## Probable Campylobacter fetus subsp. fetus Gastroenteritis

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Three strains of Campylobacter fetus subsp. fetus isolated from cases of gastroenteritis are reported. DNA-DNA hybridizations in addition to biochemical tests were used to confirm the identification of the isolates as C. fetus since all strains grew at 42°C. These isolates, like other C. fetus strains, are susceptible to cephalothin and thus would not have been detected in laboratories with Campylobacter isolation media containing this component.

It is generally agreed that Campylobacter je*juni* is a leading cause of enteritis in humans, whereas Campylobacter fetus subsp. fetus rarely, if ever, causes enteritis and is primarily an opportunistic pathogen capable of causing systemic illness in debilitated or immunosuppressed patients. We report here on three strains of C. fetus recently isolated from fecal specimens of individuals with gastroenteritis. All patients were young adult males with no history of underlying disease and whose predominant complaint was crampy abdominal pain with intermittent bloody stools. A culture of feces from all three individuals on the routine enterobacterial media vielded no pathogens. In addition, tests for ova and parasites were negative.

The three strains were isolated in three different laboratories in diverse geographical locations of southern California, and the media and methods were not identical. The one consistent feature was that all laboratories used a Campylobacter isolation medium which did not contain cephalothin. Two laboratories used Campylobacter agar (Anaerobe Systems, Santa Clara, Calif.) and the third used a Campylobacter agar manufactured by Callabs (California Laboratories Industries, North Hollywood, Calif.). Both media employed combinations of vancomycin, trimethoprim, and polymyxin as the basis for selectivity. Two of the laboratories routinely incubated their primary Campylobacter plates at 42°C in a GasPak jar with a Campy Pak envelope (BBL Microbiology Systems, Cockeysville, Md.). The third laboratory incubated Campylobacter plates at 35°C in a GasPak jar with the GasPak envelope but without a catalyst. All three laboratories isolate C. jejuni from diarrheal feces at an incidence comparable or exceeding that of Salmonella spp.; none had previously reported the isolation of a C. fetuslike organism from feces.

Identification of the strains as *C. fetus* was by biochemical tests and DNA-DNA hybridizations. Methods were those described by Harvey and Greenwood (3), except for the oxidase and hippurate tests which were performed with commercially available diagnostics (Clinical Standards Laboratory, Carson, Calif.). All three strains were tested on the battery of biochemicals (Table 1), but only the first isolate was confirmed by DNA-DNA hybridizations (Table 2).

All strains conform to the expected profile for C. fetus subsp. fetus with the exception of growth at 42°C. Although C. fetus subsp. fetus strains which are capable of growth at 42°C have been noted before (2, 4), it is considered a rare, but not inconsistent, feature. The DNA-DNA hybridizations confirmed the validity of this profile identification for isolate number 1. As Table 2 illustrates, the DNA from isolate number 1 was 98.8% related to that from the type strain of C. fetus subsp. fetus and less than 6.8% related to the DNA from the representative strains of the other Campylobacter species and Campylobacter-like organisms.

In 1979, Butzler and Skirrow (1) reported that in a survey of 22,000 stools for *Campylobacter* species, *C. fetus* was found in only three of the samples, and there was no association with gastroenteritis. Because of this extensive work in Belgium, most laboratories discount the possibility of isolating *C. fetus* from fecal specimens. Therefore, many of the commercially available *Campylobacter* isolation media contain cephalothin, an antibiotic which inhibits the growth of *C. fetus* strains but allows the growth of *C. jejuni* strains. In addition, most labora-

Strain	Oxidase	Catalase	Hippurate hydrolysis	Diam of inhibition zone (mm)		Growth at:			Growth	Growth
				Nalidixic acid	Cephalothin	25°C	35°C	42°C	in 1% glycine <sup>a</sup>	in 3.5% NaCl <sup>a</sup>
C-135	+	+	-	_b	18	+	+	+	+	
C-140	+	+	_	_	19	+	+	+	+(-)	-
C-142	+	+	-	-	20	+	+	+	+	-

TABLE 1. Results of phenotypic tests

<sup>a</sup> Tests were terminated after 5 days of incubation at 35°C. For each test, two basal media were used: fluid thioglycollate (Difco Laboratories, Detroit, Mich.), and *Brucella* broth (Difco) containing 0.16% agar. If test results differed, the test result with *Brucella* broth basal is given in parentheses.

b -, No zone of inhibition.

tories incubate Campylobacter selective media at 42°C, a temperature which enhances the growth of C. jejuni but which inhibits the growth of most strains of C. fetus. Unless laboratories

TABLE 2. DNA relatedness values between							
Campylobacter reference strains and thermophilic							
Campylobacter C-135							

Reference DNA from strain:	% Relatedness to strain C-135				
C-16, C. fetus subsp. fetus ATCC					
27374	98.8				
C-61, C. jejuni ATCC 29428	5.7				
C-38, C. coli Butzler P2	4.5				
C-32, "C. fecalis" Firehammer					
14227A	4.5				
C-133, NARTC <sup>a</sup> NCTC 11352	4.3				

<sup>a</sup> NARTC, Nalidixic acid-resistant thermophilic Campylobacter spp.

modify their Campylobacter isolation techniques to encompass the growth requirements of C. fetus as well as those of C. jejuni, it will remain impossible to adequately assess the relative frequency of C. fetus-associated gastroenteritis in this country. As with Yersinia enterocolitica, statistics for Belgium are not necessarily valid for the United States.

## LITERATURE CITED

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