

Antibiotic resistance pattern and molecular characterization of extended-spectrum β -lactamase producing enteroaggregative *Escherichia coli* isolates in children from southwest Iran

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Introduction: Enteroaggregative *Escherichia coli* (EAEC) has been implicated as an emerging cause of traveler's diarrhea, persistent diarrhea among children, and immunocompromised patients. The present study aimed to investigate the prevalence of antibiotic resistance, extended-spectrum β -lactamase (ESBL) production, and virulence factors of EAEC isolates obtained from Iranian children suffered from diarrhea.

Materials and methods: In this cross-sectional study, from March 2015 to February 2016, 32 EAEC isolates were collected from fecal samples of children aged <12 years with diarrhea in southwest of Iran. All EAEC isolates identified using phenotypic and molecular methods and the cell line adhesion assay. Antimicrobial susceptibility testing was determined using disk diffusion method. The presence of virulence factors and ESBL resistance genes were determined by polymerase chain reaction.

Results: Overall, 28.1% (9/32) of the isolates were positive for at least one of virulence genes. The most frequent gene was *aap* with a frequency of 96.9%. Neither *aafA* nor *aggA* gene was detected among all of the EAEC isolates. Antimicrobial susceptibility testing revealed the highest resistance rate to ampicillin (100%) and co-trimoxazole (100%), followed by ceftriaxone (81.3%). Further analysis revealed that the rate of ESBLs-producing isolates was 71.9% (23/32). Polymerase chain reaction screening revealed that 87.5% and 65.5% of EAEC isolates were positive for *bla*_{TEM} and *bla*_{CTX-M} genes, respectively, and 17 (53.1%) of isolates contained both *bla*_{TEM} and *bla*_{CTX-M} genes.

Conclusion: The high detection rate of ESBL-producing EAEC isolates accompanied with virulence genes highlights a need to restrict infection control policies in order to prevent further dissemination of the resistant and virulent EAEC strains.

Keywords: enteroaggregative *Escherichia coli*, diarrhea, adherence, antibiotic resistance, ESBLs, Iran

Introduction

Acute diarrheal diseases are an important health problem among children and are among the commonest causes of death among infants and children in developing countries.¹ About 70% of cases of acute diarrheal illness occurs in the first 5 years of life. Pathogenic bacteria and viruses are responsible for ~20% of the episodes of acute gastroenteritis in children.^{2,3}

Among the bacterial pathogens, diarrheagenic *Escherichia coli* (DEC) is a common cause of acute infectious diarrhea.⁴ DEC is classified into six groups based

on clinical associations, phenotypic assays, and virulence factors: enteroaggregative *E. coli* (EAEC), diffusely adherent *E. coli*, enteroinvasive *E. coli*, enteropathogenic *E. coli*, enterohemorrhagic *E. coli*, and enterotoxigenic *E. coli*.⁴ EAEC has been implicated as an emerging cause of traveler's diarrhea and persistent diarrhea among children and immunocompromised patients in both developing and developed countries.⁵ EAEC has also been associated with chronic intestinal inflammation, leading to malnutrition and growth retardation in infants.⁶

The pathogenesis of EAEC infection involves the adherence of the bacterium to the intestinal mucosa, forming a mucoid biofilm, and induces toxic effects on the intestinal mucosa, which result in diarrhea.⁷ The identification of EAEC depends on the HEp-2 adherence test, in which EAEC strains exhibit a "stacked-brick" appearance in a characteristic aggregative adherence (AA) pattern.⁷ The majority of EAEC strains carry a large (100-kb) plasmid, which encodes most putative EAEC virulence factors, including fimbrial adhesins, designated AA fimbria I (AAF/I, encoded by *aggA* gene), and AA fimbria II (AAF/II; encoded by *aafA* gene), which are responsible for the AA phenotype.⁸ The other plasmid-borne virulence factors include the enteroaggregative heat stable toxin (EAST; encoded by *astA* gene), dispersin secretory protein (encoded by *aap* gene), and plasmid-encoded toxin (Pet).⁸

In general, the first-choice agents for the treatment of EAEC infections are β -lactams, sulfonamides, and quinolones.⁹ However, in the recent years, therapeutic options are limited due to the emergence of *E. coli* strains resistant to third-generation cephalosporins, associated with the production of extended-spectrum β -lactamases (ESBLs).¹⁰ Bacterial strains producing ESBLs enzymes (TEM-1, SHV-1, and CTX-M-type) inactivate the drugs by hydrolyzing the β -lactam ring.¹⁰ The ESBL encoding genes are located on large plasmids, which can carry the genes for resistance to numerous other groups of antimicrobials. Thus, worldwide dissemination of plasmid-borne ESBLs among *E. coli* isolates is a global problem.¹⁰ The objectives of the present study was to investigate the prevalence of EAEC strains, virulence factors, antibiotic resistance, and ESBL production in children suffering from diarrhea in Ahvaz, southwest Iran.

Materials and methods

Ethics

The study was approved by the Research Ethics Committee of the Ahvaz Jundishapur University of Medical Sciences (No: IR.AJUMS.REC.1395.462), Ahvaz, Iran. Written informed consent was obtained from all the children's parents.

Study design and bacterial samples

In this cross-sectional study, from March 2015 to February 2016, 255 fecal samples were collected from children aged <12 years with diarrhea attending two teaching hospitals Golestan and Abuzar Children's Hospital, affiliated to Ahvaz Jundishapur University of Medical Sciences, southwest of Iran. The specimens were cultured on MacConkey agar and incubated at 37°C for 24 h. Subsequently, *E. coli* isolates were identified using standard microbiologic methods including Gram-staining, colony characteristics and reaction on Triple Sugar Iron agar, Simmons' citrate agar, Christensen's urea agar, Indole test, Methyl red, and Voges-Proskauer tests. The strains that confirmed as *E. coli* were stored in tripticase soy broth with 15% glycerol at -70°C for long preservation. *E. coli* strains were then screened for EAEC identification using molecular method and cell line adhesion assay.

DNA extraction and molecular assay

Genomic DNA was extracted from all *E. coli* isolates by boiling method as described previously¹¹ and subjected to polymerase chain reaction (PCR) after evaluating concentration and quality by measuring the absorbance of A260 and A280 nm with spectrophotometer and agarose gel electrophoresis, respectively. *E. coli* isolates were confirmed as EAEC by the amplification of *aggR* gene as previously described.¹² Subsequently, the presence of five virulence factors were determined by assessing the presence of *pCVD*, *aggA*, *ast*, *aap*, and *aafA* genes.¹³ Moreover, the presence of ESBL resistance genes, *bla*_{TEM}, *bla*_{PER}, *bla*_{CTX-M β} , and *bla*_{SHV} were determined by specific primers.¹⁴ Gene control strains were prepared from National *E. coli* Reference Laboratory, Pasteur Institute of Iran. The targeted genes and nucleotide sequences of the oligonucleotide primers used in this study were chosen as described in Table 1. PCR amplifications of the study genes were carried out in the following condition: initial denaturation at 95°C for 5 min, followed by 30 cycles of denaturation at 95°C for 60 s, annealing for 45 s (temperature was depending on the sequence of primers), extension at 72°C for 50 s and final extension at 72°C for 5 min. PCR amplifications for studied genes were carried out on a thermal cycler 5530 (Eppendorf master, Germany). The amplicons were separated on 1.5% agarose gel prepared in 1 \times TAE (Tris/Acetate/EDTA) buffer and visualized using ultraviolet light after staining with ethidium bromide (CinnaGen Co., Tehran, Iran).

Adhesion to HEp-2 cells

All of the *E. coli* isolates positive for the presence of *aggR* gene were confirmed as EAEC by aggregative adhesion to

Table 1 List of used primers in the present study

Primer	Oligonucleotide sequence (5' to 3')	Gene	Product size	Annealing	Reference
pCVD-F	CTGGCGAAAGACTGTATCAT	<i>pCVD432</i>	630	57	13
pCVD-R	AATGTATAGAAATCCGCTGTT				
aggR-F	GTATACACAAAAGAAGGAAGC	<i>aggR</i>	254	57	13
aggR-R	ACAGAATCGTCAGCATCAGC				
aggA-F	TTAGTCTTCTATCTAGGG	<i>aggA</i>	457	49	13
aggA-R	AAATTAATTCCGGCATGG				
aafA-F	TGCGATTGCTACTTTATTAT	<i>aafA</i>	242	56	13
aafA-R	ATTGACCGTGATTGGCTTCC				
aap-F	CTTGGGTATCAGCCTGAATG	<i>aap</i>	310	58	13
aap-R	AACCCATTCCGTTAGAGCAC				
astA-F	CCATCAACACAGTATATCCGA	<i>astA</i>	111	58	13
astA-R	GGTCGCGAGTGACGGCTTTGT				
TEM-F	GAGTATTCAACATTTCCGTGTC	<i>bla_{TEM}</i>	800	60	14
TEM-R	TAATCAGTGAGGCACCTATCTC				
PER-F	AATTTGGGCTTAGGGCAGAA	<i>Bla_{PER}</i>	925	48	14
PER-R	ATGAATGTCATTATAAAAGC				
CTX-M-F	CGCTTTGCGATGTGCAG	<i>Bla_{CTX-M}</i>	550	60	14
CTX-M-R	ACCGCGATATCGTTGGT				
SHV-F	CGCCTGTGTATTATCTCCCTGTTAGCC	<i>Bla_{SHV}</i>	843	62	14
SHV-R	TTGCCAGTGCTCGATCAGCG				

HEp-2 cells by a method described previously.¹⁵ Briefly, an overnight culture of *E. coli* was prepared, and then, a concentration of 10⁷ bacteria was incubated with monolayers of HEp-2 cells grown to 50% confluence on circular coverslips in wells of 24-well tissue culture plates. After 0.5–1 h of incubation at 37°C in 5% CO₂, the wells were gently washed three times with phosphate-buffered saline, and then, 200 µL of Dulbecco's minimum essential medium was added to each well, and the cultures were incubated at 37°C for 3 h in 5% CO₂. Fixation was done by 70% ethanol and stained with Giemsa stain. The aggregative adhesion was examined under the oil immersion lens of a light microscope. The HEp-2 cell lines were purchased from Razi Vaccine and Serum Research Institute, Karaj, Iran.

Antimicrobial susceptibility testing

Antimicrobial susceptibility testing was carried out on all EAEC isolates to 14 antibiotics by standard disk diffusion method on Mueller-Hinton agar medium (EMD Millipore, Billerica, MA, USA) as described by the Clinical and Laboratory Standards Institute (CLSI) guidelines.¹⁶ The antimicrobial agents used were gentamicin (10 µg), amikacin (30 µg), ciprofloxacin (5 µg), cefoxitin (30 µg), ceftazidime (30 µg), cefotaxime (30 µg), cefpodoxime (10 µg), ampicillin (10 µg), meropenem (10 µg), imipenem (10 µg), tetracycline (30 µg), ceftiofur (30 µg), trimethoprim/sulfamethoxazole (25 µg), and azithromycin (15 µg). *E. coli* ATCC 25922 strain was used for quality control purposes. Multiple-drug-resistant

(MDR) isolates (resistant to three or more of antimicrobials) were estimated according to previously described definitions.¹⁷ All isolates were tested for ESBL production using the combined-disk test using ceftazidime (30 µg) and cefotaxime (30 µg) disks and combination with clavulanic acid (10 µg) disk as described by CLSI guidelines.¹⁶ *E. coli* ATCC 25922 and *Klebsiella pneumoniae* ATCC 700603 were used as negative and positive control strains for ESBL production, respectively.

Statistical analysis

The analysis was performed by using SPSS™ software, version 21.0 (IBM Corporation, Armonk, NY, USA). The results are presented as descriptive statistics in terms of relative frequency. Values are expressed as the percentages of the group (categorical variables). Chi-square or Fisher's exact tests were used to determine the significance of differences. A difference was considered statistically significant if the *p*-value was <0.05.

Results

Virulence gene patterns and HEp-2 cell assay

Thirty-two EAEC isolated from 17 (53.1%) males and 15 (46.9%) females aged from 5 months to 11 years showed an overall prevalence of EAEC of 12.5% (32/255) in our region. All isolates were positive for *aggR* gene and adhered in a HEp-2 cell adherence assay in the AA pattern (Figure 1). The

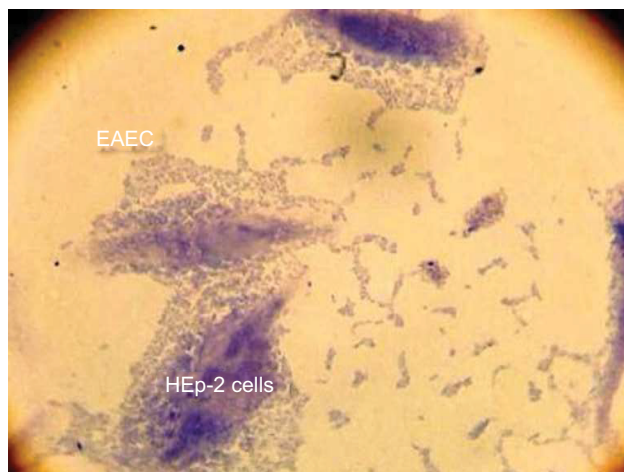


Figure 1 The AA pattern of EAEC to HEp-2 cells after 3 h of incubation.
Abbreviations: EAEC, enteroaggregative *Escherichia coli*; AA, aggregative adherence.

frequency of the detected virulence genes among EAEC isolates is shown in Table 2. The data revealed that 21 (65.6%), 7 (21.9%), and 31 (96.9%) of strains were positive for the *pCVD*, *astA*, and *aap* genes, respectively. The most frequent gene was *aap* with a frequency of 96.9%. Neither *aafA* nor *aggA* genes were detected among all of the EAEC isolates. Regarding to the coexistence of the virulence genes, our isolates showed three distinct virulence patterns (Table 2). The most prevalent combination was *pCVD-aap*, found in 21 (65.6%) strains. In 32 strains analyzed, six (18.7%) isolates were positive for *astA-aap* and four (12.5%) isolates were positive for *pCVD-astA-aap* genes.

Antibiotic resistance patterns and ESBL genes

The results of antimicrobial susceptibility testing of the 32 EAEC isolates to 14 antibiotics are summarized in Table 3.

From 32 confirmed EAEC isolates, all of them (100%) were resistant to ampicillin and trimethoprim/sulfamethoxazole, followed by 26 (81.3%) to ceftazidime, 25 (78.1%) to cefotaxime, and cefpodoxime, 25 (78.1%) to azithromycin, 17 (53.1%) to ceftazidime, 15 (46.9%) to tetracycline, 10 (31.2%) to gentamicin, 6 (18.8%) to ciprofloxacin, 5 (15.6%) to ceftazidime, 4 (12.5%) to amikacin, and 3 (9.4%) to meropenem. The results disclosed that the most effective antibiotic against EAEC isolates was imipenem with 100% susceptibility.

MDR profiles

According to the antimicrobial susceptibility testing, all 32 EAEC isolates were resistant to at least two antibiotics, and the majority of isolates ($n=31$, 96.9%) were

Table 2 The distribution of virulence and extended-spectrum β -lactamase (ESBL) genes

Virulence genes	Positive, N (%)	Negative, N (%)
<i>pCVD</i>	21 (65.6)	11 (34.4)
<i>aggA</i>	0	32 (100)
<i>astA</i>	7 (21.9)	25 (78.1)
<i>aap</i>	31 (96.9)	1 (3.1)
<i>aafA</i>	0	32 (100)
Coexistence of virulence genes		N (%)
<i>astA-aap</i>	6 (18.7)	
<i>pCVD-aap</i>	21 (65.6)	
<i>pCVD-astA-aap</i>	4 (12.5)	
ESBL genes	Positive, N (%)	Negative, N (%)
<i>bla</i> _{TEM}	28 (87.5)	4 (12.5)
<i>bla</i> _{CTX-M}	21 (65.6)	11 (34.4)
<i>bla</i> _{PER}	0	32 (100)
<i>bla</i> _{SHV}	0	32 (100)
<i>bla</i> _{TEM} <i>bla</i> _{CTX-M}	17 (53.1)	15 (46.9)

MDR with 21 different patterns (Table 4). The most prevalent resistance profile was XII (12.5%) (cefotaxime-cefpodoxime-ampicillin-ceftriaxone-trimethoprim/sulfamethoxazole-azithromycin).

Phenotypic results for ESBLs

The overall occurrence of ESBL-producing isolates was 71.9% (23/32) of EAEC. All isolates that were tested positive for ESBLs were also MDR. The results of antimicrobial susceptibility testing of the ESBL-producing EAEC isolates are summarized in Table 3.

Molecular assay of ESBL genes

PCR screening for the presence of ESBL genes showed that 28 (87.5%) and 21 (65.5%) of EAEC isolates were positive for *bla*_{TEM} and *bla*_{CTX-M} genes, respectively, and 17 (53.1%) of isolates contained both *bla*_{TEM} and *bla*_{CTX-M} genes. Neither *bla*_{SHV} nor *bla*_{PER} genes were detected among all of the EAEC isolates (Table 2). The detailed characteristics of all 32 studied EAEC isolates including virulence profile, MDR pattern, and ESBL production are summarized in Table 5.

Discussion

During the past decade, EAEC *E. coli* have been shown to cause persistent diarrhea and have received increasing attention globally.¹³ In addition to persistent diarrhea, EAEC have been isolated from acute sporadic cases and outbreaks worldwide, affecting children and adults.^{13,18} Previous studies have shown that EAEC strains are one of the most important

Table 3 The antibiotic susceptibility testing results of 32 EAEC isolates

Antibiotic	Total EAEC, N (%)			ESBL producing EAEC, N (%)		
	R	I	S	R	I	S
Gentamicin	10 (31.2)	0	22 (68.8)	6 (26.1)	0	17 (73.9)
Amikacin	4 (12.5)	1 (3.1)	27 (84.4)	3 (13)	1 (4.3)	19 (82.6)
Ciprofloxacin	6 (18.8)	1 (3.1)	25 (78.1)	5 (21.7)	1 (4.3)	17 (73.9)
Cefoxitin	5 (15.6)	0	27 (84.4)	0	0	23 (100)
Ceftazidime	17 (53.1)	4 (12.5)	11 (34.4)	12 (52.2)	3 (13)	8 (34.8)
Cefotaxime	26 (81.3)	0	6 (18.8)	20 (87)	0	3 (13)
Cefpodoxime	26 (81.3)	0	6 (18.8)	20 (87)	0	3 (13)
Ceftriaxone	26 (81.3)	1 (3.1)	5 (15.6)	21 (91.3)	0	2 (8.7)
Ampicillin	32 (100)	0	0	23 (100)	0	0
Meropenem	3 (9.4)	3 (9.4)	26 (81.3)	1 (4.3)	2 (8.7)	20 (87)
Imipenem	0	0	32 (100)	0	0	23 (100)
Tetracycline	15 (46.9)	0	17 (53.1)	7 (30.4)	0	16 (69.6)
Trimethoprim/sulfamethoxazole	32 (100)	0	0	23 (100)	0	0
Azithromycin	25 (78.1)	0	7 (21.9)	18 (78.3)	0	5 (21.7)

Abbreviations: EAEC, enteroaggregative *Escherichia coli*; ESBL, extended-spectrum β -lactamase.

Table 4 Antibiotic resistance phenotypic patterns of EAEC isolates

Resistance pattern	Phenotypic resistance	Number of resistant EAEC isolates (%)
I	AMP-SXT	1 (3.1%)
II	AMP-TET-SXT	2 (6.2%)
III	AMP-TET-SXT-AZM	1 (3.1%)
IV	GEN-CTX-CPDX-AMP-CRO-SXT-AZM	1 (3.1%)
V	GEN-CTX-CPDX-AMP-TET-SXT-AZM	1 (3.1%)
VI	GEN-AN-CIP-AMP-CRO-SXT-AZM	1 (3.1%)
VII	GEN-CAZ-CTX-CPDX-AMP-CRO-SXT-AZM	3 (9.3%)
VIII	GEN-CIP-CAZ-CTX-CPDX-AMP-CRO-SXT-AZM	1 (3.1%)
IX	GEN-FOX-CAZ-CTX-CPDX-AMP-CRO-TET-SXT-AZM	1 (3.1%)
X	GEN-FOX-CAZ-CTX-CPDX-MEM-AMP-CRO-TET-SXT-AZM	2 (6.2%)
XI	CTX-CPDX-AMP-CRO-SXT	1 (3.1%)
XII	CTX-CPDX-AMP-CRO-SXT-AZM	4 (12.5%)
XIII	CTX-CPDX-AMP-CRO-TET-SXT-AZM	2 (6.2%)
XIV	AN-AMP-TET-SXT-AZM	1 (3.1%)
XV	AN-CAZ-CTX-CPDX-MEM-AMP-CRO-SXT	1 (3.1%)
XVI	AN-CAZ-CTX-CPDX-AMP-CRO-SXT-AZM	1 (3.1%)
XVII	CIP-CAZ-CTX-CPDX-AMP-CRO-SXT-AZM	3 (9.3%)
XVIII	CIP-FOX-CAZ-CTX-CPDX-AMP-CRO-TET-SXT-AZM	1 (3.1%)
XIX	CAZ-CTX-CPDX-AMP-CRO-TET-SXT	2 (6.2%)
XX	CAZ-CTX-CPDX-AMP-CRO-TET-SXT-AZM	1 (3.1%)
XXI	FOX-CAZ-CTX-CPDX-AMP-CRO-TET-SXT-AZM	1 (3.1%)

Abbreviations: EAEC, enteroaggregative *Escherichia coli*; AMP, ampicillin; AN, amikacin; AZM, azithromycin; CTX, cefotaxime; CRO, ceftriaxone; CAZ, ceftazidime; CPDX, cefpodoxime; CIP, ciprofloxacin; FOX, cefoxitin; GEN, gentamicin; MEM, meropenem; SXT, trimethoprim-sulfamethoxazole; TET, tetracycline.

agents of diarrhea in Iranian children.^{13,19} Our study also revealed EAEC as a cause of sporadic diarrhea.

In this study, a total of 32 EAEC were isolated from 255 fecal specimens. The PCR results for the *aggR* gene disclosed good agreement with the HEp-2 cell adhesion assay, as 100% of *aggR*-positive isolates were confirmed as EAEC. Nowadays, different genes are used to recognize EAEC in molecular studies. One of them that is commonly used to

detect EAEC by PCR includes *aggR*.²⁰ In our research, the *aggR* PCR method compared to the HEp-2 cell culture assay indicated 100% sensitivity and 100% specificity.

In the present study, three diverse combinations of the virulence genes were found among the EAEC isolates and 31 (96.9%) of them were positive for *app* gene (Table 2). Neither *aafA* nor *aggA* genes were found in our study. In this regard, Aslani et al in the west of Iran reported 11 different patterns

Table 5 The detailed results of virulence genes, MDR, and ESBL gene patterns in the 32 EAEC isolates

EAEC No.	Virulence gene profile	MDR pattern	ESBL phenotype	ESBL gene pattern
1	pCVD-astA-aap	II	POS	TEM, CTX-M
2	pCVD-aap	IV	POS	TEM, CTX-M
3	pCVD-aap	II	NEG	TEM, CTX-M
4	astA	IX	NEG	TEM
5	pCVD-aap	XII	POS	CTX-M
6	pCVD-aap	XIV	NEG	TEM
7	pCVD-aap	XI	POS	TEM, CTX-M
8	pCVD-astA-aap	XV	POS	TEM, CTX-M
9	astA -aap	V	NEG	TEM
10	pCVD-aap	XVII	POS	CTX-M
11	pCVD-aap	I (not MDR)	NEG	TEM, CTX-M
12	pCVD-aap	XIX	POS	TEM, CTX-M
13	pCVD-aap	XVII	POS	TEM, CTX-M
14	pCVD-aap	III	POS	TEM
15	aap	X	NEG	TEM
16	pCVD-aap	XX	POS	TEM, CTX-M
17	pCVD-aap	VI	POS	TEM, CTX-M
18	pCVD-aap	VII	POS	TEM, CTX-M
19	pCVD-aap	VIII	POS	TEM
20	aap	XVIII	NEG	CTX-M
21	pCVD-aap	XVII	POS	TEM
22	pCVD-astA-aap	XIII	POS	TEM
23	aap	XXI	NEG	TEM, CTX-M
24	aap	XIX	POS	TEM, CTX-M
25	aap	X	NEG	TEM
26	aap	XII	POS	CTX-M
27	astA -aap	XII	POS	TEM, CTX-M
28	aap	XII	POS	TEM, CTX-M
29	pCVD-astA-aap	XIII	POS	TEM
30	aap	VII	POS	TEM, CTX-M
31	pCVD-aap	VII	POS	TEM
32	pCVD-aap	XVI	POS	TEM, CTX-M

Abbreviations: EAEC, enteroaggregative *Escherichia coli*; ESBL, extended-spectrum β -lactamase; MDR, multiple-drug resistant.

of the virulence markers in EAEC isolates.¹³ However, in contrast to our results, Aslani et al showed the prevalence of *aggA* and *aafA* in EAEC isolates.¹³ This might be due to epidemiologic differences between studied regions. A limitation of our study is that we only tried to search for the most common types of fimbriae. Our PCR targeted only AAFI and AAFII, but all isolates in this study were negative for these two fimbrial types indicating that the EAEC isolates in our region have fimbrial adhesins belonging to the other three established types or have novel types. In a study by Bouzari et al in the north of Iran, the prevalence of *astA*, *aggA*, and *aafA* among EAEC isolates obtained from children were reported as 8%, 38.8%, and 25%, respectively.¹⁹ Bafandeh et al in the north-west of Iran showed the prevalence of *aap* (88.6%), *astA* (83.5%), *aggR* (79.4%), *aafA* (46.4%), and *aggA* (5.1%) virulence determinants in EAEC isolates obtained from adult

patients with diarrhea.²¹ As a general concept from Iranian results and reports from other parts of the world, EAEC are heterogeneous, and no virulence factor has been identified as common to all EAEC strains.^{8,13,19-25}

Antimicrobial agents belonging to β -lactams family, particularly ampicillin and cephalosporins, and sulfonamides are widely used for the treatment of severe or persistent diarrhea in developing countries.⁹ However, the frequent use of these antimicrobial agents and the emergence of resistant strains have become a serious public health concern.²⁵ In our results, similar to previous reports from developing countries, the majority of isolates were resistant to ampicillin, cephalosporins, and co-trimoxazole.^{3,13,26,27} In Iran, cephalosporins are widely used due to their low degree of side effects. The high incidence of resistance to these agents may be due to the inappropriate and widespread use of antibiotics. Hopefully, based on our results, carbapenems, aminoglycosides, and fluoroquinolones can be used as an alternative for the treatment of EAEC-associated diarrhea in our area.

The emergence of MDR strains, particularly ESBL-producing *Enterobacteriaceae* is a global challenge for clinicians.²⁸ In the present study, we observed a high frequency of MDR (96.9%) and ESBL (71.9%) EAEC isolates which were resistance to the most tested antimicrobial agents. This high level of resistance is justified by the availability of medications without doctors' prescription from pharmacies in developing countries. Aslani et al closest to our findings showed the high rate of MDR EAEC (71.4%) in Iranian children.¹³ Reports from other parts of the country showed the prevalence of ESBLs producing clinical isolates of *E. coli* ranging from 22% to 74%. The ESBL production in EAEC strains from two Asian countries China and Bangladesh was reported as 50% and 49.1%, respectively.^{23,29} The differences in the prevalence of ESBLs producing isolates can be due to dissimilarities in geographical distribution, sample types, studied population, and hospital or community origin of isolates.

One of the major concern is the spread of ESBL-positive bacteria, which may mainly be due to the transfer of resistance genes via mobile genetic elements.¹⁰ ESBLs are enzymes most commonly derived from *bla*_{TEM} or *bla*_{SHV} but the prevalence of *bla*_{CTX-M} types has risen recently.³⁰ In our results, the genotype TEM was predominant with the prevalence of 87.5% followed by CTX-M type with 65.5%. The mechanisms of ESBL resistance in EAEC in Iran are poorly understood, and no similar study can be found. However, in agreement with our findings, several authors in our region showed the prevalence of *bla*_{CTX-M}, *bla*_{SHV}, and *bla*_{TEM} as the

main mechanisms responsible for ESBL production in clinical isolates of *E. coli* among Iranian patients.^{31–34} Meanwhile, some authors showed the global spread of *bla*_{CTX-M⁹} *bla*_{SHV²} and *bla*_{TEM} harboring DEC strains same as Iran.^{24,27,35,36}

Conclusion

In this study, the high detection rate of MDR and ESBL producing EAEC isolates accompanied with virulence genes highlights a need to restrict infection control policies to prevent further dissemination of the resistant and virulent EAEC strains. Hopefully, several locally available antibiotics still have promising effects against MDR isolates in our region. These findings provide experimental evidence for safe and effective management of EAEC associated infections.

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Disclosure

The authors report no conflicts of interest in this work.

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