Phylogenetic Distribution and Prevalence of Genes Encoding Class I Integrons and CTX-M-15 Extended-Spectrum \(\beta\)-Lactamases in Escherichia coli Isolates from Healthy Humans in Chandigarh, India

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Abstract

Escherichia coli is generally considered as a commensal inhabitant of gastrointestinal tract of humans and animals. The aim of this study was to gain insight on the distribution of phylotypes and presence of genes encoding integrons, extended \(\beta\)-lactamases and resistance to other antimicrobials in the commensal E. coli isolates from healthy adults in Chandigarh, India. PCR and DNA sequencing were used for phylogenetic classification, detections of integrase genes, gene cassettes within the integron and extended \(\beta\)-lactamases. The genetic structure of E. coli revealed a non-uniform distribution of isolates among the seven phylogenetic groups with significant representation of group A. Integron-encoded integrases were detected in 25 isolates with class 1 integron-encoded intI1 integrase being in the majority (22 isolates). The gene cassettes identified were those for trimethoprim, streptomycin, spectinomycin and streptothricin. The dfrA12-orf-aadA2 was the most commonly found gene cassette in intI1 positive isolates. Phenotypic assay for screening the potential ESBL producers suggested 16 isolates to be ESBL producers. PCR detection using gene-specific primers showed that 15 out of these 16 ESBL-producing E. coli harboured the blaCTX-M-15 Gene. Furthermore, molecular studies helped in characterizing the genes responsible for tetracycline, chloramphenicol and sulphonamides resistance. Collectively, our study outlines the intra-species phylogenetic structure and highlights the prevalence of class 1 integron and blaCTX-M-15 in commensal E. coli isolates of healthy adults in Chandigarh, India. Our findings further reinforce the relevance of commensal E. coli strains on the growing burden of antimicrobial resistance.

Introduction

Escherichia coli, a commensal of the intestinal tract of vertebrate gut, is one of most well studied model organisms and remains a workhorse in molecular biology. Though a prominent member of gut microbiota, the organism is causeative agent of life threatening diarrheal diseases, extra-intestinal infections and found to be associated with systemic diseases such as inflammatory bowel syndrome and colorectal cancer [1-5]. The composition and genetic structure of E. coli is largely dependent on climate conditions, dietary habits and host genetic factors. At present, E. coli populations are structured in seven major phylogenetic groups namely A, B1, B2, C, D, E and F. Strains responsible for extra-intestinal infection are far more likely to be member of phylogroups B2 or D than A or B1 [6]. Phylo-groups E and F contains strains, of which O157:H7 is the best known member, and form a sister group to phylo-group B2 respectively. The phylogroup C is closely related but distinct from phylo-group B1 [7].

Much of the studies on antimicrobial resistance and integron associated antimicrobial resistance has focused on pathogenic E. coli, even though preponderance of data has clearly documented the frequent presence of genes encoding extended-spectrum-\(\beta\)-lactamases and class 1 integrons in commensal E. coli strains from healthy human populations of diverse geographic locations [8–10]. In addition research has documented the exchange of the genes which confer drug resistance between normal flora and pathogenic E. coli and Salmonella and in turn the dissemination and the development of multidrug resistance in bacterial population [8,11,12].

Until now, data on the distribution of various phylotypes and occurrence of antimicrobial resistance determinants in commensal E. coli strains from healthy humans of Indian population are quite scarce; this motivated us to embark on the present study.

Materials and Methods

Ethics statement

This study and its procedures were approved and were in accordance with the guidelines of Institutional Bioethics Committee (IBSC), clearance number IBSC/2012-2/19. All the participants

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were explained the purpose of the study. Participation of all was voluntary and their written consent was obtained. The IBSC committee approved the consent procedure, and the records of the written consent are maintained in our lab.

Study design, sampling and E. coli identification

This study analyzed E. coli fecal samples of 102 healthy adults of age 18–55 years (80 males and rest are female) collected during the period of November 2012 to December 2013 from Chandigarh, an urban city in Northern region of India with humid subtropical climate. In the population under study 49 adults were either lacto-ovo-vegetarian or lactovegetarian and 53 were semi vegetarian that is they eat meat once a week. The maximum volunteers were research scholars and remaining was working class. None of the volunteers were suffering from any systemic or diarrheal diseases. They had not been treated with any antibiotics for at least 3 months preceding the collection of fecal sample.

In order to harvest E. coli isolates, freshly voided stools were collected from the subjects, placed into sterile sealable plastic containers and transported to the laboratory for processing within 2–4 h of collection. Stool samples were streaked on MacConkey agar (Difco, USA) and incubated aerobically for 18 h at 37°C. One lactose fermenting pink colony per sample was randomly picked. E. coli strains were further confirmed by complete 16S rDNA sequencing coupled with standard biochemical test obtained from National Institute of Cholera and Enteric Disease (NICED), Kolkata, India (a WHO accredited laboratory). The phylogenetic group of all 102 isolate was determined according to Clermont et al. [7].

Antimicrobial susceptibility testing and Screening for ESBL

All 102 isolates were tested for the susceptibility using the standard Kirby-Bauer disk diffusion method against 6 classes of antimicrobials at their breakpoint concentration in accordance with the Clinical and Laboratory Standards [13]. The 6 classes of antimicrobials used were aminoglycosides (kanamycin, gentamycin, streptomycin), sulfonamides (sulphafurazole, co-trimoxazole), tetracycline, fluoroquinolones (nalidixic acid, ciprofloxacin), chloramphenicol and beta-lactams (ampicillin, cefotaxime, ceftriaxone, cefazidime, azotreonam, cepfime imipenem meropenem). Isolates that were not susceptible to any of the oxymimic-cephalosporins were tested for ESBL production by the double disc synergy test and the combination disk test in accordance with the recommendations of the CLSI. Resistance to antimicrobials belonging to ≥5 classes of antimicrobials was considered as multidrug resistance (MDR). The isolates that showed resistance towards the meropenem was checked for carbapenemase production by modified Hodge test [14]. The antibiotic disks used were purchased from HiMedia Lab. Ltd., India.

Identification of resistance genes

Isolates that screened positive for ESBL production were subjected to molecular screening for five β-lactamase genes namely, blaTEM-1, blaOXA-10, blaCTX-M, blaTEC and blaSHV. DNA was isolated and PCR was performed using specific primer pairs to screen for β-lactamase genes [15]. DNA sequencing of resistance genes was performed to validate their identities and further subgrouping. All Sequences were analyzed using the BLAST software (http://www.ncbi.nlm.nih.gov/blast).

All the isolates that showed resistance or moderate susceptibility to tetracycline, chloramphenicol and sulfonamides were checked for the presence of genes encoding resistance to these antimicrobials. Tetracycline resistant samples were checked for presence of tetA, tetB, tetC, tetD, tetE and tetG whereas chloramphenicol resistant strains were examined for the presence of cat1, cat2, cat3 and cmlB [16]. Sulfonamides resistant samples were tested for sul1, sul2 and sul3 [16].

Integron analysis and characterization of inserted gene cassettes

PCR was performed to check the presence of class 1, 2 and 3 integrons in all 102 isolates using primers specific for the integron integrase genes intI1, intI2 and intI3 [17]. Strains that were positive for intI1 and/or intI2 gene/s were subsequently subjected to PCR for amplification of the variable region of class 1 and class 2 integrons respectively [16]. Amplified products corresponding to the gene cassettes within the integrons were then subjected for nucleotide sequencing and confirmed by BLAST software.

Results

Phylogenetic grouping of E. coli

Our data reveal the predominance of phylogenetic group A (55 isolates) followed by phylogenetic groups B1 (24 isolates), B (7 isolates), F (7 isolates), B2 (4 isolates), C (3 isolates) and E (1 isolate). One isolate remain unclassified as it was negative for all the genes in quadruplex PCR.

Prevalence of antimicrobial resistance and distribution of CTX-M extended-spectrum β-lactamases in E. coli from fecal samples

Susceptibility test on all 102 isolates against 18 antimicrobial drugs showed the carriage of multidrug resistance in 37 isolates. Resistance was most frequently observed to ampicillin (57 isolates), nalidixic acid (48 isolates), sulphafurazole (41 isolates), amoxyclav (36 isolates), co-trimoxazole (32 isolates) and tetracycline (31 isolates). In contrast, resistance to ciprofloxacin, streptomycin, azotreonam, kanamycin, gentamicin and chloramphenicol were found in 26, 16, 14, 7, 5 and 2 isolates respectively (Figure 1A). It should be highlighted that, resistance to the antimicrobials belonging to the class of fluoroquinolones and sulphonamides was much higher than the aminoglycosides and chloramphenicol. 25 isolates were resistant to at least one of the third generation cephalosporins used in the study. 14 isolates were resistant to all of the third and fourth generation cephalosporins tested in this study. Isolates resistant to cefotaxime, ceftriaxone and ceftazidime were 24, 16 and 17 respectively, whereas 14 isolates were resistant to cepfime- a fourth generation cephalosporin. Resistance towards carbapenem was not common, 6 isolates were resistant to meropenem however, none of the isolate was positive in modified Hodge test used to screen carbapenemase. No isolate showed imipenem resistance.

We analyzed the antimicrobial resistance combination pattern in special reference to classes of antimicrobial. The most common resistance pattern were fluoroquinolones-beta lactams (33 isolates), sulphonamides-beta lactams (30 isolates) and sulphonamides-flur- oquinolones (23 isolates).

Phenotypic assay (double disc synergy test and the combination disk test) for screening the potential ESBL producers suggested 16 isolates to be ESBL producers. We determined that 15 out of these 16 ESBL-producing E. coli harbored the blaCTX-M-15 gene (Table 1). blaTEM-1 enzyme was also identified in seven of these blaCTX-M-15 producers. blaOXA-10, blaSHV-like and blaVEB-like were not found in any of the isolates. Apart from resistance to beta
lactams, ESBL producers showed higher resistance to fluoroquinolone, tetracycline and sulfonamide class of antibiotics (Figure 1B).

Integron and gene cassettes in E. coli isolates: Prevalence of class 1 integrons

Previous studies have clearly evident the emergence of integron associated antibiotic resistance in intestinal E. coli strains. We observed the prevalence of class 1 integron (22 isolates) over class 2 integron (4 isolates) while rarely described class 3 was absent in our commensal E. coli isolates. It should be noted that one isolate from our study harbored both class 1 and 2 integron genes. Blast analysis of sequenced products of variable region of class 1 and class 2 integrons demonstrated the presence of six different cassette combinations for nine different genes. These genes encode for the resistance to trimethoprim (dfrA1, dfrA5, dfrA7, dfrA12 and dfrA17), aminoglycosides (aadA1, aadA2 and aadA5) and streptothricin (satA1) (Table 1). Out of 22 intI1-positive isolates 7 carried dfrA12-oxf-aadA2 cassettes; three isolate harbored dfrA17-aadA5 gene cassette; one was found to contain dfrA11-aadA1 cassette, two isolates contained dfrA5 and one had dfrA7. All 4 intI2-positive strains showed the presence of cassette array usually associated with the integron on Tns7: dfrA1-sat1-aadA1. In our isolates, gene cassettes found in the integron conferred resistance to trimethoprim, streptomycin and streptothricin. Taken together, our analysis revealed that presence of integron in commensal E. coli isolates from healthy subjects are free of recent antimicrobial exposure.

Detection of genes encoding resistance to tetracycline, chloramphenicol and sulfonamide

Out of 41 tetracycline resistant or moderate susceptible isolates, 25 isolates were carrying tetA. 16 and 1 isolate was PCR positive for tetB and tetD respectively. One of these isolate was carrying both tetA and tetB. Both of the chloramphenicol resistant isolates were carrying cat1 gene. Isolates that were resistant for either of both co-trimoxazole and sulphafurazole were tested for three allelic variation of the sul gene. The sul gene was found in 36 isolates. The sul2 was the most common gene present in 26 isolates, whereas sul1 was found in 18 isolates. 8 of these isolates were carrying both sul1 and sul2 genes. No isolate was tested positive for the presence of sul3. Further classification of sul allele showed that 20 out of 25 integrase 1 and/or 2 positive isolates carried atleast one sul gene. 16 out of 36 sul positive isolates were not tested positive for the presence of any intI1 genes.

Discussion

In this work, we were driven by a desire to understand the phylogenetic architecture and antimicrobial resistance pattern of commensal E. coli from healthy population in Chandigarh, India. In this connection, our data are in congruence with previous reports and clearly witnessed the predominance of strains belonging to phylogeny A as well as prevalence of class 1 integron and CTX-M-15 ESBL positive strains in healthy Indian population [18-21]

It should be highlighted that our geographically distinct population is significantly different in terms of percentage of isolates in each phylogenetic groups [18-20]. We observed the prevalence of E. coli strains belonging to phylogeny A and B1 over other phylytype reflecting similar pattern as seen in case of other population [18,22,23]. Interestingly, we found more D over B2 whereas earlier study describes more B2 phylotype than D [7,20]. This alteration could be explained in terms of differences in dietary habits, environmental conditions or host genetic factors. Our data further support the earlier reports where it was concluded that in tropical climate phylogroup A dominate over B1 and prevalence of B2 and D is very low, suggesting the role of environmental factors in defining the genetic structure of E. coli in a given population [19].

The prevalence of class I integron in our isolates is in agreement with previous report where frequency of class 1 integron is reported much more than class 2 [21]. In the class 1 integron-positive isolates we found eight different gene cassettes in five
Table 1. Distribution of isolates positive for ESBL gene, integron and gene cassettes among phylogenetic groups of E. coli.

| Phylogenetic group | A | B1 | B2 | C | D | E | F | All groups |
|--------------------|---|----|----|---|---|---|---|-----------|
| Extended spectrum b-lactamase | | | | | | | | |
| CTX-M-15 | 91 | - | - | 3 | - | 2 | 1 | 5 |
| Integrons and Gene cassette array | | | | | | | | |
| Gene cassette arrays in positive isolates | | | | | | | | |
| intI1 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| dfrA12-orfF-aadA2 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| dfrA17-aadA5 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| dfrA1-aadA1 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| dfrA5 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| dfrA7 | 22 | 3 | - | 1 | - | - | 1 | 7 |
| Gene cassette arrays in positive isolates | | | | | | | | |
| intI2 | 22 | 3 | - | 1 | - | - | 1 | 7 |

Data are given as numbers of isolates.

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The simple, low-cost survey of resistance in commensal bacteria from patient hospitalized in New Delhi, India [29]. Since then, increasing occurrence of this particular type of ESBL has been reported in clinical E. coli isolates from different parts of the world [30]. The prevalence of CTX-M-15 producers in this study is strikingly high. As CTX-M-15 is known to have peculiar association with community-onset E. coli infections [31] therefore it may be concluded that CTX-M-15-producing E. coli have already been established in our area. Phylogenetic B group and serotype O25:H4 has been particularly associated with CTX-M-15, but, in this study 9 of the 15 CTX-M-15 were from phylogroup A and only isolate was of phylogenetic group B. It should be noted that one isolate of phenotypically confirmed ESBL do not have any ESBL encoding gene, this isolate probably produce other ESBL enzyme that were not checked during this study. In addition like earlier studies, we observed high prevalence of fluoroquinolones resistance among ESBL-producing isolates [32,33]. Many reasons have been suggested in literature for this association but the most possible explanation is the presence of genes of the two resistance mechanisms on the same plasmid. We also observed that all six meropenem resistant isolates are carbenapenemase negative but ESBL producers. This acquired resistance could be attributed to many factors including the loosening of outer membrane porins in ESBL positive isolates resulting in reduced carbenapenem uptake [34].

Acquisition of tetracycline, sulphonamide and chloramphenicol genes are also of concern. In contrast to earlier reports where tetB is reported as the most common gene found [35,36], our population observed the prevalence of tetA. One isolate tested positive for both tetA and tetB. The difference in the distribution pattern of tet gene suggest ecosystem-specific reservoir for resistance gene. Our isolates showed the prevalence of sul2 gene over sul1, the observed pattern of gene frequency distribution (sul2>sul1>sul3) corroborate previous studies [37]. Though literature has witnessed the sul1 gene as consistent marker for the presence of class 1 integron, still we found 11 isolates that were carrying the mtl1 but not the sul1 gene [38]. Similar finding has been reported earlier where it has been suggested that it is either because of the loss of sul1 gene region from class 1 integrons or that this gene is carried on another genetic context in the isolates [39]. 14 isolates that were tested resistant or moderately susceptible against the sulphonamides did not harbor any sul gene pointing to the probability of other sul alleles in these isolates or other resistance mechanism.

There are reports documenting ESBL positive E. coli isolates primarily from clinical settings in India [40–42]. In this regard, our study remains first report to reveal the status of commensal E. coli strains from healthy Indian population and supports the emerging theme of commensal flora mediated antibiotic resistance burden. The simple, low-cost survey of resistance in commensal bacteria
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Author Contributions

Conceived and designed the experiments: SR CD. Performed the experiments: CD SM. Analyzed the data: SR CD. Contributed reagents/materials/analysis tools: SR CD. Wrote the paper: SR.
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