ORIGINAL ARTICLE

Increased stool sampling during a waterborne outbreak of cryptosporidiosis does not increase the detection of other faecal pathogens

A O Qamruddin, M G L Keaney, R McCann, P R Chadwick

J Clin Pathol 2002;**55**:271–274

See end of article for authors' affiliations

Correspondence to:
Dr A O Qamruddin,
Department of
Microbiology, Manchester
Royal Infirmary, Oxford
Road, Manchester
M13 9WL, UK;

Accepted for publication
29 October 2001

Aims: To ascertain the effect of enhanced surveillance following an outbreak of waterborne cryptosporidiosis on the number of faecal specimens submitted to the local microbiology laboratory and the number positive for common enteric pathogens. The outbreak provided an opportunity to estimate the extent of routine under ascertainment of common enteric pathogens.

Method: Retrospective search of the computerised microbiology system database for details of faecal examination requests for the period 26 April to 6 June in 1998 and 1999 (period of outbreak).

Results: Specimens were received from 378 community patients during the six week period 26 April to 6 June 1999. This was double that for the same period in 1998 (a non-outbreak year). Oocysts of *Cryptosporidium parvum* were detected in 59 patients, an eightfold increase compared with the same period in 1998. Despite the greater number of patients tested, the detection of other pathogens in patients with gastroenteritis was not altered when compared with the same period in the previous year.

Conclusion: This study found no evidence of under ascertainment of gastrointestinal infection (common bacterial pathogens and rotavirus) by local general practitioners.

Cryptosporidiosis¹ is caused by the protozoan *Cryptosporidium parvum* of which there are human and animal associated genotypes. Infection in the immunocompetent individual is characteristically a self limiting diarrhoea, which may take up to a few weeks to resolve. However, in immunocompromised individuals, infection may not be cleared and can result in severe intractable diarrhoea, with associated morbidity and mortality. Transmission occurs by animal to human (zoonotic infection), person to person, and waterborne routes.²

An outbreak of cryptosporidiosis occurred in the north west of England in 1999.³ This followed the detection on the 23 April of 30 cryptosporidium oocysts (normal, < 2) in a routine weekly 10 litre grab sample taken on the 20 April from the raw inlet water for the Dunmail Raise water treatment works. This plant supplies water to many areas in the north west of England, including Salford (which receives almost 100% Thirlmere aqueduct water). The population of Salford is approximately 240 000.

"In immunocompromised individuals, infection may not be cleared and can result in severe intractable diarrhoea, with associated morbidity and mortality"

Following an outbreak meeting on 26 April it was decided not to issue a general "boil water" notice because any contaminated water would have already passed through the water distribution system and daily water samples for cryptosporidium oocysts from the 23 April onwards had been within normal limits. A press statement was released and a letter was sent to all general practitioners in Salford, requesting them to send faecal specimens from patients with suspected infective diarrhoea.

A sharp increase in laboratory confirmed cases of cryptosporidiosis was noted in subsequent weeks and affected mainly four health authority areas: Salford and Trafford, Wigan and Bolton, South Lancashire, and Northwest Lancashire. The outbreak subsided by the first week of June 1999.

Cryptosporidium parvum from patients associated with this outbreak has been typed as genotype 2 (the animal associated genotype) by the Public Health Laboratory Service. *Cryptosporidium parvum* was also detected in sheep around the Thirlmere reservoir.

We analysed microbiology laboratory results and demographic data on community patients from whom faeces specimens had been processed during the period 26 April to 6 June 1999 by the microbiology laboratory, based at Hope Hospital, which serves the population of Salford. Comparisons were made with results and data from the same period in 1998, during which no outbreaks of infectious diarrhoeal disease were reported. The aim was to ascertain what effect a combination of the cryptosporidiosis outbreak, enhanced general practitioner surveillance, and increased public awareness of diarrhoeal disease would have on the number of community patients providing faecal specimens for processing by the microbiology laboratory and on the number of patients testing positive for common enteric pathogens.

METHODS

Routine testing of all community faecal specimens in Salford involves selective culture for *Salmonella* spp (DCA agar and XLD agar; LabM, Bury, UK), *Shigella* spp (DCA agar), *Campylobacter* spp (Campylobacter selective agar; Oxoid, Basingstoke, UK), and *Escherichia coli* 0157 (cefexime tellurite sorbitol MaConkey agar; Oxoid), auramine staining for cryptosporidium oocysts, and a direct wet preparation examination for ova, cysts, and parasites. Rotavirus antigen detection by latex agglutination (Rotagen; Biokit, Barcelona, Spain) is performed on samples from patients under 5 years of age. There were no changes in laboratory protocols during the two study periods.

The microbiology computer system database (TelePath CDS; TelePath Ltd, Birmingham, UK) was retrospectively searched

Abbreviations: CCDC, consultant in communicable disease control; CI, confidence interval

Figure 1 Illness onset dates for cryptosporidium positive individuals between 26 April and 6 June 1999.

Figure 2 Number of community faecal specimens received between 5 April and 13 June, 1998 and in 1999 (weekly intervals).

Table 1 Number of community patients positive for enteric pathogens from 1998 to 1999 for the period 26 April to 6 June

	1998		1999		Percentage change	95% CI
	N	%	N	%		
Faeces negative patients	143		290			
Faeces positive patients	32	18	88	23	+5	-2 to 12
<i>Cryptosporidium</i> sp	7	4	59	16	+12	7.3 to 16.7
Rotavirus	10	6	10	3	-3	-6.8 to 0.6
<i>Campylobacter</i> sp	12	7	15	4	-3	-7.2 to 1.4
<i>Salmonella</i> sp	2	1	3	1	0	-2.1 to 1.5
<i>Shigella</i> sp	1	1	0	0	-1	
<i>Giardia</i> sp	0	0	1	0	0	
All patients	175		378			

CI, confidence interval.

for details of faecal examination requests from Salford residents for the period 26 April to 6 June in 1998 and in 1999. Data were analysed on a patient basis rather than on a specimen basis (that is, multiple samples from the same patients, of which there were 21 and three during the period 26 April to 6 June in 1999 and in 1998, respectively, were treated as a single request). Statistical analysis included calculation of the 95% confidence intervals for the percentage difference in positive isolates between the two time periods. Illness onset dates for cryptosporidium positive individuals were obtained from the local consultant in communicable disease control (CCDC; RM).

Some general practitioners serving Salford patients use microbiology laboratory services other than the Hope Hospital microbiology laboratory. Of all Salford residents notified to the CCDC with laboratory confirmed cryptosporidiosis between 26 April and 6 June 1999, 77% had been tested at Hope Hospital. Our data therefore refer to most, but not all, of the Salford population.

RESULTS

Figure 1 shows illness onset dates for cryptosporidium positive individuals between 26 April and 6 June 1999. Two peaks can be seen, one around the 24 April and the other

Table 2 Microbiology results, with age breakdown, on general practice patients from whom faeces specimens were received during the period 26 April to 6 June, 1998 and 1999

Age (in years)	Population of Salford as estimated at 1/1/99	Patients positive for cryptosporidium 26–6/6		Patients positive for other tested pathogens 26/4–6/6		Patients negative for all tested faecal pathogens 26/4–6/6	
		1998	1999	1998	1999	1998	1999
0–4	14224	1	17	12	14	35	64
5–9	16285	2	10	0	1	6	14
10–14	15541	0	8	0	2	3	9
15–19	16008	2	5	0	1	3	8
20–24	17601	0	6	1	2	8	16
25–29	19190	0	7	1	1	8	23
30–34	19680	0	3	2	1	4	23
35–39	17929	0	0	0	0	4	23
40–44	14883	0	1	1	1	9	12
45–49	13754	0	0	1	1	11	9
50–54	14426	0	1	2	1	12	10
55–59	11998	0	0	2	1	5	3
60–64	11238	1	0	1	1	8	9
65–69	10130	0	1	1	0	3	12
70–74	9162	0	0	1	0	5	12
74+	17225	1	0	0	0	13	25
Not known	–	0	0	0	2	6	18
Totals	239274	7	59	25	29	143	290

around the 7 May. Onset dates were not available for six patients. Figure 2 shows the number of general practice patients from whom faeces specimens were received during seven day intervals spanning the period 5 April to 13 June, both for 1998 and 1999. The number of samples increased during the period 26 April to 16 May in 1999 compared with the same period in 1998. Patients with cryptosporidiosis submitted faeces for testing between 0 and 23 days (median, six) after the onset of symptoms.

Table 1 shows the percentage change in the number of individuals positive for enteric pathogens between 1999 and 1998 for the period 26 April to 6 June. The total number of patients tested more than doubled and the number of patients positive for cryptosporidium increased more than eightfold. There was no appreciable change in the number of patients with *Campylobacter* spp, salmonella, shigella, giardia, or rotavirus. Because only one isolate of shigella and one isolate of giardia were obtained over the two time periods, statistical analysis was not undertaken. Calculation of 95% confidence intervals demonstrates that only the increase in cryptosporidium was significant. Table 2 shows the analysis, with age stratification, of microbiology results on community patients from whom faeces specimens were received during the period 26 April to 6 June in 1998 and in 1999. The observed increase in cryptosporidium positive patients in the 1999 period was weighted towards the younger ages, particularly those under 10 years, whereas the increase in patients testing negative for the routinely tested enteric pathogens was more evenly distributed.

DISCUSSION

This outbreak of cryptosporidiosis was confirmed by both the absolute and proportionate (relative to the total number of patients tested for enteric pathogens) rises in the numbers of cryptosporidium positive patients, compared with the same period the previous year. Despite a twofold increase in the total number of patients tested during the outbreak period, there was no proportionate increase in the numbers of patients positive for other routinely tested enteric pathogens. The rise in the number of patients tested during the outbreak period is more than can be accounted for by all the patients testing positive for the routinely tested pathogens. These excess patients may represent patients with cryptosporidiosis (or other recognised pathogens) testing laboratory negative,

Take home messages

- Double the number of specimens were received in 1999 compared with the same period in 1998 (a non-outbreak year) and there was an eightfold increase in the number of patients positive for *Cryptosporidium parvum*
- However, the detection of other pathogens in patients with gastroenteritis was not altered when compared with the same period in the previous year
- Therefore, there was no evidence of under ascertainment of gastrointestinal infection (common bacterial pathogens and rotavirus) by local general practitioners

patients with non-infective causes of diarrhoea, or patients with viral gastroenteritis (for example, Norwalk-like virus, adenovirus, calicivirus). However, following the enhanced surveillance request and press publicity, the rise was probably the result of a lower threshold for individuals with diarrhoea (regardless of cause) seeking medical advice and a lower threshold for requesting faeces examination by general practitioners.

“The rise in the number of patients tested during the outbreak period is more than can be accounted for by all the patients testing positive for the routinely tested pathogens”

General practitioners were not given explicit advice for obtaining a faecal specimen but were asked to sample patients with suspected infective diarrhoea. It is accepted that there is wide variation in faecal sampling between general practitioners. It was not possible to determine the number of patients presenting with symptoms of gastroenteritis to all general practitioners during the two periods. This would have required a separate study and manual retrieval of data in many practices. One possible explanation for the absence of an increase in the isolation of other pathogens is that local general practitioners are correctly sampling patients who have bacterial pathogens. However, this hypothesis is not borne out by other studies, although they did demonstrate that patients with illness caused by common bacterial pathogens, such as *Salmonella* sp and *Campylobacter* sp, are more likely to see their general practitioners, to have faeces examined, and to be

reported to local and national surveillance networks than individuals with viral gastroenteritis.^{4,5} In one of these studies the ratio of cases in the community to cases reported to national surveillance was calculated.⁴ This ratio was lower for bacteria (salmonella, 3.2 : 1) than for viruses (rotavirus, 35 : 1).

However, our study is not directly comparable with these because our objective was to determine whether enhanced surveillance (as a result of an outbreak) picked up additional cases of gastrointestinal infection and not to determine the background incidence of gastrointestinal infection in the community. We acknowledge, however, that our study has limitations because of the small sample size.

A coincidental community wide outbreak of diarrhoea caused by pathogens other than those routinely tested (such as Norwalk-like viruses) cannot be ruled out, but seems unlikely.

ACKNOWLEDGEMENT

The authors gratefully acknowledge the help of R Philip and C Bryan in gathering data for this paper.

.....

Authors' affiliations

A O Qamruddin, M G L Keaney, P R Chadwick, Department of Medical Microbiology, Salford Royal Hospitals NHS Trust, Hope Hospital, Salford M6 8HD, UK

R McCann, Public Health Department, Salford and Trafford Health Authority, Salford M30 0NJ, UK

REFERENCES

- 1 **Ungar BLP**. Cryptosporidium. In: Mandell GL, Bennet JE, Dolin R, eds. *Principles and practice of infectious diseases*. London: Churchill Livingstone, 1995:2500–10.
- 2 **Meinhardt PL**, Casemore DP, Miller KB. Epidemiologic aspects of human cryptosporidiosis and the role of waterborne transmission. *Epidemiol Rev* 1996;**18**:118–36.
- 3 **Communicable Disease Surveillance Centre**. Outbreak of cryptosporidiosis in north west England. *Commun Dis Rep CDR Weekly* 1999;**9**:175–8.
- 4 **Wheeler JG**, Cowden JM, Sethi D, et al. Study of infectious intestinal disease in England: rates in the community, presenting to GPs and reported to national surveillance. *BMJ* 1999;**318**:1046–50.
- 5 **Tompkins DS**, Hudson MJ, Smith HR, et al. A study of infectious intestinal disease in England: microbiological findings in cases and controls. *Commun Dis Public Health* 1999 Jun;**2**:108–13.

New JCP online submission and review system

We are pleased to inform authors and reviewers of the new online submission and review system at JCP. Developed by High-Wire Press (CA, USA), Bench Press is a fully integrated electronic system that utilises the web to allow rapid and efficient submission of manuscripts. It also allows the peer review process to be conducted entirely online. We are one of the first journals in the BMJ Special Journals group to go online in this way. The aim, apart from saving trees, is to speed up the often frustratingly slow process (for both authors and editors) from submission to publication. Many reviewers might appreciate this too. Authors may submit their manuscript in any standard word processing software. Acceptable standard graphic formats include: jpeg, tiff, gif, and eps. The text and graphic files are automatically converted to PDF for ease of distribution and reviewing purposes. Authors are asked to approve their submission before it formally enters the reviewing process. On approval by the authors, the submission is passed to the editor and/or reviewers via the web. All transactions are secure.

To access the system click on "SUBMIT YOUR MANUSCRIPT HERE" on the JCP homepage: HYPERLINK <http://www.jclinpath.com>, or you can access Bench Press directly at HYPERLINK <http://submit-jcp.bmjournals.com>.

We are very excited with this new development and would encourage authors and reviewers to use the online system whenever possible. As editors, we will use it all the time, the up side being lack of need to travel to the editorial office to deal with papers, the down side having no more excuses to postpone decisions on papers because we are "at a meeting"!

The system is very easy to use and should be a big improvement on the current peer review process. Full instructions can be found on Bench Press <http://submit-jcp.bmjournals.com> and JCP online at <http://www.jclinpath.com>. Please contact Natalie Davies, Project Manager, HYPERLINK <mailto:ndavies@bmjgroup.com> for any further information.

H Holzel, P van Diest