Abstracts

922 PHYLLOGENIC 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 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, phylogenic 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 IV356(85 vs. 41), PAI ICF703(78 vs. 25%), kpsMTHH (71 vs. 42%), ompT (68 vs. 53%), usp (52 vs. 3%), sucA (41 vs. 1%), PAI ICF703 (88 vs. 13%), PAI ICF703 (57 vs. 7%), fliC (H7 (24 vs. 9%)), hlyD (22 vs. 2%), focC (21 vs. 3%), cnfI (21 vs. 0%), gafD (20 vs. 2%), cdtB (19 vs. 7%), bmaE (17 vs. 5%), PAI II (96 vs. 11 vs. 3%), were more frequent virulence markers in urinary E. coli isolates than commensal E. coli, respectively. The frequency of sfa (7 vs. 10%), intimH (92 vs. 97%), kpsMT (K1) (41 vs. 57%), kpsMTHH (11 vs. 5%), rfc (O4 LPS) (9 vs. 2%), PAI IIS56 (20 vs. 24%), PAI IIIS56 (3 vs. 0%), PAI I (96 vs. 53 vs. 43%), bvaC (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.

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 dominant 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, fundoscopy 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±2 weeks).

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...