Case Report

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First reported case of *Campylobacter lanienae* enteritis in a human

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Introduction: Campylobacters are the most frequently identified bacteria causing diarrhoea in humans worldwide. *Campylobacter lanienae* was isolated for the first time in 2000 from faecal samples of two asymptomatic abattoir workers in Switzerland during a routine hygiene screen, but has never been associated with human disease.

Case presentation: At hospital admission, the patient reported diarrhoea, lower abdominal cramps, nausea, one episode of bilious vomiting and low-grade fever of 38 °C. The patient was having 10 or more diarrheic stools per day as well as during the night, and had noticed blood mixed with the stools on several occasions. Stool cultures were negative for species of *Salmonella* and *Shigella*, *Escherichia coli* O157:H7 and *Yersinia enterocolitica*, but were positive for *C. lanienae*. Identification was made by classical biochemical testing, as well as 16S rRNA gene and *cpn60* sequencing. The patient slowly improved without antibiotic treatment and was discharged nine days after admission with complete resolution of symptoms.

Conclusion: On the whole it seems very likely that *C. lanienae* was the causative agent. Clinical microbiologists should be aware of this micro-organism which can be identified by phenotypic and molecular methods. The real burden of *C. lanienae* infection in humans might be underestimated and should be further investigated as a potential cause of human diarrhoea disease.

Received 7 March 2016 Accepted 29 April 2016

Introduction

Campylobacters are the most frequently identified bacteria causing diarrhoea in humans worldwide (Allos, 2001). *Camplylobacter lanienae* was isolated for the first time in 2000 from faecal samples of two asymptomatic abattoir workers in Switzerland during a routine hygiene screen (Logan *et al.*, 2000), but has never been reported in humans since. *C. lanienae* has also been isolated from pig and wild boars faeces in Japan (Sasaki *et al.*, 2003, 2013), Hungary (Schweitzer *et al.*, 2011), Korea (Shin & Lee, 2009) and the USA (Jay-Russell *et al.*, 2012), from wild ruminants in Spain (Carbonero *et al.*, 2014; Navarro-Gonzalez *et al.*, 2014), from sheep in Turkey (Acik *et al.*, 2013) and Spain

The GenBank/EMBL/DDBJ accession number for the 16S rRNA gene sequence of strain 001A-0718 is HM770742.

(Oporto & Hurtado, 2011), and from bovine faeces in Alberta (Inglis *et al.*, 2003, 2004, 2005) as well as in Quebec (Guevremont *et al.*, 2008). Recently, the bacterium was also isolated from laboratory chinchillas in the USA (Turowski *et al.*, 2014). In this study we report what seems to be, to our knowledge, the first case of a symptomatic human infection with *C. lanienae*.

Case report

In March 2009, a 39-year-old female sought treatment at the emergency department of the Centre Hospitalier Universitaire de Sherbrooke, reporting a 5-day history of diarrhoea, lower abdominal cramps, nausea, one episode of bilious vomiting and low-grade fever of 38 °C. She was having 10 or more diarrheic stools per day as well as during the night, and had noticed blood mixed with the stools on

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Keywords: Campylobacter; enteritis; campylobacteriosis.

several occasions. She denied any infectious contact, recent antibiotic treatment or travel outside the province of Quebec. She was living on a hog farm and usually drank water from a private well. One year prior to the consultation, the patient had a diagnosis of irritable bowel syndrome and a colonoscopy performed then was normal. For the past few months, she was experiencing three loose stools a day. The patient had also been diagnosed with gastroesophageal reflux disease, hypothyroidism and refractory interstitial cystitis, and had a total hysterectomy. Her daily medication included esomeprazole, levothyroxine, phenazopyridine, amitriptyline, pentosan polysulfate sodium and pregabalin. Physical examination showed sinus tachycardia with normal blood pressure and a temperature of 37.5 °C. The abdomen was diffusely tender but soft with normal bowel sounds. The complete blood count, serum electrolytes,

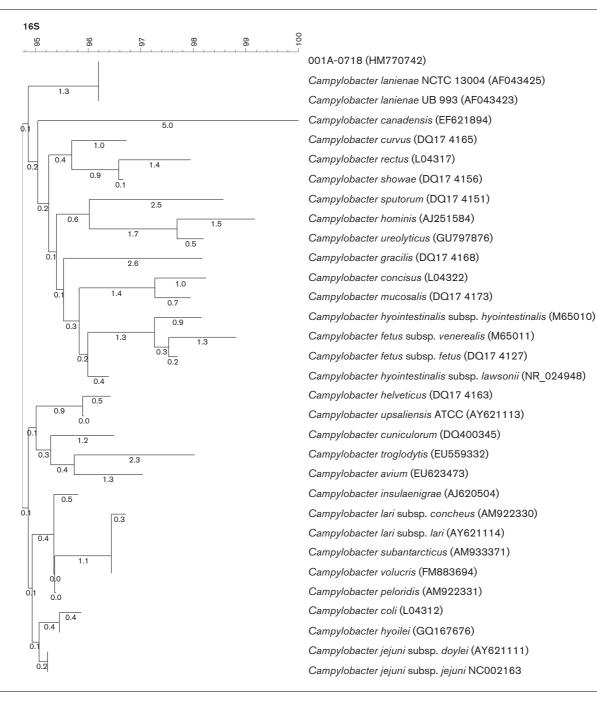


Fig. 1. Phylogenetic tree based on the neighbour-joining method representing the genetic relatedness between partial 16S rRNA gene sequences for campylobacters. The bar represents the percentage of similarity, and numbers on the tree indicate internal branch distances within the resulting trees obtained by bootstrap analysis (1000 replicates). GenBank accession numbers are in parentheses.

creatinine, and liver and pancreatic enzymes were normal. A pelvic and abdominal ultrasound showed only a left ovarian cyst. The patient was admitted to our hospital for intravenous hydration and observation. Because of the persistence of symptoms five days after admission, a colonoscopy was performed and revealed mild inflammation in the left colon. Biopsies confirmed discrete acute focal colitis. The *Clostridium difficile* toxin assay and a stool examination for ova and parasites were negative. Blood cultures were also negative. Stool cultures done two days apart were negative for species of *Salmonella* and *Shigella*, *Escherichia coli* O157:H7 and *Yersinia enterocolitica*, but showed a '*Campylobacter*-like' organism. Because the patient's symptoms were not compatible with a viral infection, no viral detection tests were performed. The patient slowly improved without antibiotic treatment and was discharged nine days after admission with complete resolution of her symptoms.

16S		GenBank		
26	98 66 00 100	accession no.	Country	Source
		AY288304	Canada	Bovine
		EU007896	Canada	Bovine
		EU007902	Canada	Bovine
		EU181124	Canada	Bovine
		EU181125	Canada	Bovine
		EU181126	Canada	Bovine
		EU181127	Canada	Bovine
		EU181128	Canada	Bovine
		EU181130	Canada	Bovine
		EU181131	Canada	Bovine
		EU181132	Canada	Bovine
		EU181135	Canada	Bovine
		EU181136	Canada	Bovine
		EU181137	Canada	Bovine
		EU007903	Canada	Bovine
		EU007904	Canada	Bovine
		EU007901	Canada	Bovine
		EU181123	Canada	Bovine
		EU007900	Canada	Bovine
		HM770742 (001A-0718)	Canada	Human
		AB076675	Japan	Swine
		AB076676	Japan	Swine
		AB076677	Japan	Swine
		AF550664	Austria	Swine
		AY165045	Ireland	Swine
		AF043425	Switzerland	Human
		AF043423	Switzerland	Human
		EU007897	Canada	Swine
		EU007898	Canada	Swine
		EU181129	Canada	Swine
		EU181133	Canada	Swine
		EU181134	Canada	Swine
		EU007899	Canada	Swine
		EU007905	Canada	Swine
		NC002163 (C. jejuni)		

Fig. 2. Phylogenetic tree based on the unweighted pair group method with arithmetic mean (UPGMA) method representing the genetic relatedness between sequences of the partial 16S rRNA gene for *C. lanienae*. The bar represents the percentage of similarity. *C. jejuni* was used as an outgroup.

Investigations

The stool sample was initially streaked on Karmali agar and incubated under microaerobic conditions generated in a jar with a CampyGen sachet (Oxoid) at 42 °C for 48 h. Small, grey, flat, irregular or round and moist colonies were observed but growth was poor compared to the *Campylobacter jejuni* control strain. The Gram stain revealed slender, curved Gram-negative rods. The organism was oxidase- and catalase-positive, hippurate-negative, indoxyl acetate-negative, resistant to both nalidixic acid and cephalothin, and susceptible both to erythromycin and ciprofloxacin by the disc diffusion method.

The Laboratoire de Santé Publique du Québec (LSPQ) further identified the isolate by sequencing the cpn60 gene (Hill et al., 2006) and the 16S rRNA gene (Bekal et al., 2006). BLAST analysis of the partial 16S rRNA gene sequence obtained (1221 bp), using the taxonomy browser of the National Center for Biotechnology Information (www.ncbi.nlm.nih.gov), revealed between 98 % and 100 % sequence identity with available C. lanienae sequences. The sequence was deposited in the GenBank database under accession number HM770742. Fig. 1 shows the phylogenetic position of the patient's isolate (001A-0718) in a neighbour-joining tree representing the genetic relatedness between DNA sequences of the partial 16S rRNA gene of Campylobacter lineage. The isolate represented a distinct monophyletic lineage within the genus Campylobacter and was related to human strains NCTC 13004 and UB 993, the only two strains of C. lanienae previously isolated from humans (Logan et al., 2000). Sequence comparison of our isolate with NCTC 13004 showed 99.43% similarity for 1220 nucleotides. The partial sequence of the cpn60 gene was also compared with cpnBD (Hill et al., 2004) and the isolate was identified as C. lanienae.

Fig. 2 shows the phylogenetic position of our isolate in an unweighted pair group method with arithmetic mean (UPGMA) tree representing the genetic relatedness between DNA sequences of the partial 16S rRNA gene from *C. lanienae* isolated worldwide from human, bovine and swine faeces. Our isolate clustered with *C. lanienae* previously isolated from swine in Japan, Austria, Ireland, Switzerland and Canada, and the sequence analysed showed the same five nucleotide differences in regard to bovine isolates, as previously described by Guevremont *et al.* (2008). The fact that our isolate was resistant to cephalothin also supported the relationship with swine isolates, as bovine strains are typically susceptible to cephalothin (Shin & Lee, 2009).

Discussion

In a PCR-based survey of *Campylobacter* associated with faeces collected from beef cattle, *Campylobacter* DNA was detected in 83 % of the samples (Inglis *et al.*, 2003). The most frequently detected species was *C. lanienae* (49 %), followed by *C. jejuni* (38 %), *Cmpylobacter hyointestinalis* (8 %) and *Campylobacter coli* (0.5 %). In another study,

some cattle were found to be shedding exceptionally large numbers of cells (approximately 10^8 cells g⁻¹), and numbers of *C. lanienae* released in the environment in cattle faeces could exceed 10^{12} cells per animal per day(Inglis *et al.*, 2004). Inglis *et al.* reported that *C. lanienae* dwells in the large intestine, including the caecum, proximal ascending colon, distal descending colon, and rectum of cattle (Inglis *et al.*, 2005) and suggested that *C. lanienae* may be pathogenic to cattle (Inglis *et al.*, 2004).

Inglis et al. also observed that PCR-based detection methods were substantially more effective for detecting C. lanienae from faecal specimens than conventional culture methods (Inglis et al., 2003). They isolated C. lanienae from cattle faeces, but found that cattle strains of the bacterium were very fastidious and did not grow on many of the commercial media commonly used to isolate campylobacters (Inglis & Kalischuk, 2003). The first isolation of C. lanienae was obtained on Campylosel agar, consisting of 5 % (v/v) blood in a Columbia agar base with the selective antibiotics cefoperazone, vancomycin and amphothericin B (Logan et al., 2000). When comparing different culture media for growing C. lanienae from cattle faeces, Inglis and Kalischuk were only able to recover isolates from Karmali medium at 40 °C (Inglis & Kalischuk, 2003). C. lanienae is susceptible to polymyxin B (Logan et al., 2000), which might explain why the organism cannot be recovered from Preston and Campy-Line media. Also, bovine strains of C. lanienae in Canada typically cannot be isolated on CCDA, the most common medium used in hospitals to isolate campylobacter from human stool samples (Inglis et al., 2004). As our hospital laboratory uses Karmali agar as the routine media for the detection of Campylobacter in human faeces, we were able to detect the presence of C. lanienae in our patient.

In 2009, nearly 20% of the 2025 Campylobacter isolates notified in Quebec were not identified to the species level and were reported as Campylobacter sp. (Bureau de surveillance et vigie, 2010), possibly due to the inability (or the lack of interest or resources) of the clinical laboratory to perform phenotypic identification tests for species of Campylobacter other than C. jejuni. On the whole it seems very likely that C. lanienae was the causative agent. Furthermore, given that C. lanienae would not typically be detected in human diagnostic facilities, despite its prevalence in cattle and pig faeces, the real burden of C. lanienae in humans might be underestimated and should be further investigated as a potential cause of human diarrhoea disease.

Acknowledgements

We thank the personnel of the clinical microbiology laboratory of the Centre Hospitalier Universitaire de Sherbrooke for the strain isolation and biochemical analysis.

This study was conducted with funding from the Centre de recherche du Centre hospitalier universitaire de Sherbrooke (CHUS), Sherbrooke, Québec, Canada. On behalf of all authors, the corresponding author states that there is no conflict of interest.

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