Original research

Molecular investigation of extended-spectrum β-lactamases (ESBLs) genes in the *Salmonella* isolates obtained from children with acute diarrhea

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Abstract

Salmonellosis is an important public health concern among children in worldwide. Extended-spectrum β-lactams (ESBLs) cause resistance to clinically important beta-lactams which are generally used to treat invasive *Salmonella* infections. Therefore, the aim of this study was to investigate the presence of *SHV*, *TEM* and *CTX-M* genes in different strains of *Salmonella* isolated from children with acute diarrhea and to determine their resistance profile. In this cross-sectional study, 300 fecal samples were collected from children referred to the Amirkola Children’s Hospital, Babol, Iran. Antibiotic susceptibility testing was done according to the CLSI guideline. ESBLs-producing strains were identified using double disk synergy test method on the Mueller-Hinton agar plates. Multiplex-PCR was performed using oligonucleotide specific primers to detect of *SHV*, *TEM* and *CTX-M* genes. In total, 7% (n; 21/300) salmonella were isolated, which 61.9%, 28.6% and 9.5% were *Salmonella typhimurium*, *Salmonella enteritidis* and *Salmonella typhi*, respectively. The prevalence of the ESBL-producing isolates were 52.4%. M-PCR results showed that 42.8%, 38.1% and 14.3% of isolates were carried *CTX-M*, *TEM* and *SHV* genes, respectively. Also, 18.2% of isolates harbored *CTX-M*, and *TEM* genes, simultaneously. The high rate of ESBLs-producing *Salmonella* strains in the pediatric patients is an alarm. It is also recommended that alternative drugs be used with less resistance, which requires further investigation.

**Keywords:** Beta-lactamase, *Salmonella*, Diarrhea, Pediatric, PCR

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1. Introduction

*Salmonella*, a member of the *Enterobacteriaceae* family, are Gram-negative bacilli and spore-free that most normally transmitted from animals to humans [1]. These intestinal organisms, which are often considered potential pathogens for humans, have more than 2,500 different serovars that are divided into three distinct species including, *Salmonella typhi*, *Salmonella choleraesuis*, and *Salmonella enterica* [2]. *Salmonella enterica* sub-species *enterica* serovar *Enteritidis* (briefly *S. enteritidis*) is the most important cause of Non-typhoidal Salmonellosis. This disease is one of the most important infectious diseases between humans and animals, which is mostly related to the consumption of meat, poultry, eggs and milk [3]. So this bacterium is a food-borne pathogen. Salmonellosis can develop into one of the forms of typhoid fever, septicemia, and gastrointestinal infections (gastroenteritis). Gastroenteritis is the most common form of salmonellosis, often associated with symptoms such as fever, abdominal cramps, and diarrhea. The most common serotypes involved in this form is *S. typhimurium* and *S. enteritidis* [4]. The course of the disease lasts 4 to 7 days and many people recover without the need for antibiotics [5].

Today, one of the most important antimicrobials used in treatment of Salmonellosis is β-lactam agents. Unfortunately, for some reason, the addition of antibiotics to the diet of livestock, improper and arbitrary use of antibiotics, and lack of careful surveillance of drug administration have led to antibiotic-resistant strains [6]. The main problem in treating infections caused by these organisms is the emergence of multidrug resistance (MDR) strains, which often lead to prolonged hospitalization, increase mortality and mobility rates and increase the cost of treatment [7].

β-lactamases are a class of enzymes that hydrolysis the amide bond in the β-lactam ring of penicillins and cephalosporins. A common classification system is Ambler system which categorizes these enzyme into 4 dissimilar groups (A, B, C, D), based on the structure. Ambler type A (broad-spectrum beta-lactamases (ESBL)), C (Cephalosporinase), and D (oxacillinase) β-lactamases are categorized as a serine β-lactamases because they have ser amino acid at the enzyme's active site [8]. Ambler type B enzyme is classified as a metalloenzyme since the type need a divalent cations, usually zinc in active site to function. The spread of ESBLs is a global public health concern. ESBLs can develop resistance to antimicrobials such as 3th and 4th generation cephalosporins and monobactams [9]. The most common type of ESBLs in clinical samples is SHV, whose coding gene is located on a transferable plasmid and is therefore easily distributed among bacterial strains [10]. SHV β-lactamases are inhibited by clavulanic acid but not by EDTA. The TEM-1 was the first broad-spectrum β-lactamase, found not only in *Enterobacteriaceae* but also in *Pseudomonas aeruginosa* [11]. Among ESBLs, the CTX-M group has been described globally. The CTX-M gene is located on a transferable plasmid and is effective on the cefotaxime (a 3th-generation cephalosporin). These enzymes have the ability to hydrolyze cephalosporins and are inhibited by clavulanic acid, sulbactam, and tazobactam. CTX-M types are widely spread around the world. Although *Salmonella* serotypes are a significant etiologic agents of diarrhea in various parts of the world, very few reports are obtainable about the incidence of ESBLs among them [12]. So, the aim of the current study is to detect ESBL-encoded genes (TEM, SHV and CTX-M) in the *Salmonella* isolated from children with acute diarrhea and determination of the resistance pattern of these strains.

2. Materials and Methods

2.1. Bacterial isolation

In this cross-sectional study, which was performed over a period of 9 months (from the February to October 2019), in total 300 stool samples were obtained from the children’s suspected to Salmonellosis at the Amirkola Children's Hospital, Babol, Iran. In laboratory, each sample was enriched using culturing on the Selenite-F Broth (Mercury, Germany). After incubation on the 37°C for 24h, samples were cultured on the Eosin Methylene Blue (EMB) agar and Xylose Lysine Deoxycholate (XLD) agar (Merck, Germany) plates. Lactose negative colonies were identified by API 20E system (BioMerieux, France). All *Salmonella* spp. Isolates were serotyped with *Salmonella* specific O and H antisera using the slide agglutination test.

2.2. Antimicrobial susceptibility test

In according to the Standard Institute of Laboratory and Clinical Standards (CLSI), Antimicrobial susceptibility test was determined using agar disk diffusion method on the Mueller-Hinton agar (MHA) (Merck Co, Germany) for ceftazidime (30
µg), Aztreonam (30 µg), Imipenem (10 µg), Cefotaxime (30 µg), Ofloxacin (5 µg), Amikacin (30 µg) and Tetracycline (30 µg) (HiMedia, India) [13].

2.3. Double disk synergy test (DDST)
DDST was used for ESBL-producing isolates, cefotaxime 30 µg and ceftazidime 30 µg with and without clavulanic acid 10 µg (Mast, UK). The 0.5 McFarland standard turbidity of isolates were spread on the Muller Hinton agar (Merek, Germany). DDST was done by comparing the inhibition zone of disks containing cefotaxime and ceftazidime with and without clavulanic acid. When zones were distended < 5 mm around the disk comprising clavulanic acid the strain were considered as ESBL positive. *Klebsiella pneumoniae* ATCC 7006039 was used as a positive control [14].

2.4. Multiplex PCR
Cellular DNA was extracted by the boiling way formerly defined previously [15]. The M-PCR was done using specific primers to identify *SHV*, *TEM* and *CTX-M* genes (Table 1). The PCR reaction was performed at a volume of 25 µl. Each PCR reaction includes, 1.5 µL of template DNA, 12.5 µL of CinnaGen PCR Master Mix, 1.0 µL of each primer, and 9 µL of ddH2O. The M-PCR reaction was performed for 33 cycles in an Eppendorf MasterCycle Gradient thermocycler (Eppendorf, Germany) with the falling program, initial denaturation at 94°C for 6 min, denaturation at 95°C for 45 s, annealing at 57°C for 60 s, extension at 72°C for 72 min. M-PCR amplicons were electrophoresed in a 1% agarose/0.5 × TBE (45 mM-Tris-borate, 1 mM-EDTA) gel stained with 0.1 µl/ml Gel Red™ (Biotium, USA), then photographed under an UV trans-illuminator. The 100-bp DNA ladder (CinnaGen, Iran) was used as a molecular size marker.

### 3. Results
The mean age of the patients was 13.6±1.2 years (range from 4 to 14 years), which 45.3% (n; 163) were female and 45.6% (n; 137) male. In total, 7% (n; 21/300) *salmonella* species were collected, which 61.9% (n; 13/21), 28.6% (n; 6/21) and 9.5% (n; 2/21) were *S. typhimurium*, *S. enteritidis* and *S. typhi*, respectively. Antibiotic susceptibility test showed that all isolates were susceptible to imipenem (Table 2). Moreover, 42.8% (n; 9/21) of isolates were resistant to at least 3 different antibiotic classes and therefore considered MDR. Most MDR strains belonged to *S. typhimurium* serotype.

### Table 1. Oligonucleotide primer sequences used in this study

<table>
<thead>
<tr>
<th>Target gene</th>
<th>Amplicon size (bp)</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>bla</em>SHV</td>
<td>747</td>
<td>[16]</td>
</tr>
<tr>
<td><em>bla</em>TEM</td>
<td>445</td>
<td>[17]</td>
</tr>
<tr>
<td><em>bla</em>CTX-M</td>
<td>593</td>
<td>[17]</td>
</tr>
</tbody>
</table>

The results of DDST showed that 52.4% (n; 11) of isolates had ESBL-positive phenotype. The results of M-PCR showed that 42.8% (n; 9), 38.1% (n; 8) and 14.3% (n; 3) of isolates were carried *CTX-M*, *TEM* and *SHV* genes, respectively. Also, 18.2% (n; 2) of isolates carried genes *CTX-M*, and *TEM* simultaneously.
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4. Discussion

Diarrheal diseases caused by Salmonella are a worldwide health problem. Most people improve without specific treatment. Antibiotics are usually used only to treat cases suffering from severe infection. Patients should drink extra liquids as long as diarrhea lasts. In some cases, diarrhea may be so severe that the person needs to be hospitalized. Occasionally, infection may spread from the intestines to the bloodstream, and then to other parts of the body. In these cases, Salmonella can cause death unless the person is treated promptly with antibiotics. β-lactamases are extensively used in the treatment of salmonellosis [1]. In the present study, out of 300 stool samples, 7% Salmonella spp. were obtained, which is consistent with the study directed by Nodeh Farahani et al. [18]. This contradiction can be related to the number of samples and the year of study. In agreement with Soltan Dallal et al. [19], slide agglutination test results showed that 61.9%, 28.6% and 9.5% of isolates were belonged to the S. typhimurium, S. enteritidis and S. typhi serotypes, respectively. This variation may be resulted from distribution of different serotypes in various geographical areas.

Antimicrobial susceptibility test showed that all strains were susceptible to the imipenem, which was fully consistent with the study of Eshraghi et al. [20]. Also, 80.9% of isolates were resistant to tetracycline which was consistent with findings of Siourimé et al. [21]. In our study, with the exception of only one strain (S. enteritidis), all of them (20 isolates; 95.2%) were sensitive to ciprofloxacin, which is almost consistent with the results of Eshraghi et al. [20]. The researchers showed that the favorable effect of quinolones in combination with cephalosporins. Spiliopoulou et al. showed that all isolates were sensitive to ceftriaxone and ciprofloxacin, which is consistent with the present study [22]. This agreement could be due to the lack of need for antibiotics in the treatment of salmonellosis and its self-limitation.

In the DDST test, 52.4% of isolates were ESBLs producing phenotype. Also, the prevalence of CTX-M, TEM and SHV genes were 42.8%, 38.1%, and 14.3%, respectively. These data are inconsistent with the study of Tajbakhsh et al., since that all 174 Salmonella species were negative for TEM, CTX-M and SHV genes [23]. These results could be due to the geographical distance and the presence of other resistance mechanisms such as, efflux pumps, chromosomal beta-lactamase and porin loss. Of 138 S. enterica, 29.6% (n; 40) ESBL-producing isolates were collected by Ranjbar et al. [24]. The frequency of CTX-M, TEM and SHV genes was 12.3% (n; 17), 29.9% (n; 40) and 2.9% (n; 4), respectively, which was different with the present study. These results are important as an alarming in the development of resistance genes to the other susceptible strains and special attention to these strains. All 92 Salmonella spp. collected by Boisramé-Gastrin et al. were positive for TEM gene. So, 22.8% (n; 21/92) and 5.4% (n; 21/92) of isolates were harbored SHV and CTX-M genes, respectively [25]. The present study provided information on the frequency and antimicrobial susceptibility profile of Salmonella isolates in Iran. The high rates of ESBLs positive-Salmonella strains collected from pediatric

<table>
<thead>
<tr>
<th>Antimicrobial agents</th>
<th>S. typhimurium (n=13)</th>
<th>S. enteritidis (n=6)</th>
<th>S. typhi (n=2)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>S  I  R</td>
<td>S  I  R</td>
<td>S  I  R</td>
</tr>
<tr>
<td>Ceftazidime</td>
<td>3 (23.1) 1 (7.7) 9 (69.2)</td>
<td>1 (16.6) 0 5 (83.3)</td>
<td>0 0 2 (100)</td>
</tr>
<tr>
<td>Aztreonam</td>
<td>4 (30.7) 1 (7.7) 8 (61.5)</td>
<td>2 (33.3) 0 4 (66.6)</td>
<td>0 0 2 (100)</td>
</tr>
<tr>
<td>Imipenem</td>
<td>13 (100) 0 0</td>
<td>6 (100) 0 0</td>
<td>2 (100) 0 0</td>
</tr>
<tr>
<td>Cefotaxime</td>
<td>4 (30.7) 0 9 (69.2)</td>
<td>2 (33.3) 0 4 (66.6)</td>
<td>0 0 2 (100)</td>
</tr>
<tr>
<td>Oxofloxcin</td>
<td>8 (61.5) 1 (7.7) 4 (30.7)</td>
<td>4 (66.6) 1 (16.6) 1 (16.6)</td>
<td>1 (50) 1 (50) 0</td>
</tr>
<tr>
<td>Amikacin</td>
<td>2 (15.4) 2 (15.4) 9 (69.2)</td>
<td>5 (83.3) 0 1 (16.6)</td>
<td>0 0 2 (100)</td>
</tr>
<tr>
<td>Tetracycline</td>
<td>3 (23.1) 0 10 (76.9)</td>
<td>1 (16.6) 0 5 (83.3)</td>
<td>0 0 2 (100)</td>
</tr>
<tr>
<td>Ceftriaxone</td>
<td>12 (92.3) 0 1 (7.7)</td>
<td>5 (83.3) 0 1 (16.6)</td>
<td>2 (100) 0 0</td>
</tr>
<tr>
<td>Ciprofloxacin</td>
<td>12 (92.3) 0 1 (7.7)</td>
<td>6 (100) 0 0</td>
<td>2 (100) 0 0</td>
</tr>
</tbody>
</table>

Results presented as No. (%)
Abbreviations: S; susceptible; I: intermediate-resistant; R: resistant
cases is alarming and indicates a necessity to substitute the cephalosporins with an appropriate alternative.

Current limitations include the limited number of serotypes identified and resistance genes. It is also suggested that other resistance genes and other types of beta-lactamases should be identified.

Accurate diagnosis and rapid identification of ESBLs-producing Salmonella strains from pediatric patients is very important. Therefore, continuous monitoring of these resistances, especially at the endemic region, is an obvious thing that can prevent the spread of resistance to other strains and reduce the cost of treatment.

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Author Contributions
All authors contributed similarly to this work, and confirm the final version of manuscripts.

Conflict of Interests
Authors declare there is no conflict of interest.

Ethical declarations
All samples were obtained from patients as parts of routine sampling during their hospitalization period, so the regional ethical committee waived the need for informed consent.

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