

Antimicrobial susceptibility patterns against *Escherichia coli* and prevalence of extended-spectrum β -lactamases

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Abstract

Background: There is rapid increase in bacterial resistance to antibiotics resulted in increased morbidity and mortality among patients in hospitals. **Aims:** monitoring of antibiotic resistance to provide data for antibiotic therapy and resistance control prescription programs. **Materials and Methods:** Sixty seven clinical samples were collected from urine, stool, pus, wound of the patients and operating theater from some of Baghdad hospitals in 2010. Sixty two isolates of *Escherichia coli* were detected by bacteriological and biochemical tests, these isolates were submitted to hemolysis test and antibiotic susceptibility to 10 kinds of antibiotic disks on Mueller Hinton agar by disk diffusion method. The detection of β -lactamase production was also done and Extended-spectrum beta-lactamase (ESBL) for all the isolates. **Results:** Hemolysis test was positive for two isolates. All the isolates of *E. coli* were resistant to ampicillin and cephalothin (100%) and high resistance was observed to cephalixin (95.1%), tobramycin (90.3%), doxycycline (82.2%) and nalidixic acid (70.9%). Both ciprofloxacin and nitrofurantoin was (67.7%). Low resistance was noticed to amikacin (11.2%) and trimethoprim (8.1%). β -lactamase test was positive for 57 isolates (91.9%), while three isolates (4.8%) showed positive result for extended-spectrum beta-lactamase.

Conclusions: Isolates of *E. coli* showed high resistance to ampicillin, cephalothin and cephalixin. Low resistance was revealed to amikacin and trimethoprim. Most of the isolates were positive for β -lactamase test (91.9%) and (4.8%) of the isolates were positive for extended-spectrum beta-lactamase.

Key Words: *E. coli*, antimicrobial susceptibility, β -lactamase, ESBL.

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INTRODUCTION

Escherichia coli is one of the commensals in the human intestinal tract. As a commensal, it contributes to the maintenance of health of a person. However, *E. coli*, when enters into unnatural sites, can cause variety of infectious diseases such as urinary tract infections, wound infections, bacteraemia, meningitis and other soft tissue infections.^[1] Resistance to antibiotics is highly prevalent

in Bacterial isolates worldwide, particularly in developing countries. Routine monitoring of antibiotic resistance provides data for antibiotic therapy and resistance control prescription programs, making policy decisions and assessing the effectiveness of both.^[2] The rapid increase in bacterial resistance to antibiotics has resulted in increased morbidity and mortality among patients in hospitals, and in intensive care units in particular. Various measures have been proposed for alleviating this

situation, such as increased surveillance, improved physical controls like hand washing, and the substitution of empirically employed broad-spectrum antibiotics by narrow-spectrum drugs to which the infecting organism is known to be sensitive.^[3] Strains of Enterobacteriaceae that produce extended-spectrum β -lactamases (ESBLs) have emerged as significant pathogens. First reported in the mid-1980s, they were mainly found in *Klebsiella pneumoniae* and *Escherichia coli* although they can now be found in many other species.^[4]

The aim was to study the prevalence of ESBL producers among *Escherichia coli*, and their susceptibility pattern to antimicrobial agents.

MATERIALS AND METHODS

Isolation and Detection

Between April 2010- July 2010, a total of 67specimens were collected (urine, stool, pus, wound and operating theater) from some of Baghdad hospitals. Clinically 62 Gram negative bacilli belonging to *E.coli*.

The isolates were identified by standard techniques, bacteriological and biochemical tests. The specimens were plated on blood agar for hemolysis test, MacConkey agar and triple sugar iron agar. Catalase test, oxidase test, EPI system (Oxoid) were also used.^[5]

Antimicrobial agents susceptibility testing

Susceptibility to antimicrobial agents was assessed by the disc diffusion technique on Mueller-Hinton agar. Ten kinds of antibiotic discs (Amikacin, Ampicillin, Cephalixin, Cephalothin, Ciprofloxacin, Doxycycline, Nalidixic acid, Nitrofurantion, Tobramycin, Trimethoprim). Antimicrobial susceptibility testing was performed by Kirby-Bauer method and interpretation of results was as recommended by NCCLS.^[6] *E. coli* ATCC25922 was used as standard strain.

Beta-lactamase assay

Production of beta-lactamase test was done for the 62 isolates by using the rapid idometric method of WHO.^[7]

Extended-spectrum β -lactamase production test

The 62 isolates were subjected to ESBL screening. ESBL production was tested by using cefotaxime (30 μ g) ceftazidime (30 μ g) and augmentin (30/10 μ g) discs on Mueller-Hinton agar.

Organism was considered as an ESBL producer if there was a ≥ 5 mm increase in zone diameter around ceftazidime/augmentin disc compared to zone around ceftazidime disc alone.^[8]

RESULTS AND DISCUSSION

Among 67 clinical specimens, 62 isolates (92.5%) were identified as *E.coli* and their distribution as shown in table

1. The prevalence found in our study was similar to those found in a study conducted in *E. coli* prevalence in 2008 (92%),^[9] and (63%) in Nigeria from different clinical specimens.^[10] While the prevalence of *E.coli* in a recent study in Iraq from urine was (52%),^[11] and the current study was (75.8%). These wide variations in the prevalence, bacterial species, and antibiotic sensitivity could be due to variation in the study methodology, agent, host and environmental factors that exist.^[12]

Table 1. Distribution of 62 *E.coli* isolates among clinical specimens.

Specimens	Isolates Number	%
Operating theater	2	3.2
Pus	3	4.8
Stool	8	12.9
Urine	47	75.8
Wound	2	3.2

Hemolysis test was positive for the two isolates (3.2%) as shown in table 2 and these results are similar to the results demonstrated by Drews^[13] who stated that a lower prevalence of beta-hemolysis among nalidixic acid or ciprofloxacin-resistant *E. coli* urine isolates compared with susceptible isolates. Fluoroquinolone resistance is linked to loss of beta-hemolysis.^[13,14]

Table 2. Hemolysis positive *E. coli* isolates.

Specimens	Hemolysis positive isolates	(%)
Operating theater	1	1.6
Pus	-	0
Stool	-	0
Urine	1	1.6
Wound	-	0
Total	2	3.2

In antimicrobial agents susceptibility test all the isolates of *E. coli* (100%) were resistant to ampicillin and cephalothin. High resistance was observed to cephalixin (95.1%), tobramycin (90.3%), doxycycline (82.2%) and nalidixic acid (70.9%). Both ciprofloxacin and nitrofurantoin was (67.7%) as showed in table 3 and 4.

Table3. Antimicrobials susceptibility patterns of 62 *E. coli* isolates.

Antibiotics	Resistant isolates	Sensitive isolates
Amikacin (30 μ g)	7	35
Ampicillin (10 μ g)	62	-
Cephalixin (30 μ g)	59	3
Cephalothin (30 μ g)	62	-
Ciprofloxacin (5 μ g)	42	10
Doxycycline (30 μ g)	51	10
Nalidixic acid (30 μ g)	44	9
Nitrofurantion (300 μ g)	42	12
Tobramycin (10 μ g)	56	4
Trimethoprim(5 μ g)	5	11

Alhelfi study in Iraq^[12] and another study in Oman^[13] mentioned a high resistance rate in *E. coli* to ampicillin (99%) which is similar to our results (100%). Trimethoprim and amikacin showed the lowest resistance among the rest antimicrobials in this study, a new study in Kuwait showed similar result.^[15]

Table 4. The resistant *E. coli* isolates and their percentage.

Antimicrobial agents	Resistant isolates	(%)
Amikacin (30 μ g)	7	11.2
Ampicillin (10 μ g)	62	100
Cephalexin (30 μ g)	59	95.1
Cephalothin (30 μ g)	62	100
Ciprofloxacin (5 μ g)	42	67.7
Doxycycline (30 μ g)	51	82.2
Nalidixic acid (30 μ g)	44	70.9
Nitrofurantion (300 μ g)	42	67.7
Tobramycin (10 μ g)	56	90.3
Trimethoprim (5 μ g)	5	8.1

Table 5. Multiple resistance among *E. coli* isolates.

Number of antimicrobial agents	Number of resistant isolates
5	5
6	7
7	8
8	12
9	12
10	6

Table 6. Prevalence of β -lactamase and ESBL producing isolates.

Specimens	β -lactamase positive isolates		ESBL positive isolates	
	No.	(%)	No.	(%)
Operating theater	2	3.2	-	-
Pus	3	4.8	-	-
Stool	8	12.9	-	-
Urine	42	67.7	3	4.8
Wound	2	3.2	-	-
Total	57	91.9	3	4.8

The *E. coli* isolates in this study showed multiple resistances to more than 6 kinds of antimicrobials as displayed in table 5.

β -lactamase test was positive for 57 isolates (91.9%), while three isolates (4.8%) showed positive result for extended-spectrum beta-lactamase as in table 6. The prevalence of β -lactamase isolates found in our study was not similar to those found in a study conducted in 1997 in Iraq investigating *E. coli* producing β -lactamase prevalence, as the percentage was (60%) of the isolates.^[16] These wide variations in the prevalence could be due to variation in the study methodology, agent, host and environmental factors that exists.

An ESBL-producing *E. coli* prevalence of 4.8% was observed in this study.

AL-Helfi in a recent study in Iraq has reported that the prevalence of ESBL producing *E. coli* was (18%). ESBL producing *E. coli* was varied from (62.3%) as in mentioned by Shivaprakasha^[17] and (57%) in Ståle study.^[18] Direct comparisons are difficult due to selective sampling and different handling of the samples in the clinical laboratories. In UK the prevalence of ESBL producing isolates was (1.3%) by using the same technique.^[19] In this study ESBL producing isolates were resistant to more than 6 kinds of antimicrobial agents.

The available therapeutic options for the treatment of ESBL-associated infections are limited by drug resistance conferred by the ESBLs, along with frequently observed co-resistance to various antibiotic classes, including cephamycins, fluoroquinolones, aminoglycosides, tetracyclines, and trimethoprim /sulfamethoxazole.^[20]

The extended spectrum beta lactamase (ESBL) enzymes are plasmid-mediated enzymes capable of hydrolyzing and inactivating a wide variety of beta lactams, including third generation cephalosporins, penicillins and aztreonam.

Plasmids responsible for ESBL production carry resistance to many antibiotics like aminoglycosides, fluoroquinolones, tetracyclines, chloramphenicol and cotrimoxazole. The ESBL producing organisms are reported in increasing numbers worldwide.^[21]

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