

Supplementary material

Table S1. Primers used for determining the *uidA* and *mutS* genes, and for phylogrouping.

Primer	Sequence (5' – 3')	Size	Reference
<i>uidA</i>	CATTACGGCAAAGTGTGGGTCAAT (F) TCAGCGTAAGGGTAATGCGAGGTA (R)	658 bp*	[1]
<i>mutS</i>	GGCCTATACCCTGAACTACA (F) GCATAAAGGCAATGGTGTC (R)	596 bp	[1]
<i>chuA</i>	ATGGTACCGGACGAACCAAC (F) TGCCGCCAGTACCAAAGACA (R)	288 bp	[2]
<i>yjaA</i>	CAAACGTGAAGTGTGTCAGGAG (F) AATGCGTTCCTCAACCTGTG (R)	211 bp	[2]
<i>TspE4</i>	CACTATTCGTAAGGTCATCC (F) AGTTTATCGCTGCGGGTCGC (R)	152 bp	[2]
<i>ArpA</i>	AACGCTATTCGCCAGCTTGC (F) TCTCCCCATACCGTACGCTA (R)	400 bp	[2]
<i>Group E</i>	GATTCCATCTTGTCAAAATATGCC (F)	301 bp	[2]
<i>ArpAgpE</i>	GAAAAGAAAAAGAATTCCCAAGAG (R)		
<i>Group C</i>	AGTTTTATGCCAGTGCAG (F)	219 bp	[2]
<i>trpAgpC</i>	TCTGCGCCGGTCACGCC (R)		

* bp = base pair.

Table S2. Primers used for amplification of virulence genes.

Primers	Sequence (5' – 3')	Size (bp)	Reference
<i>stx1</i>	Forward ACACTGGATGATCTCAGTGG Reverse CTGAATCCCCCTCCATTATG	614	[3]
<i>stx2</i>	Forward CCATGACAACGGACAGCAGTT Reverse CCTGTCAACTGAGCAGCACTTTG	779	[3]
<i>eaeA</i>	Forward GTGGCGAATACTGGCGAGACT Reverse CCCCATTCCTTTTCACCGTCG	890	[3]
<i>hlyA</i>	Forward ACGATGTGGTTTATTCTGGA Reverse CTTCACGTGACCATACATAT	165	[3]
STa	Forward GCCTATGCATCTACACAATC Reverse TGAGAAATGGACAATGTCCG	278	[4]
LTb	Forward TATCCTCTCTATATGCACAG Reverse CTGTAGTGGAAGCTGTTATA	480	[4]

* bp = base pair.

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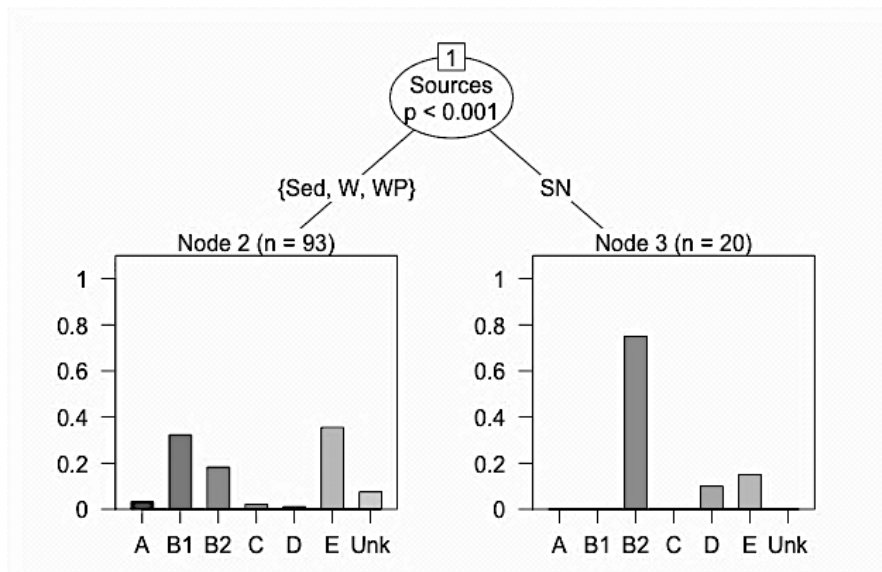


Figure S1. Multinomial log-linear regression analysis of phylogroup distribution of isolates across sample types. Phylogrouping was performed according to the scheme of Clermont *et al.*, 2013. The X axis denotes phylogroups and the Y-axis represents proportion of isolates. Sed – sediment, W – water, WP – water plant, SN – snail.

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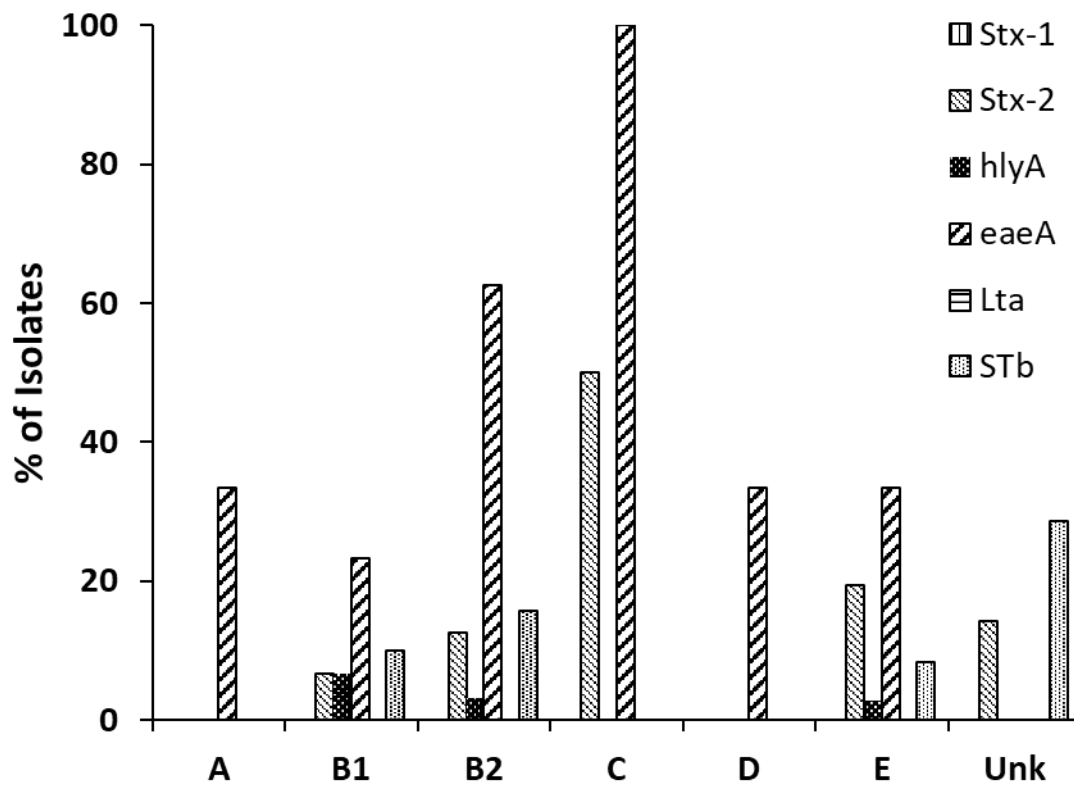


Figure S2. Virulence gene distribution across isolates allocate to phylogroups based on the scheme of Clermont *et al.*, 2013. The number of isolates for each phylogroup is given in parentheses on the x axis.

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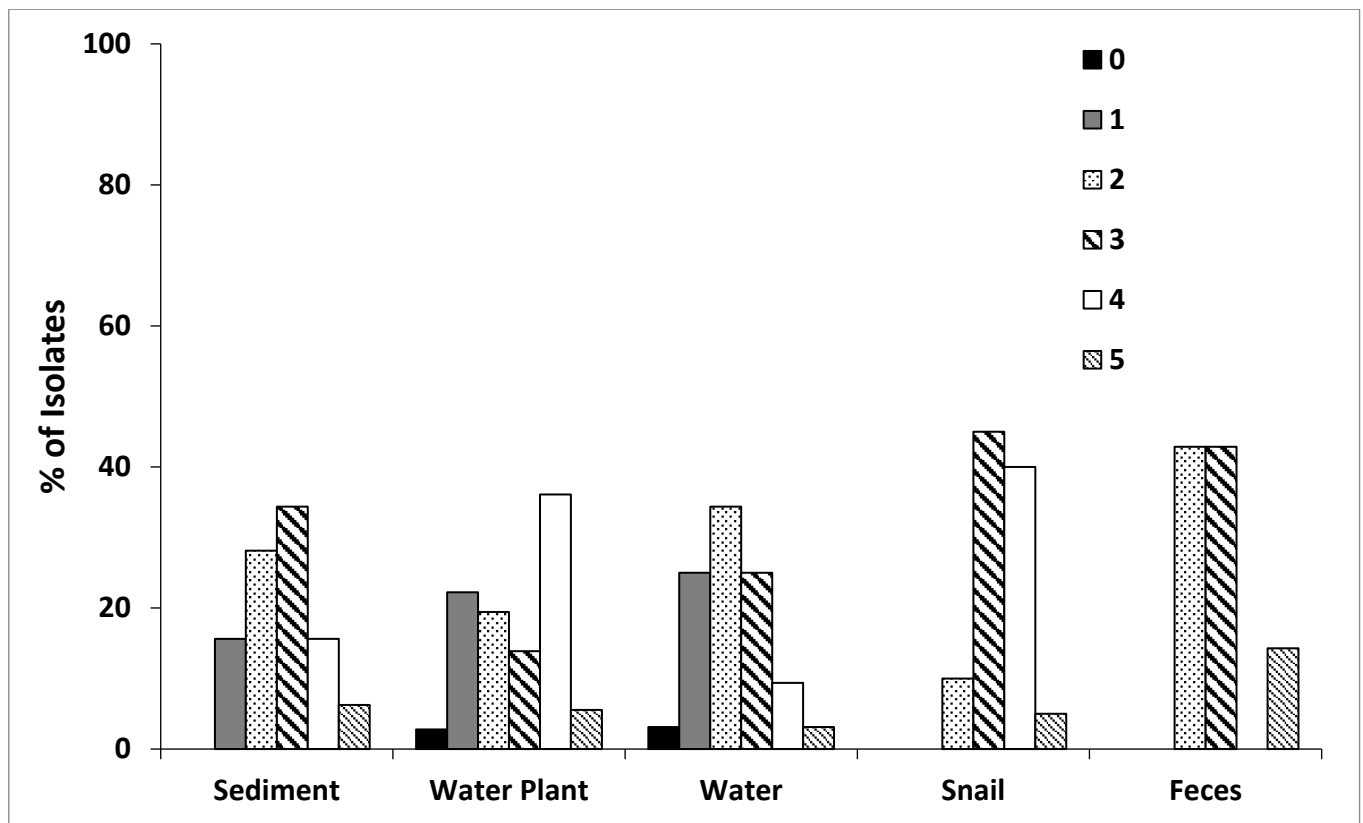


Figure S3. Distribution of sensitive (0) and isolates displaying Intermediate resistance to 1, 2, 3, 4 or 5 antibiotics from the five sampling sites.

1. Walk ST, Alm EW, Gordon DM, Ram JL, Toranzos GA, Tiedje JM, et al. Cryptic lineages of the genus *Escherichia*. *Appl Environ Microbiol.* **2009**; *75*. doi: 10.1128/aem.01262-09.
2. Clermont O, Christenson JK, Denamur E, Gordon DM. The Clermont *Escherichia coli* phylo-typing method revisited: improvement of specificity and detection of new phylo-groups. *Environ Microbiol Rep.* **2013**; *5*(1):58–65. doi: 10.1111/1758-2229.12019.
3. Fagan PK, Hornitzky MA, Bettelheim KA, Djordjevic SP. Detection of Shiga-Like Toxin (stx(1) and stx(2)), Intimin (eaeA), and Enterohemorrhagic *Escherichia coli* (EHEC) Hemolysin (EHEC hlyA) Genes in Animal Feces by Multiplex PCR. *Appl Environ Microbiol.* **1999**; *65*(2):868–872. PubMed PMID: 9925634.
4. Osek J. Multiplex polymerase chain reaction assay for identification of enterotoxigenic *Escherichia coli* strains. *J Vet Diagn Invest.* **2001**; *13*(4):308–311. Epub 2001/08/02. doi: 10.1177/104063870101300405. PubMed PMID: 11478602.

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