EDITORIAL BOARD

EDITOR IN CHIEF Professor G I Muguti

ASSOCIATE EDITORS

Professor IT Gangaidzo Dr S P Munjanja

EDITORIAL BOARD MEMBERS

Professor MM Chidzonga (Zimbabwe) Professor P Jacobs (South Africa) Dr R A Kambarami (Zimbabwe) Professor S A Latif (Zimbabwe) Professor P R Mason (Zimbabwe) Professor CT Musabayane (Zimbabwe) Professor KJ Nathoo (Zimbabwe) Mr L Nystrom (Sweden) Dr S Siziya (Zambia)

PAST EDITORS

Professor M Gelfand (1953-1985) Professor H M Chinyanga (1985-1990) Professor J A-Matenga (1991-1999)

ADMINISTRATIVE AND OFFICE STAFF

Director of Publications: Mr. Munani S Mtetwa Administrative Manager: Mr Christopher Mashavira

Technical Editor: Mrs Ling M Cooper Statistical Advisor: Mr S Rusakaniko Secretary: Mrs Patricia Bhunu

All manuscript will be prepared with the International Committee of Medical Journal Editors-Uniform requirements for manuscript submitted to Biomedical Journals 1993.

Manuscript submitted for publication are accepted on the understanding that they are contributed exclusively to the *Central Journal of Medicine*. A statement to that effect should be included in the letter accompanying the manuscript.

Communications concerning editorial matter, advertising, subsriptions, change of address, etc. should be addressed to the Administrative Manager, P.O. Box A195 Avondale, Harare, Zimbabwe.

The subsription rate including surface transmission postage is:-

	SURFACE TRANSMISSION			AIR-MAIL TRANSMISSION		
	INDIVIDUAL	INSTITUTE	POSTAGE	INDIVIDUAL	INSTITUTE	POSTAGE
ZIMBABWE	Z\$40 000	Z\$50 000	Z\$13 200		I	
AFRICA	US\$281	US\$379	US\$90	US\$281	US\$379	US\$200
REST OF THE WORLD	US\$351	US\$379	US\$90	US\$351	US\$379	US\$278

Owned and published by the Central African Journal of Medicine in conjuction with the Faculty of Medicine



University of Zimbabwe

Cryptosporidiosis in Harare, Zimbabwe

C SIMANGO, S MUTIKANI

Abstract

Objective: To determine the prevalence of Cryptosporidium parvum in diarrhoeal patients.

Design: This was a laboratory-based cross sectional study on cryptosporidiosis in diarrhoeal patients. **Setting:** Department of Medical Laboratory Sciences, College of Health Sciences in Harare, Zimbabwe.

Subjects: People of all ages with diarrhoea presenting at primary level health centres in Harare.

Main Outcome Measures: Patient's age, laboratory results.

Results: Cryptosporidium parvum was the commonest enteric pathogen and was detected in 5.8% of the 500 diarrhoeal patients of all ages followed by Shigella species (3.8%) and Salmonella species (2.0%). The highest detection rate of C.parvum oocysts was observed in children less than five years old (11.2%) followed by children between six and 10 years old (6.3%) and then the 31 to 40 year age group (5.9%).

Conclusions: Cryptosporidiosis affects people of all ages in Harare but is more common in children, particularly those under five years. The *C.parvum* oocysts should be looked for routinely in diarrhoeal stool specimens particularly those from children less than five years since *C.parvum* may be one of the causative agents of diarrhoea in this age group.

Cent Afr J Med 2004;50(5/6):52-4

Introduction

Coccidia of the genus Cryptosporidia are small protozoan parasites found in many animals such as chickens, turkeys, mice, guinea pigs, calves, sheep and rhesus monkeys.1 Many human infections are zoonotic but person to person spread through the faecal-oral route in cryptosporidiosis is common especially in crowded environments. Epidemiological studies have indicated a higher prevalence of cryptosporidiosis in developing countries than in developed countries.² A study carried out during the rainy season at Baragwanath Hospital in South Africa showed that 17 of the 92 (18.4%) children less than two years old with diarrhoea and one of the 29 (3,4%) children without diarrhoea excreted C. parvum oocysts in their faeces.³ In a study conducted over a period of one year in South Africa, Moodley et al.⁴ showed that Cryptosporidium parvum infections were detected in 9.0% of children under 10 years who presented with diarrhoea, and were significantly more prevalent during the rainy season. A study on cryptosporidiosis in Zambian children, carried out during the rainy season, showed a high prevalence rate of 18% of 222 children with diarrhoea.⁵ The detection rates of C.parvum in diarrhoeal patients in other African countries were 7.8% in Rwanda,6 6.1% in Sudan7 and 3.8% in Kenya.8

C.parvum is recognised as an important enteropathogen in children and in immunocompromised persons. Cryptosporidiosis has been strongly linked to Human Immunodeficiency Virus (HIV) infection, especially in the advanced stages. In immunocompetent people, C. parvum infection causes acute self-limiting diarrhoea9 but chronic cryptosporidiosis is common in immunosuppressed patients such as those with HIV/AIDS¹⁰ as well as those receiving chemotherapy. In a study carried out in Lusaka, Zambia, cryptosporidiosis was found more frequently in HIV-seropositive (14%) than in HIV-seronegative (8%) children. 11 C. parvum has also been shown to be an important aetiological agent of acute diarrhoea in HIV seropositive patients in Zaire. 10,12 The aim of this study was to determine the prevalence of C.parvum oocysts in patients with diarrhoea presenting at health centres in Harare.

Materials and Methods

The present study was conducted over a period of four months (November to February) during the hot and wet season. Cryptosporidiosis has been shown to occur more frequently during the rainy season.⁴ Five hundred stool specimens were collected in sterile containers from people of all ages and sexes with all forms of diarrhoea, presenting as outpatients at primary level health centres in Harare. No information was available on the HIV status of the patients

Department of Medical Laboratory Sciences College of Health Sciences University of Zimbabwe P O Box A178 Avondale, Harare, Zimbabwe Correspondence and reprint requests to: Dr C Simango

and the duration of the diarrhoeal illness. Names, ages and sex of all patients were checked in order to avoid multiple specimens from the same patients. All the samples were processed on the same day of collection from the patients.

For the detection of C. parvum, the stool specimens were concentrated for the C. parvum oocysts using the formolether concentration technique.¹³ The sediments of the concentrated stool specimens were used to prepare smears on standard glass microscope slides. The slides were allowed to air dry before being fixed in absolute methanol for three minutes. The faecal smears were stained for C. parvum oocysts using the modified Ziehl-Neelsen technique.13 The smears were stained with cold carbol fuchsin for 15 minutes and then rinsed in distilled water. The smears were decolourised with 1% acid alcohol for 10 to 15 seconds, then rinsed with distilled water. The smears were then counterstained with methylene blue for 30 seconds, rinsed with distilled water and allowed to air dry. The stained slides were examined microscopically for C. parvum oocysts at 1000X magnification. A stool specimen was considered positive for C. parvum oocysts if the stained smear had brightly red or pink stained round or oval oocysts.13

Other parasitic pathogens were sought using the formolether concentration technique. ¹³ All the specimens processed during this study were also cultured for *Shigella* species, *Salmonella* species, *Campylobacter* species and enteropathogenic *Escherichia coli* using standard procedures. ¹⁴ Viruses were not looked for in the diarrhoeal stools.

Results

The distribution of enteric pathogens detected in the stool samples is shown in Table I. *C.parvum* oocysts were detected in 5.8% of the patients and it was the most common enteric parasitic pathogen isolated followed by *G.lamblia*, and then *Entamoeba histolytica*. *Shigella* species were the most common bacterial enteric pathogens isolated followed by *Salmonella* species, then *Campylobacter* species and enteropathogenic *Escherichia coli*.

Table I: Enteric pathogens detected in the diarrhoeal stools.

Pathogen	Number of isolat (n = 500)	es Percentage
Shigella species	19	3.8
Salmonella species	10	2.0
Campylobacter species	6	1.2
Enteropathogenic Escherichia coli	5	1.0
Cryptosporidium parvum	29	5.8
Giardia lamblia	3	0.6
Entamoeba histolytica	1	0.2
No pathogen	429	85.8

The distribution of C. parvum infection by age group is shown in Table II. The highest percentage of C. parvum

infection was observed in patients of the zero to five year age group (11.2%), followed by the six to 10 year age group (6.3%). The lowest percentage was observed in the 11 to 20 year age group (1.4%). The occurrence of coinfection by *C. parvum* and pathogenic bacteria was observed in two patients. In one patient, *C. parvum* was detected together with *Campylobacter* species and in another patient the parasite was detected together with *Salmonella* species.

Table II: Age distribution of patients with diarrhoea due to C. parvum infection.

Age group in years	Number of specimens processed	Number of specimens positive for <i>C.parvum</i>	%
0 - 5	134	15	11.2
6 - 10	32	2	6.3
11-20	71	1	1.4
21-30	85	3	. 3.5
31-40	85	5	5.9
> 40	93	3	3.2

Discussion

Epidemiological studies have demonstrated that cryptosporidiosis is more prevalent in developing countries than in developed countries where prevalence is low.² In the present study, it was observed that 5,8% of the people of all ages with diarrhoea excreted *C. parvum* oocysts in their faeces. The present study was carried out during the rainy season, when transmission of cryptosporidiosis was likely to be high.⁴ The prevalence of *C. parvum* associated diarrhoea which was observed in the present study is higher than those which were observed in previous surveys for enteropathogens in Harare urban area.^{15,16}

This may be due to the fact that the present study was carried out during the hot and wet months which has been associated with a high prevalence of *C.parvum* infection whereas the other studies included the dry months.

The 5,8% prevalence of C. parvum infection noted in the present study is almost similar to other cryptosporidiosis prevalence rates of some of the developing African countries such as Rwanda (7,8%), Sudan (6,1%), and South Africa (4,1%).² In the present study the highest prevalence of cryptosporidiosis (11.3%) was in the zero to five year age group and similar observations were made by Moodley et al.4 in South Africa where prevalence of C. parvum in children was 9,0%. A high detection rate of C. parvum oocysts was observed in four crowded townships of Lusaka, Zambia, where the prevalence of cryptosporidiosis in 222 children with diarrhoea was 18%.5 The observed high prevalence of C.parvum infection in children aged less than 10 years strongly implicates C. parvum infection as an important cause of diarrhoea in children, especially those aged less than five years. However, asymptomatic infection with C. parvum occurs in some people, as is the case with

many enteric parasites. Viruses such as rotaviruses, which have been shown to be important causative agents of diarrhoea in young children, 17 were not looked for in the present study.

The HIV status of the patients in the present study was not known. The 31 to 40 year age group also had a high prevalence of *C. parvum* infection (5,9%). Although the HIV status of the patients was not known in the present study, the high prevalence of cryptosporidiosis in this age group could be due to HIV related immunosuppression since these persons are at high risk of acquiring HIV infection. Sub-Saharan Africa has the highest HIV infection prevalence in the whole world, and AIDS related cryptosporidiosis could as well be very high in Zimbabwe. Zimbabwe is a country in southern Africa with a very high prevalence of HIV infected people.

The present study has shown that cyptosporidiosis affects outpatients of all ages in Harare but the infection occurs mainly in children less than 10 years old, with the peak prevalence in children aged under five years. *C. parvum* is a parasite which should be looked for routinely in diarrhoeal stool specimens in clinical laboratories particularly those specimens from children under five years old.

References

- 1. Nime FA, Burek JD, Page DL, Holscher MA, Yardley JH. Acute enterocolitis in a human being infected with the protozoan *Cryptosporidium*. *Gastroenterology* 1976;70:592-8.
- 2. Current WL. Garcia LS. Cryptosporidiosis. Clin Microbiol Rev 1991;4:325-58.
- 3. Berkowitz FE, Vallabh W, Buqwana A, Heney C. Cryptosporidiosis in black South African children. S Afr Med J 1988;74:272-3.
- 4. Moodley D, Jackson TFHG, Gathiram V, Van den Ende J. *Cryptosporidium* infection in children in Durban: seasonal variation, age distribution, and disease status. *S Afr Med J* 1991;79:295-7.
- 5. Nchito M, Kelly P, Sianougo S, Juo PN, Feldman R, Farthing M. Sri-Baboo K. Cryptosporidiosis in urban Zambian children. An analysis of risk factors. *Am J Trop Med Hyg* 1998;59:435-7.
- Bogaerts J, Lepage P, Rouvroy D, Vandepitte J. Cryptosporidium spp. a frequent cause of diarrhoea in Central Africa. J Clin Microbiol 1984;20: 874-6.
- 7. Robinson M, Hart CA, Baxby D, Battin M, Sulliman GI, el Seed AM, Coulter JB. *Cryptosporidium* as a cause of gastroenteritis in Sudanese children. *Ann Trop Paediatr* 1986;6:155-6.
- 8. Simwa JM, Chunge RN, Kinoti SN, Karumba PN, Wanola I, Kabiru P. Cryptosporidiosis and childhood diarrhoea in a rural community in Kenya. *East Afr Med J* 1989; 66:520-5.
- 9. Hunt DA, Shannon R, Palmer SR, Jephcott, AE. Cryprosporidiosis in an urban community. *BMJ* 1984;289:814-16.

- 10. Colebunders R, Lusakumuni K, Nelson AM, Gigase P, Lebughe I, van Marck E, et al. Persistent diarrhoea in Zairian AIDS patients: an endoscopic and histological study. Gut 1988;29:1687-91.
- 11. Chintu C, Luo C, Baboo S, Khumalo-Ngwenya B, Matthewson J, DuPont HL, *et al*. Intestinal parasites in HIV-seropositive Zambian children with diarrhoea. *J Trop Pediatr* 1995;41:149-52.
- 12. Kelly P, Nchito M, Sri-Baboo K, Ndubani P, Nchito M, Okeowo NP, et al. Cryptosporidiosis in adults in Lusaka, Zambia and its relationship to oocyst contamination of drinking water. J Infect Dis 1997;176:1120-3.
- 13. Cheesbrough M. Medical Laboratory Manual for Tropical Countries. Volume I: Microbiology. Cambridge:ELBS/Butterworth, 1987.
- 14. Cheesbrough M. Medical Laboratory Manual for Tropical Countries. Volume II: Microbiology. Cambridge:ELBS/Butterworth, 1985.
- 15. Simango C, Nyahanana M. Campylobacter infection in children in an urban community. Cent Afr J Med 1997;43:172-5.
- 16. Simango C, Mbewe C. Salmonella enteritidis diarrhoea in Harare, Zimbabwe. Trop Med Intern Health 2000;5:503-6.
- 17. Cruickshank JG, Zilberg B. Winter diarrhoea and rotaviruses in Rhodesia. *S Afr Med J* 1976;50:1895-6.



This work is licensed under a Creative Commons
Attribution – NonCommercial - NoDerivs 3.0 License.

To view a copy of the license please see: http://creativecommons.org/licenses/by-nc-nd/3.0/

