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Microbe Profile: *Escherichia coli* O157 : H7 – notorious relative of the microbiologist’s workhorse

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**Graphical abstract**

Summary of enterohaemorrhagic *Escherichia coli* O157 pathogenesis using basic symbols. The right-hand illustration shows enterohaemorrhagic *E. coli* interactions with an epithelial cell.

**Abstract**

*Escherichia coli* O157 : H7 is a zoonotic diarrhoeal pathogen of worldwide importance. It belongs to a subset of Shiga toxin-producing *E. coli* that can form attaching and effacing lesions on intestinal epithelia via the action of a type 3 secretion system that injects bacterial effectors into enterocytes. Infections in humans often arise from contaminated food or direct environmental exposure and can involve life-threatening Shiga toxin-dependent sequelae. In the three decades since *E. coli* O157 : H7 was first recognized intensive research has helped to unravel the basis of pathogenesis, but few effective options for prevention and treatment of infections exist.
TAXONOMY
Domain Bacteria, phylum Proteobacteria, class Gammaproteobacteria, order Enterobacteriales, family Enterobacteriaceae, genus Escherichia, species E. coli, serotype O157:H7.

PROPERTIES
E. coli O157:H7 is a chemoorganotrophic facultative anaerobe with both respiratory and fermentative metabolism identified by biochemical tests and selective/chromogenic media. Cells are Gram-negative straight rods about 1 μm × 2–6 μm and typically motile via peritrichous flagella. The enterohaemorrhagic E. coli (EHEC) pathotype describes strains that can express a type 3 secretion system (T3SS) and produce Shiga toxin subtypes (Stx 1a, 2a, 2c, etc.), although the original definition also required an association with human disease [1]. Different EHEC serotypes are associated with severe human disease but O157:H7 is the most prevalent in North and South America and the UK.

GENOME
EDL933 was the first O157 strain to be sequenced with a 5.4 Mb chromosome encoding 1.34 Mb (1387 genes) in 177 ‘O-islands’ which were not present in the laboratory-adapted E. coli K-12 strain MG1655, although the strains share 4.1 Mb of highly homologous DNA [2]. The majority of E. coli O157:H7 isolates also contain a 92 kb plasmid. Lambdaoid bacteriophages, integrative elements and plasmids have introduced the same or similar genes into diverse serotypes that have distinct evolutionary histories, indicating parallel evolution of EHEC by independent acquisition of mobile elements. Sequencing of E. coli O157:H7 catalysed our understanding of the vast diversity that can exist in the ‘accessory genome’, with just 39.2% of non-redundant proteins shared by EDL933, MG1655 and uropathogenic E. coli CFT703. The most recent evidence, based on analysis of over 1000 E. coli O157:H7 genomes, is that the more recent acquisition of Stx2a prophages was critical to emergence of EHEC O157:H7 as a life-threatening zoonotic pathogen in the last few decades [3]. Machine-learning approaches applied to whole-genome sequences indicate that only a minority of E. coli O157:H7 isolates may have zoonotic potential with prophage variation being critical to this capacity [4].

PHYLOGENY
Evidence supports evolution of E. coli O157:H7 from a non-toxigenic EPEC O55:H7 by acquisition of Stx2c-encoding prophage followed by the pO157 plasmid and the switch to the O157 somatic antigen via recombination. There was divergence from this β-glucuronidase-positive last common ancestor approximately 400 years ago with the common ancestor of the current diversity appearing an estimated 175 years ago. There are considered to be three main lineages of EHEC O157:H7, viz. I, I/II and II. The common ancestor was lineage II, β-glucuronidase negative and sorbitol non-fermenting. Lineage I split from this approximately 125 years ago and lineage I/II approximately 95 years ago [3, 5].

KEY FEATURES AND DISCOVERIES
In 1983, Karmali et al. noted that Stx-producing E. coli may be a common cause of haemorrhagic colitis preceding haemolytic uremic syndrome [6]. Stx is produced and released following induction of the bacterial SOS response which induces activation of the integrated Stx-encoding prophage(s). As specific classes of antibiotics can induce this response, there is debate over their use to treat EHEC infections. Stxs share their catalytic mechanism with ricin with an A subunit that arrests translation by depurination of 28S rRNA as well as five B subunits involved in cell binding and trafficking. Globotriaosylceramide (Gb3/CD77) toxin receptors are present on the surface of endothelial cells throughout the human body, particularly capillaries in the kidneys and brain. For intestinal colonization, E. coli O157:H7 forms ‘attaching and effacing’ lesions on intestinal epithelium in a manner dependent on the T3SS encoded by the locus of enterocyte effacement. Attaching and effacing lesions are characterized by intimate bacterial adherence to the apical surface of enterocytes on actin-rich ‘pedestals’ and destruction of proximal microvilli (Graphical abstract figure). Remarkably, intimate adherence is facilitated by injection of Tir, which serves as a receptor for the bacterial outer membrane protein intimin [7]. Prototype E. coli O157:H7 strains inject around 40 ‘effector’ proteins, expressed from the locus of enterocyte effacement and assorted prophage regions, into enterocytes to subvert cellular responses [8].

Ruminants are a key reservoir of E. coli O157:H7. Colonization of cattle is generally asymptomatic, as Gb3 is not expressed on the surface of bovine endothelial cells. The bacteria primarily colonize follicle-associated epithelium of the terminal rectum and super-shedding animals are responsible for the majority of on-farm transmission and the threat of zoonosis [9]. The primary role of Stxs in reservoir hosts or the environment remains to be determined but may include killing of grazing protozoa, immune modulation and enhancement of bacterial adherence. Selection of E. coli O157:H7 in the bovine host may be towards innate and adaptive immune silencing mechanisms that promote persistence and transmission, whereas these mechanisms may have different outcomes in humans as an incidental host leading to life-threatening pathology.

Control measures have focused on the ruminant reservoir, including vaccines based on secreted proteins or an extract of iron-limited E. coli O157:H7, but to date these have not received sufficient uptake. Treatment in humans remains largely supportive, with Stx-specific neutralizing antibody and receptor mimics showing promise for control of systemic sequelae in animal models. A serious outbreak in 2011 caused by an Stx2-producing E. coli O104:H4 hybrid with features of EHEC and enterohaemorrhagic E. coli serves to highlight the dynamic evolution of Stx-producing E. coli
and compels us to look beyond *E. coli* O157:H7 for future threats.

**KEY QUESTIONS**

- What is the purpose of Stx in the environment and/or ruminant host? Ultimately, the selective pressure maintaining Stx also drives the evolution of EHEC strains and so this knowledge is critical.

- What factors influence the zoonotic potential of EHEC strains and disease severity? Despite the common presence of Stx-encoding *E. coli* in ruminants and the environment, why do only a small subset of strains actually cause serious human disease?

- What processes lead to the genetic convergence of multiple lambdoid prophages encoding both effector proteins and Stx in strains with a T3SS, and how likely are similar convergences with pathogenic potential?

- How do you generate an effective immune response against an organism that manipulates host responses to remain ‘under the radar’?

- What level of reduction of shedding in cattle is required to reduce incidence of human disease? If only a small subset of strains cause human disease, would widely applied cattle-based interventions ever represent a cost-effective method of controlling human infections?

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The authors declare that there are no conflicts of interest.

**References**


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