiella spp.* in our study showed resistance to imipenem which is similar to the study of Hawser et al24 where 10% of ESBL positive *Klebsiella spp.* were resistant to imipenem; but it contrasts with the study of Sasirekha16, Yasmin19 and Paterson23 who found all ESBL producing *Klebsiella spp.* sensitive to imipenem. Imipenem resistance of ESBL positive *Klebsiella spp.* is increasing all over the world day by day.22

In this study all isolates except ESBL positive *Klebsiella spp.* were found sensitive to tigecycline; 80.6% of ESBL +ve *Klebsiella spp.* were sensitive to tigecycline. In other studies, it was found that tigecycline had excellent in vitro sensitivity but data reflecting clinical outcome is lacking.20,23,25 It was also reported that there was increased mortality rate in tigecycline-treated patients than in patients treated by other antibiotics.20 FDA warned health professionals and their medical care organizations about the increased risk of death when intravenous tigecycline is used. They suggested to reserve tigecycline for use only in situations when alternative treatments are not suitable.26

About 94% of ESBL positive *E. coli* were found sensitive to colistin in this study. All ESBL negative *E. coli* and all *Klebsiella spp.* were sensitive to colistin. Other studies also demonstrated its efficacy in treating multidrug resistant organisms including ESBL producers.20,23 Although recent studies have shown that it has acceptable effectiveness and has been used to treat infections due to multiresistant Gram-negative organisms, its use should be reserved as a last resort mainly for ESBL producing bacteria that are also resistant to aminoglycosides and carbapenems.25

In this study, <20% ESBL producers and >80% non-ESBL producers (both *E. coli* and *Klebsiella*) were sensitive to amoxycillin/clavulanic acid. In the study of Dalela17, 15.6% of ESBL producers and 23.8% of non-
ESBL producers were sensitive to amoxycillin/clavulanic acid.

In our study all (100%) ESBL negative *E. coli* and *Klebsiella spp.* were found sensitive to piperacillin/tazobactam whereas 90.6% of ESBL positive *E. coli* and 80.6% of ESBL positive *Klebsiella spp.* were sensitive to piperacillin/tazobactam. Dalela\textsuperscript{17} found 65.5% of non-ESBL producers and 72.6% of ESBL producers sensitive to piperacillin/tazobactam. Sasirekha\textsuperscript{16} found about 70% of non-ESBL producers and 60% of ESBL producers sensitive to piperacillin/tazobactam. Other studies showed that clinical experience with β-lactam/β-lactamase inhibitor combinations in the treatment of severe infections due to ESBL producers is limited and showed mixed results; but it may be an option of treatment for infections like urinary tract infections (UTI) caused by susceptible ESBLs.\textsuperscript{20-23}

The cephamycins (e.g. cefoxitin) which are also 2nd generation cephalosporins demonstrate markedly greater resistance to microbial degradation by β-lactamases than is normally displayed by the cephalosporins.\textsuperscript{22} In our study, 85.9% of ESBL positive and 100% of ESBL negative *E. coli* and 86.1% of ESBL positive and 93.3% of ESBL negative *Klebsiella* were sensitive to cefoxitin. However, clinical data regarding its use is scarce and clinical failure has been documented due to co-resistance.\textsuperscript{4,20}

In this study about 76.5% of ESBL positive and 77% ESBL negative strains of *E. coli* showed sensitivity against amikacin and comparatively it was higher in ESBL positive (83.3%) and ESBL negative (88.9%) *Klebsiella spp.* These findings are consistent with that of Yasmin\textsuperscript{19} who found 83% of *E. coli* and 81% of *Klebsiella spp.* sensitive to amikacin. Dalela\textsuperscript{17} found 64.5% of ESBL producers and 46.4% of non-ESBL producer organisms sensitive to amikacin. About 78% ESBL producers and 60% of non-ESBL producers were susceptible to amikacin in the study of Ahmed et al.\textsuperscript{1} But in contrast to these findings less sensitivity was observed in the study of Sasirekha\textsuperscript{16} where only 30% of ESBL producers and about 40% of non-ESBL producers were sensitive to amikacin.

In our study, 77.5% of ESBL positive *E. coli* and 55.6% of ESBL positive *Klebsiella spp.* were sensitive to gentamicin whereas 94.4% of ESBL negative *E. coli* and 97.8% of ESBL negative *Klebsiella spp.* were sensitive to this drug. In the study of Ahmed et al\textsuperscript{1} 34.8% of ESBL positive and 21.5% of ESBL negative *E. coli* were sensitive to gentamicin. However, aminoglycosides may be potentially useful in the treatment of complicated urinary tract infections due to ESBL-producing organisms; but these are not recommended as routine monotherapy for severe infections at other sites.\textsuperscript{20,23}

Nitrofurantoin was used in case of urine isolates only; both ESBL positive and ESBL negative *E. coli* were found almost equally sensitive (~85%). But *Klebsiella spp.* showed different figure – 31.6% of ESBL positive *Klebsiella spp.* and 65.8% of ESBL negative strains were sensitive to nitrofurantoin. Our findings are almost similar to that of Yasmin\textsuperscript{19} in case of *E. coli* (about 96% sensitive) but differed in case of *Klebsiella spp.* (about 95% sensitive). Ahmed et al\textsuperscript{1} found 54% cases sensitive in case of both ESBL positive and negative *E. coli*. Sasirekha\textsuperscript{16} found about 40% cases sensitive in case of both ESBL producer and non-producer organisms.

Sensitivity of ESBL positive organisms to other antibiotics was very low (doxycycline 29.1% and zero, cotrimoxazole 24.9% and 19.4% and ciprofloxacin 11.3% and 16.7% for *E. coli* and *Klebsiella spp.* respectively). But sensitivity of ESBL negative strains were significantly higher than ESBL positive organisms (p<0.05). These findings are consistent with that of Yasmin\textsuperscript{19}, Dalela\textsuperscript{17}, Ahmed et al\textsuperscript{1} and Sasirekha\textsuperscript{16}.

While reporting the culture and sensitivity tests, the ESBL positive organisms should be pointed out with comment like this – “The organisms are ESBL positive and resistant to penicillins, cephalosporins and monobactams”. In this study imipenem was found as the most effective drug for treatment of ESBL positive as well as ESBL negative organisms followed by colistin, tigecycline, piperacillin/tazobactam. However, these drugs should be kept reserved and used only when other sensitive drugs are not available so that emergence of resistance against these drugs is deferred. While selecting antibiotic against ESBL positive organisms, it should be remembered that this group of organisms are relatively less sensitive than ESBL negative organisms. Also in cases of ESBL positive organisms, it seems prudent to avoid β-lactam antibiotics as far as possible.
References


