

## Abstracts

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**Background** The present study is comparison of commensal and uropathogenic *Escherichia coli* (UPEC) pathotypes according to multiple virulence genes by multiplex PCR.

**Methods** Thirty two virulence factors including adhesions group, protectin related genes, common toxins related to UPEC, phylogenetic typing, Pathogenicity Island and miscellaneous genes which are involved in *Escherichia coli* pathogenicity mechanisms were examined by PCR.

**Results** The PCR assay results identified *vat*(96 vs. 4%), PAI IV536(85 vs. 41%), PAI ICFT073(78 vs. 25%), *kpsMTII*(71 vs. 42%), *ompT*(68 vs. 53%),*usp*(52 vs. 3%), *sat*(47 vs. 3%), *picU* (41 vs. 1%), PAI IICFT073(38 vs. 13%), PAI I536(37 vs. 7%), *fliC* (*H7*(24 vs. 9%)), *hlyD*(22 vs. 2%), *focG*(21 vs. 3%), *cnf1*(21 vs.0%), *gafD*(20 vs. 2%), *cdtB* (19 vs. 7%),*bmaE*(17 vs. 5%), PAI II J96(11 vs.3%), were more frequent virulence markers in urinary *E.coli* isolates than commensal *E.coli*, respectively. The frequency of *afa*(7 vs.10%), *fimH*(92 vs. 97%), *kpsMT(K1)* (41 vs. 57%),*kpsMTIII*(11 vs.5%), *rfc* (O4 LPS) (9 vs. 2%), PAI II536(20 vs. 24%), PAI III536(3 vs.0%), PAI I J96(53 vs. 43%), *ibeA*(17 vs. 26%) markers was almost similar in both of them. *cvaC*(64 vs. 18%)was most frequent marker in commensal *E.coli*. PCR phylotyping revealed higher prevalence of commensal *E. coli* in groups A and D while uropathogenic strains were mainly found in subgroup B2.

**Conclusion** Based on these results, the potential for Commensal *E.coli* to act as human UPEC or as a reservoir of virulence genes for UPEC should be considered.

922 **PHYLOGENIC TYPING OF URINARY *E. COLI* ISOLATES AND COMMENSAL *E. COLI* IN PATIENTS WITH UTI IN MOFID CHILDREN HOSPITAL IN IRAN**

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**Background** The objectives of this study were to define the phylogenetic groups in urinary and commensal *E. coli* isolated from urine and stool samples of hospitalized children and to determine the pattern of resistance to antibiotics.

**Method** A total of 100 urine and stool samples were processed during the study period from September 2009 to August 2010. Samples were cultured using standard microbiological techniques. Biochemical testing was used to identify the organisms, *E. coli* isolates were tested for phylogenetic grouping by using triplex PCR and antibiotic susceptibility test done by the Kirby Bauer method.

**Results** Phylogenetic group B2 and D were predominant in urinary samples (54% and 34% respectively). Phylogenetic group A, D and B2 were found in decreasing order of 41%, 26% and 16% respectively in the stool samples.

Following resistance patterns were observed in urinary *E. coli* isolates vs. commensal *E. coli*, respectively: nitrofurantoin(2% vs. 8%); imipenem (2% vs. 1%); amikacin (4% vs. 3%); ciprofloxacin(8% vs. 5%); nalidixic acid (8% vs. 27%); amoxicillin (16% vs. 20%); ceftazidime (12% vs.7%); amoxicillin-clavulanic acid (14% vs. 9%); gentamicin (22% vs. 12%); cefpodoxime (48% vs. 19%); cefotaxime (45% vs. 27%); co-trimoxazole (61% vs. 82%); and aztreonam (78% vs. 16%); Multi-drug resistance (MDR = resistance in>3 drugs) was most commonly associated with UPEC isolates.

**Conclusion** Although group B2 *E. coli* strains were uncommon in stool samples, as they are highly virulent they still represent a potential reservoir for urinary tract infection. Resistance to most antimicrobials is high both in UPEC and commensal strains.

923 **MULTIPLEX PCR FOR DETECTION OF SHIGA-LIKE TOXIN AND PLASMID ANTIGEN H GENES OF DIARRHEAGENIC *ESCHERICHIA COLI***

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**Background and Aims** Despite the fact that Shiga-like toxin *E. coli* has been identified as a major etiologic agent of children with *diarrhea worldwide*, few studies have been performed to evaluate the etiology of Shiga-like toxin-producing *Escherichia coli* (SLTEC) in children with diarrhea, in Iran. The aim of this study was to evaluate the etiology of Shiga-like toxin-producing *Escherichia coli* (SLTEC) in children with diarrhea, in Iran.

**Method** A total of 300 stool specimens from children of 300 children with diarrhea were tested for the detection of *E. coli*, according to standard methods. Out of 300 specimens, 39 were identified as diarrheagenic *E. coli*, and subjected to multiplex polymerase chain reaction (MPCR) for detection of *stx1/stx2*, *eae* and *ipaH* genes. We designed a single multiplex polymerase chain reaction (MPCR) for the detection of target genes in diarrheagenic *Escherichia coli*.

**Results** EPEC was the dominated strain (55.6%) among the tested isolates, followed by EHEC (25%) and EIEC (19.4%) strains.

**Conclusions** Shiga-like toxin *E. coli* has been identified as a major etiologic agent of children with diarrhea in Iran. our method proved to be specific and rapid in detecting virulence genes from Shiga toxin-producing (*stx1*, *stx2*, and *eae*), enteropathogenic (*eae*), enteroinvasive (*ipaH*) *Escherichia coli* in stool samples, and able to simultaneously detect of diarrheagenic *E. coli* strains in a single reaction.

924 **DIAGNOSIS AND PROGNOSIS OF CONGENITAL TOXOPLASMOSIS**

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**Aims** Congenital toxoplasmosis can cause neurological impairment and ocular disease. To describe clinical profile of infants with suspected congenital toxoplasmosis.

**Methods** Observational study of infants born to mothers with a suspected infection with *Toxoplasma gondii* during pregnancy between 2002 and 2011.

Serological tests were performed at birth: Toxoplasma specific antibodies IgA, IgM, IgG by Enzyme Immune Assay (EIA), Enzyme Linked Fluorescent Assay (ELFA), Western Blot (WB) tests and WB-IgG compared analysis for mother-infant pairs. Infants underwent cranial Ultrasound Scanning, funduscopy examination, Auditory Brainstem Response, and periodic clinical evaluations.

**Results** One hundred thirty-one infants Toxoplasma IgG-positive at birth were evaluated; 118/131 (90%) become IgG-negative at 12 months of life.

Congenital toxoplasmosis was confirmed in 13/131 infants (9.9%). Transmitters pregnant women seroconverted in the third trimester (mean 28±8weeks).

IgM-ELFA test was positive in 9/13 infants; in 4/13 infants IgM positivity was detected by WB test (negative IgM-EIA/ELFA). Three of 6 infants had a different IgG-WB reactivity compared to their mothers.

Six of 13 infected infants (46%) were symptomatic at birth: 2/13 infants developed chorioretinitis; 4/13 had a pathological neuroimaging (4/4 cerebral calcifications, 1/4 ventriculomegaly). None had



## 923 Multiplex PCR for Detection of Shiga-Like Toxin and Plasmid Antigen H Genes of Diarrheagenic *Escherichia Coli*

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