

Shiga toxin-producing *Escherichia coli* and their associated serotypes and virulence factors.

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Abstract

Shiga toxin-producing *Escherichia coli* (STEC) is a group responsible for great part of gastrointestinal diseases in humans and for the high calf mortality rate. It is often associated with certain serotypes and other virulence factors. Here we tested STEC strains from feces collected from cohabitants farm animals to determine if they were serving as reservoirs of pathogenicity.

Key words:

Escherichia coli, colibacillosis, serotyping, virulence factors

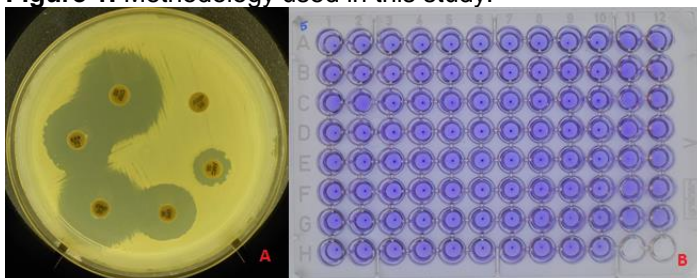
Introduction

Shiga toxin-producing *Escherichia coli* (STEC) is a group responsible for great part of gastrointestinal diseases in humans and for the high calf mortality rate¹. Healthy animals, especially bovines, might have STEC strains and be reservoirs that can later affect humans. STEC is often associated with a great variety of O:H serotypes, the most commonly related to human outbreaks being O157:H7. But when it comes to rough numbers, non-O157 are the most prevalent serogroups². In this study, Shiga toxin-producing *E. coli* strains were tested to determine their serogroups and other virulence factors that might be associated with the pathology brought by the Shiga toxin, such as adherence factors.

Results and Discussion

For this project, 675 *E. coli* strains were isolated from healthy animals' feces from two dairy farms (calf, cow, horse, mule, dog, lamb and sheep) and tested for Shiga toxin genes (*stx1* and *stx2*) using PCR. A total of 108 (16%) strains were positive for *stx1* and/or *stx2*, all of them originated from calf, lamb or sheep. Antimicrobial susceptibility testing for those 108 strains showed 60 phenotypically distinct isolates, which were then tested against 31 different "O" antisera (Fig.1). Out of the 60 strains tested, 12 were positive for O22, 10 for O4, 8 for O5, 4 for O91, 3 for O26, 2 for O103, 2 for O126, 1 for O2, 1 for O109, 1 for O118 and 17 couldn't be serotyped (Table 1). The antimicrobial susceptibility testing (Fig.1) showed that five strains were classified as Multidrug Resistant (MDR, resistant to ≥ 3 antimicrobial classes). All 108 strains were tested for O157 and H7 genes using PCR and none was positive. As for the associated virulence factors, 67 (62%) strains were positive for the adherence factor genes (*eae*).

Figure 1. Methodology used in this study.



A: Disc diffusion method. B: Serotyping.

Table 1. Serogroups found and their respective strains.

Serogroup	<i>E. coli</i> positive
O2	BA36c
O4	BA19c*; BA19d*; BA26b*; BB1a; BB1b; BB1c; BB1d; BB1e; CO1a; CO1e
O5	BA24b; BA32a; BA36b; BA42c; BA44a*; BA56e; BA57b; BA58c*
O22	BA5a; BA5b; BA6a; BA6b; BA9a; BA9b; BA16b; BA18a; BA18d; BA19c*; BA19d*; BA26b*
O26	BA19c*; BA19d*; BA26b*
O91	BA44a*; BA58c*; CO1a; CN1d
O103	BA30e; BA36a
O109	BA42b
O118	BA33c
O126	BA2a; BA3b

*Strains positive for more than one serogroup, needs further investigation. *MDR isolate*: ampicillin, amoxicillin + clavulanic acid, ciprofloxacin, enrofloxacin, chloramphenicol, tetracycline and sulfamethoxazole + trimethoprim.

Conclusions

Our findings indicate that the animals from which the isolates came from were serving as reservoirs for pathogenic *E. coli*, since they weren't sick themselves, but still carried pathogenic strains. This raises concerns about workers involved with the animals and about milk's contamination.

Acknowledgement

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¹ García, A., Ruiz-Santa-Quiteria, J.A., Orden, J.A. *Prod. Animal.* **1999**; 147: 19-372

² Blanco, M., Blanco, J.E., Mora, A., Dahbi, G., Alonso, M.P., González, E.A., Bernárdez, M.I., Blanco, J. *J. Clin. Microbiol.* **2004**; 42(2): 645-651.