

# Recap Exercise 2

# ResFinder

## Known Mutations

**parE**

No mutations found in parE

**parC**

No known mutations found in parC

**folP**

No mutations found in folP

**gyrA**

Mutation	Nucleotide change	Amino acid change	Resistance	PMID
gyrA p.S83A	TCG → GCG	S → A	Quinolones, Fluoroquinolones	<a href="#">15848289</a>

**pmrB**

No known mutations found in pmrB

**pmrA**

No mutations found in pmrA

**16S\_rrsB**

No known mutations found in 16S\_rrsB

**16S\_rrsH**

No known mutations found in 16S\_rrsH

**gyrB**

No mutations found in gyrB

**ampC**

No known mutations found in ampC

**16S\_rrsC**

No known mutations found in 16S\_rrsC

**23S**

No known mutations found in 23S

*GyrA* and *GyrB* in *E. coli* are difficult cases as it is often a **combination** of mutations that leads to resistance

- According to ResFinder-3.0, the *E. coli* strain that caused the German outbreak has **one** point mutation in *gyrA* (S83A)
- Mutations at this position are usually related to resistance to quinolones/ flouroquinolones (ciprofloxacin, nalidixic acid)
- According to Grad et al, 2012, the *E. coli* strains are resistant to nalidixic acid, while susceptible to ciprofloxacin
- For ciprofloxacin, studies have shown that single mutations in *gyrA* only lead to modest increment in resistance - isolates would still be considered clinically susceptible. Only a second mutation in *gyrA* or a mutation in *parC* leads to a clinical level of resistance (PMID: 15942878)
- ResFinder-3.0 currently outputs resistance according to the environmental break point, not the clinical breakpoint

# ResFinder

## Unknown Mutations

parC		
Mutation	Nucleotide change	Amino acid change
parC p.E62K	GAA → AAA	E → K

gyrA		
No unknown mutations found in gyrA		

pmrB		
Mutation	Nucleotide change	Amino acid change
pmrB p.D283G	GAC → GGC	D → G
pmrB p.Y358N	TAC → AAC	Y → N

16S_rrsB	
Mutation	Nucleotide change
16S_rrsB r.80A>C	A → C
16S_rrsB r.89T>G	T → G
16S_rrsB r.93T>C	T → C
16S_rrsB r.250A>T	A → T
16S_rrsB r.253A>T	A → T
16S_rrsB r.273T>A	T → A
16S_rrsB r.474G>A	G → A

16S_rrsH	
Mutation	Nucleotide change
16S_rrsH r.80A>C	A → C
16S_rrsH r.89T>G	T → G
16S_rrsH r.90C>T	C → T
16S_rrsH r.93T>C	T → C
16S_rrsH r.250A>T	A → T
16S_rrsH r.253A>T	A → T
16S_rrsH r.273T>A	T → A
16S_rrsH r.1036A>G	A → G
16S_rrsH r.1120T>C	T → C

ampC		
Mutation	Nucleotide change	Amino acid change

## SerotypeFinder-1.1 Server - Results

H type						
Serotype gene	%Identity	Query/HSP length	Contig	Position in contig	Predicted serotype	Accession number
<i>fliC</i>	100.00	1050 / 1050	Supercontig_1.4	70365..71414	H4	<a href="#">AJ605764</a>

O type						
Serotype gene	%Identity	Query/HSP length	Contig	Position in contig	Predicted serotype	Accession number
<i>wzx</i>	100.00	1278 / 1278	Supercontig_1.4	215774..217051	O104	<a href="#">KB021482</a>
<i>wzy</i>	99.64	1113 / 1113	Supercontig_1.4	218280..219392	O104	<a href="#">AF361371</a>

**Predicted Serotype: O104:H4**

[extended output](#)

[Results as text](#)

[Results tab separated](#)

[Hit in genome sequences](#)

[Serotype gene sequences](#)

**Selected %ID threshold: 85.00 %**

**Selected minimum length: 60 %**

**Input Files: *escherichia\_coli\_c227-11.fsa***

### CITATIONS

For publication of results, please cite:

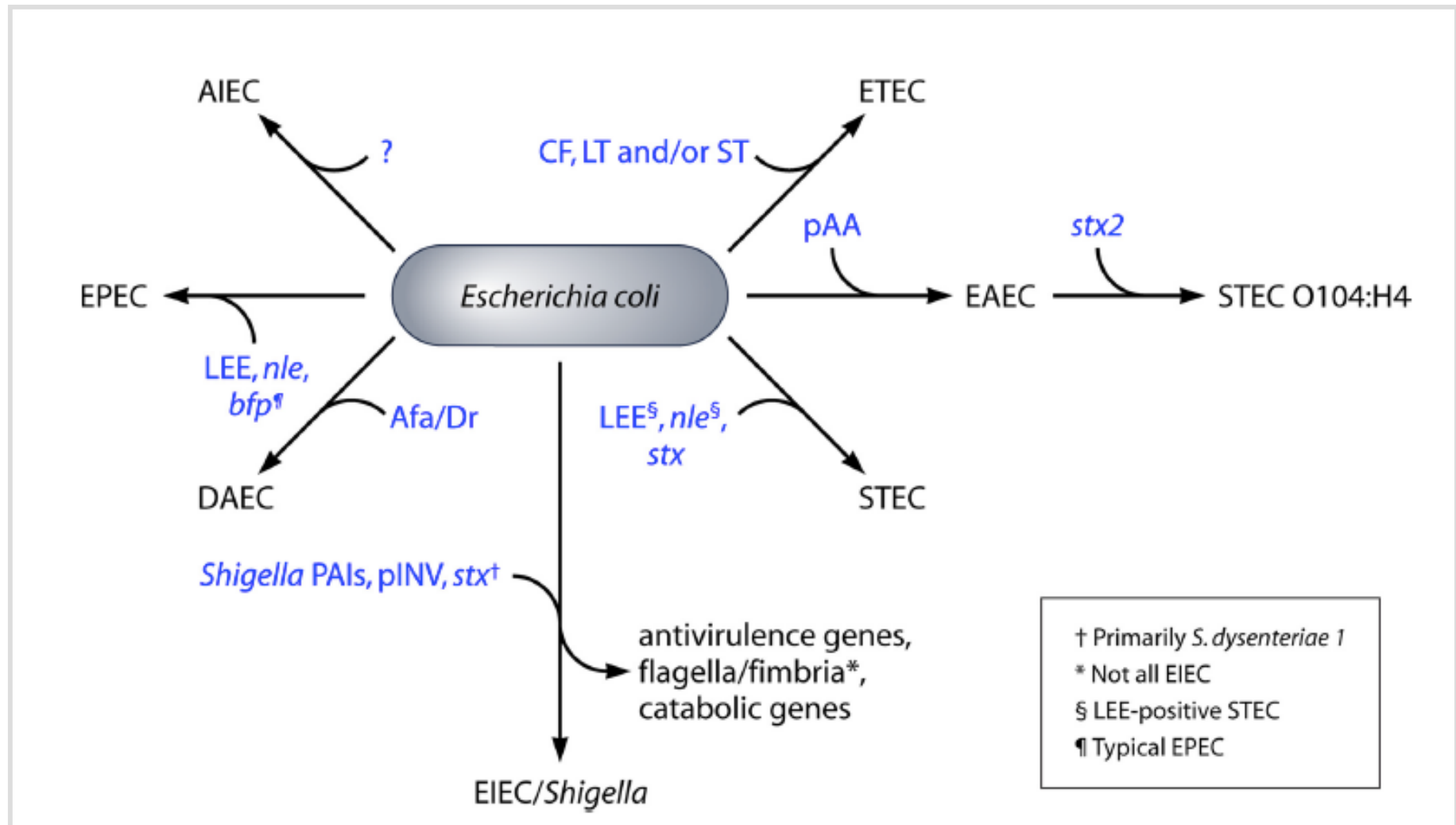
- Joensen, K. G., A. M. Tetzschner, A. Iguchi, F. M. Aarestrup, and F. Scheutz. 2015. Rapid and easy in silico serotyping of *Escherichia coli* using whole genome sequencing (WGS) data. *J.Clin.Microbiol.* 53(8):2410-2426. doi:JCM.00008-15 [pii];10.1128/JCM.00008-15 [doi]

Link to the [article](#)

# VirulenceFinder results

Gene	Associated pathotype	Present in isolate
<i>iroN</i>	UPEC	-
<i>papG</i>	UPEC	-
<i>Stx1a</i>	EHEC	-
<i>Stx1b</i>	EHEC	-
<i>Stx2a</i>	EHEC	+
<i>Stx2b</i>	EHEC	+
<i>pic</i>	EAEC	+
<i>pet</i>	EAEC	-
<i>astA</i>	EAEC	-
<i>aggR</i>	EAEC	+
<i>sta1</i>	ETEC	-
<i>stb</i>	ETEC	-
<i>ltaA</i>	ETEC	-

# Pathogenic gene acquisition and loss for different pathotypes



# The positive control

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### MyDbFinder-1.1 Server - Results

User database				
Fasta header	%Identity	Query/HSP length	Contig	Position in contig
<i>PapG</i>	98.32	1011 / 1011	CFT073_NC_004431	3429593..3430603
<i>PapG</i>	98.52	1011 / 1011	CFT073_NC_004431	4940831..4941841
<i>dsbD</i>	98.17	1698 / 1698	CFT073_NC_004431	4972379..4974076

[extended output](#)[Results as text](#)[Results tab separated](#)[Hit in genome sequences](#)[Database gene sequences](#)

**Selected %ID threshold:** *98.00 %*

**Selected minimum length:** *60 %*

**Input Files:** *EC4.fsa*



escherichia\_coli\_c227-11.fsa

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## MyDbFinder-1.1 Server - Results

User database				
Fasta header	%Identity	Query/HSP length	Contig	Position in contig
<i>dsbD</i>	99.06	1698 / 1698	Supercontig_1.6	34344..36041

**Selected %ID threshold: 98.00 %**

**Selected minimum length: 60 %**

**Input Files: *escherichia\_coli\_c227-11.fsa***