

Difference of virulence factors, clinical presentation according to phylogenetic group in uropathogenic *E. coli* strains isolated Korean patients

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Introduction

- *Escherichia coli* (*E. coli*) is most predominant organism causing 95% of community acquired urinary tract infection.
- *E. coli* is made up of four phylogenetic groups; A, B1, B2, D. It has been demonstrated that most uropathogenic *E. coli* belong to phylogenetic group B2 and D.
- But there are only a few studies about the difference of virulence factors, clinical characteristics according to phylogenetic group.
- We aimed to find out the prevalence of virulence factors of uropathogenic *E. coli* and difference of clinical manifestation according to phylogenetic group in Korean urinary tract infection patients.

Methods

- The study was performed on 133 *E. coli* isolates recovered from blood, urine specimens from patients with urinary tract infections who were diagnosed at Keimyung university of Dongsan medical center from February 2015 through May 2016.
 - The bacteria were confirmed as *E. coli* by VITEK system.
 - Phylogenetic analysis was conducted by targeting three genetic markers *chuA*, *yjaA*, *TSPE4.C2*.
 - Twenty nine virulence factors were identified by multiplex PCR.
 - Primers' sequence were as published, provided by other investigators.
- Primers were sorted into 5 pools according to amplicon size.
- Pool 1: *PAI*, *papA*, *fimH*, *kpsMT III*, *papEF*, and *ibeA*
- Pool 2: *fyuA*, *bmaE*, *sfa/focDE*, *iutA*, *papG allele III*, and *K1*
- Pool 3: *hlyA*, *rfaE*, *nfaE*, *papG allele I*, *kpsMT II*, and *papC*
- Pool 4: *gafD*, *cvaC*, *cdtB*, *focG*, *traT*, and *papG allele II*
- Pool 5: *papG allele I*, *papG alleles II and III*, *afa/draBC*, *cnfI*, *sfaS*, *K5*

Results

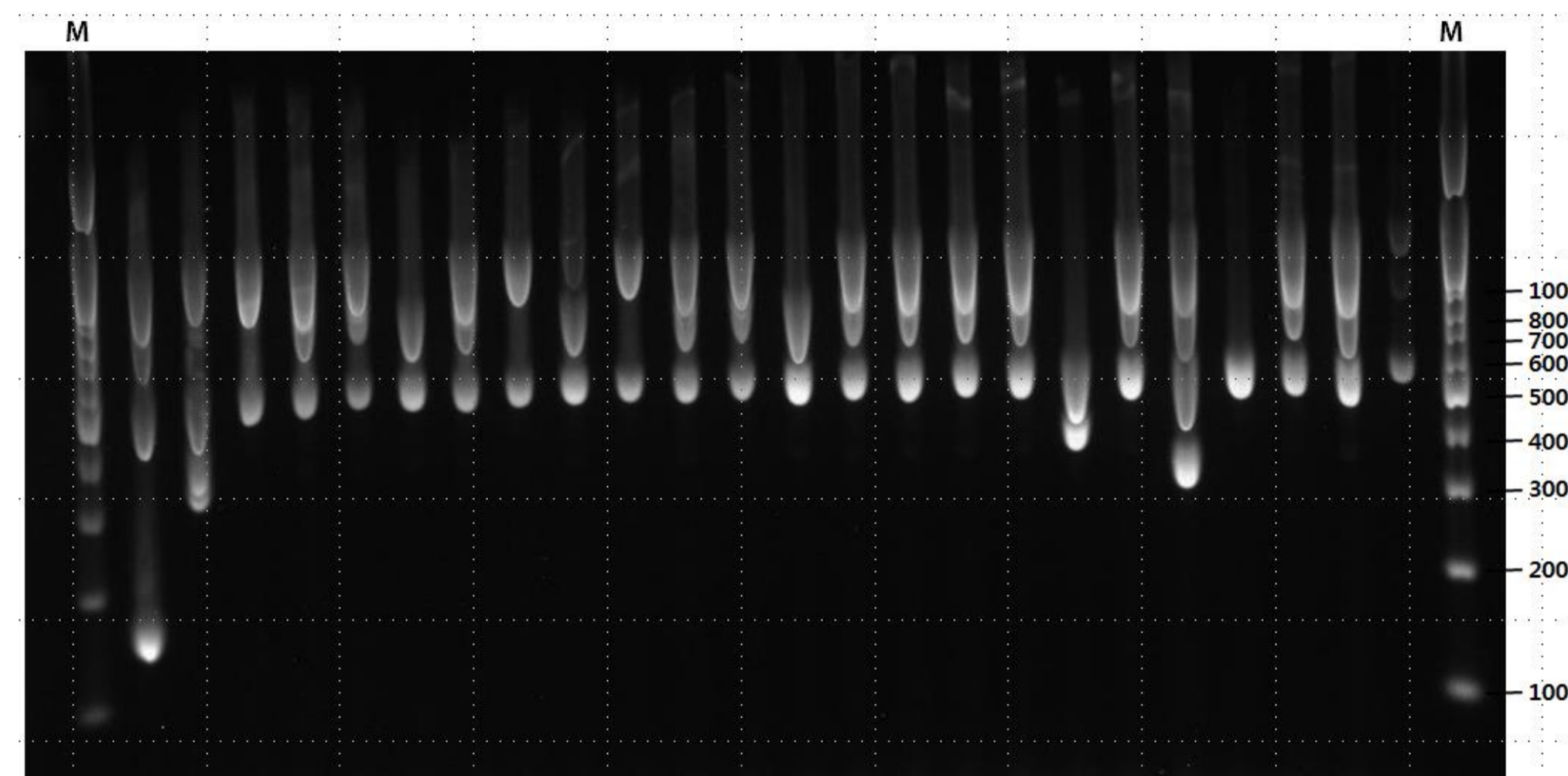


Figure 1. Agarose gel showing polymerase chain reaction (PCR) products from multiplex virulence factors PCR of pool 1 set.

- The phylogenetic group analysis reveals that most of uropathogenic *E. coli* are group B2 and D.
- The proportion of phylogenetic group of uropathogenic *E. coli* as follows: B2 (101, 75.9%), D (27, 20.3%), B1 (3, 2.3%), A (2, 1.5%).

Gene name	N (%)	Gene name	N (%)	Gene name	N (%)
<i>fimH</i>	131(98%)	<i>K5</i>	41(31%)	<i>kpsMT III</i>	3(2%)
<i>fyuA</i>	129(97%)	<i>hlyA</i>	40(30%)	<i>papG allele III</i>	3(2%)
<i>traT</i>	102(77%)	<i>cnfI</i>	40(30%)	<i>bmaE</i>	2(2%)
<i>PAI</i>	99(74%)	<i>K1</i>	35(26%)	<i>rfaE</i>	2(2%)
<i>iutA</i>	95(71%)	<i>sfa/focED</i>	18(14%)	<i>nfaE</i>	2(2%)
<i>papC</i>	92(69%)	<i>focG</i>	15(11%)	<i>sfaS</i>	1(1%)
<i>papG</i>	91(68%)	<i>afa/draBC</i>	13(10%)	<i>gafD</i>	0
<i>papA</i>	88(66%)	<i>cvaC</i>	10(8%)	<i>cdtB</i>	0
<i>papG allele II</i>	88(66%)	<i>papEF</i>	8(6%)	<i>papG allele I</i>	0
<i>kpsMT II</i>	77(58%)	<i>ibeA</i>	8(6%)		

Table 1. Prevalence of virulence factors of uropathogenic *E. coli*

- Among the Virulence factors, *fimH*, *fyuA*, *traT*, *PAI*, *iutA*, *papC*, *papG*, *papA* were most frequently observed.

Results

- We compared clinical presentations and virulence factors of group B2 (n=101) and D (n=27).
- Phylogenetic group B2 was more related with virulence factors which were *sfa/focED*, *focG*, *fyuA*, *hlyA*, *cnfI*. Group D was more related with *nfa* (Table 2). Number of virulence factor was higher in group B2 (11(7-12) vs. 8(6-10), $p=0.001$).
- Group B2 and D were showed similar clinical presentation and complication. But, Group D was mostly community origin infection (26(96.3%) vs. 80(79.2%), $p=0.043$). Group B2 was more related with healthcare-associated infection and antimicrobial resistance.
- Most of extended spectrum β -lactamase producing *E. coli* was group B2 (42/45, 93%). K1 serotype was prevalent in group B2 and K5 was higher in group D.

	Virulence factor	Group B2 n=101	Group D n=27	<i>p</i>	Virulence factor	Group B2 n=101	Group D n=27	<i>p</i>	
Adhesion	<i>papA</i>	70 (69.3%)	16 (59.3%)	0.36	Iron metabolism	<i>fyuA</i>	101 (100%)	26 (96.3%)	0.211
	<i>fimH</i>	100 (99%)	26 (96.3%)	0.379		<i>iutA</i>	74 (73.3%)	19 (70.4%)	0.81
	<i>papEF</i>	5 (5%)	3 (11.1%)	0.364	Toxin	<i>hlyA</i>	38 (37.6%)	1 (3.7%)	0.001
	<i>sfa/focED</i>	18 (17.8%)	0	0.013		<i>cvaC</i>	9 (8.9%)	0	0.203
	<i>nfaE</i>	0	2 (7.4%)	0.043		<i>cnfI</i>	40 (39.6%)	0	0.001
Protection	<i>afa/draBC</i>	8 (7.9%)	5 (18.5%)	0.146	<i>kpsMTIII</i>	2 (2.0%)	1 (3.7%)	0.512	
	<i>focG</i>	15 (14.9%)	0	0.039	<i>kpsMTII</i>	60 (59.4%)	17 (63%)	0.827	
	<i>papC</i>	73 (72.3%)	17 (63%)	0.353	<i>traT</i>	81 (80.2%)	19 (70.4%)	0.299	
	<i>papG</i>	73 (72.3%)	17 (63%)	0.353	Others	<i>PAI</i>	96 (95%)	3 (11.1%)	0.001
	<i>papG alleleII</i>	70 (69.3%)	17 (63%)	0.643		<i>ibeA</i>	8 (7.9%)	0	0.202

Table 2. Association of virulence factors of uropathogenic *E. coli* according to phylogenetic group B2 and D.

conclusion

- In our study, phylogenetic group B2 and D show different characteristics.
- Phylogenetic group B2 has more virulence factors than group D. Group B2 showed high presentation of adhesion related virulence factors(S fimbriae, F fimbriae). Higher presentation of resistance and healthcare-associated infection also noted.
- Presentation of virulence factors in Korean patients was different from other geographic regions. We need more studies to find out the microbiologic characteristics of uropathogenic *E. coli*. It may help to find out new target of therapy for urinary tract infection.