

# ROTAVIRAL AND PROTOZOAL INFECTIONS AMONG MALIGNANT CHILDREN IN BASRAH GOVERNORATE

Maysloon A . Al-Sadoon \*

## SUMMARY

Over five months period , the incidence of rotaviral and protozoal infections in patients receiving treatment for malignancy was investigated in the present study. A total of 58 patients [38 males and 20 females] were involved in this study. Formalin – ether concentration method And direct smear method were used for diagnosis of stool samples. Higher rates of infection were Found among those <2 years old ,and also among males than Females . In this study higher rate of infection in rural than in urban area was found and majority of them with mixed parasitic infections . All them with symptomatic and most patient which positive for Rotavirus with chronic diarrhea (66.7 % ) higher than discontinuous diarrhea (33.3 % ).

## INTRODUCTION

Rotavirus is the most common cause of severe gastroenteritis in infants and young children worldwide(1). In developing countries, rotavirus gastroenteritis is a major cause of childhood death and it is responsible for approximately half a million deaths per year among children aged <5 years (2) Rotaviruses are shed in high concentrations in the stools of infected children and are transmitted by the fecal-oral route, both through close person-to-person contact and through fomites (3-4). Rotaviruses also might be transmitted by other modes, such as water and respiratory droplets (5-6) The spectrum of rotavirus illness ranges from mild, watery diarrhea of limited duration to severe, dehydrating diarrhea with vomiting and fever, which results in death(7-).8 Rotaviral gastroenteritis may result in mortality for populations at risk such as infants, the elderly, and immunocompromised patients . Previous

study indicates that may produce a chronic infection in immunodeficient Children( 9 ) the chronic infection is accompanied the generation of extra viral immunodeficient host (10) unusual assortments in an genomic segments and their children and adults with impaired immunity, such as those with congenital immunodeficiency, or post haematopoietic or solid organ transplantation, are at increased risk of severe, prolonged, and even fatal rotavirus gastroenteritis.(<sup>11,12</sup>) Rotavirus is an important cause of nosocomial gastroenteritis,<sup>(13,14)</sup> and can also cause disease in adults, especially those caring for children, and outbreaks of gastroenteritis in aged care facilities.<sup>(12)</sup> Patients with some type of immunocompromised condition and those submitted to immunosuppressive therapy have an increased probability of acquiring parasitic infections, generally with a high degree of severity (15,16 )

---

\* Department of Microbiology , College of Medicine , University of Basrah, Basrah - IRAQ

The intestinal non-opportunistic pathogenic parasites most frequently encountered in immunocompromised hosts include *Entamoeba histolytica/ Giardia lamblia* and others. Parasitic infections that cause auto-limited diarrhea in immunocompetent patients may cause profuse diarrhea in immunocompromised individuals(17,18). The aim of the present study is detect of Rotaviral and protozoal infections in malignant children in Basrah

## MATERIAL & METHODS

This study was conducted in Basrah Teaching Hospital (IRAQ), over a period of five months from March to August 2010.(58) malignancy affected patients (38) males and (20) females were studied . These subjects who were presumably immunocompromised patients received cytotoxic drugs . Their age varied from 1 to >13 years . The control group was composed of (35) apparently healthy individuals (25 males and 10 females ) their age range from 1 to 13 years none had any past or present history suggesting malignant disease nor had they received any type of anticancer therapy . who were randomly selected . Fecal samples were examined microscopically ( wet mounts) by the direct method using normal saline and lugol,s iodine . Formalin – ether concentration method was used according to procedure described in diagnostic microbiology and diagnostic medical parasitology (19-20) . The principle of one step Rotavirus test Device - (FECES) is the mixture migrates upward on the test line region of the test . During testing the specimen reacts with the particle coated with anti - rotavirus antibody .The mixture migrates up word on the

membrane chromate graphically by capillary action to react with anti-rotavirus . antibody on the membrane and generate a colored line .The presence of this colored line in the test region indicates a positive result . While the absence indicates a negative result .

## RESULTS & DISCUSSION

The types of malignant diseases among tested patients are present in (table 1).The results showed that acute lymphoblastic leukemia (ALL) was the major type (55.1%) of malignant cases in the studied subjects .The other types ranged from (1.7% to 8.6 %). These result similar to the result of various studies.(21-22).Age of most patient with ALL are within age group of (3-11) years and the male higher number than female .

The explanation of this result is possibly due to the changing age structure of such cases with a shift towards the younger age group (23), or may be due to viral infection especially at this age group which lead to changes in immune system. Sex of my patients ,males were more affected than females by malignant. Diseases including leukemia ,similar result was obtained by two studies done in united states ( 22 ,24)

The clinical symptoms among tested patients was illustrated in table (2) showed that (75.8%) of patients suffered from weight loss , while fever and diarrhea were noticed in 28.3% and 24% respectively . while other study done in southern Iran Showed 74% of patients with fever .(25)

TABLE (3) show the distribution of rotaviral infection among patient & control group . It is clear that patient had the

highest rate of rotaviral infection & the male more affected. In this study the prevalence of rotavirus was 6(10.34)%. Alfalahi2002(26) documented a prevalence rate 24% while Aahmed et al associated 2006(27) Erbil recovered prevalence rate are 37% other study mentioned by Ali et al 2008 (28), the percentage was 19% and higher than the result of study reported by Vesikari et al, 1981 (29), the percentages was 11%/. The Distribution of cases according to sex show high prevalence rate in male than female. May be attributed to (x) linked recessive susceptibility pabset et al & Branth et al, 2003 (30-31).

in Table-4- Parasitic infections among patients with malignant disease and control group according to sex out of (58) patients with malignant diseases (24) 41.37% with parasitic infection (18) 31.03% male and (6) 10.34% female. It is clear that, patient had the highest rate of parasitic infections than control group. These result may be related to the fact; that males are more active, mobile and integrated in the

environment especially among agricultural community.

Higher percent of infected patient with rotavirus was found in rural area (5) 13.9% than in urban area (1) 4.5%. That agree with the result was observed by AL-Ani in Ramadi city (32). These differences might be due to poor water supply, poor sewage disposal and low education.

Table -6 The distribution of the parasitic infections is presented in this table. Giardia lamblia was the most frequent infection among children (15.5%), followed by Entamoeba histolytica (10.3%) and Blastocystis hominis (8.6%). The most prevalent parasites in our study were G. lamblia and E. histolytica. A similar finding has been reported by Azab. et al (33).

This table 7 show higher percent in patients with chronic diarrhea ( ) than patients with acute diarrhea (%). The same result was observed in some study (9,33).

## TABLES

Table -1-Number of cases in relation of type of malignant diseases with age and sex:

	Type of Malignant cases	Age		Sex		Total	
		< 2	3 – 11	Male	Female	No.	%
Leukemia	1- ALL	6	26	20	12	32	55.1%
	2- AML	1	2	1	2	3	5.2%
	Sarcoma	-	1	1	-	1	1.7%
	Neuroblastoma	2	3	3	2	5	8.6%
	Hepatoblastoma	1	-	-	1	1	1.7%
	Wilm's Tumor	-	1	1	-	1	1.7%
	Lymphoblastic Lymphoma	-	4	2	2	4	6.8%
	H.L	-	3	3	-	3	6.1%
	N.H.L	-	4	3	1	4	6.8%
	Retinoblastoma	-	1	1	-	1	1.7%
	Rhabdomyosar-coma		1	1		1	1.7%
	Ewing's Sarcoma	-	1	1	-	1	1.7%
	Histocytosis	-	1	1	-	1	1.7%
	Total	10	48	38	20	58	100%

ALL = Acute lymphocytic leukaemia; AML = Acute myelocytic leukaemia; HL = Hodgkin lymphoma; NHL = Non-Hodgkin lymphoma.

Table -2- :Clinical Symptoms among tested patients

Symptoms	No.	%
Weight loss	44	75.8%
Fever	15	28.3%
Diarrhea	14	24.1%
Abdominal Pain	10	18.8%
Vomiting	6	11.3%
No. Symptoms	0	0%

TABLE – 3-: Rotavirus infections among patients with malignant disease and control group:

		INFECTION		+VE	-VE	TOTAL
<b>PATIENTS</b>	<b>MALE</b>	<b>NO</b>		<b>4</b>	<b>34</b>	<b>38</b>
		<b>%</b>		<b>10.53</b>	<b>89.47</b>	<b>100.00</b>
	<b>FEMALE</b>	<b>NO</b>		<b>2</b>	<b>18</b>	<b>20</b>
		<b>%</b>		<b>10%</b>	<b>90%</b>	<b>100.00</b>
		<b>TOTAL</b>		<b>6</b>	<b>52</b>	<b>58</b>
<b>CONTROL</b>	<b>MALE</b>	<b>NO</b>		<b>0</b>	<b>15</b>	<b>15</b>
		<b>%</b>		<b>0.00</b>	<b>100.00</b>	<b>100.00</b>
	<b>FEMALE</b>	<b>NO</b>		<b>1</b>	<b>19</b>	<b>20</b>
		<b>%</b>		<b>5%</b>	<b>95%</b>	<b>100.00</b>
		<b>TOTAL</b>		<b>1</b>	<b>34</b>	<b>35</b>

Table-4-: Parasitic infections among patients with malignant disease and control group according to sex

	No.examined	Parasitic infection			
		male	%	female	%
Patient group	58	18	31.03%	6	10.34%
Control group	35	2	5.71%	2	5.71%

Table -5-:Rotavirus infection in relation to residence:

Infection			+ ve	- ve	Total
Patient group	Urban	No.	1	21	22
		%	4.5%	95.5%	100.00
	Rural	No.	5	31	36
		%	13.9%	86.1%	100.00
Total			6	52	58
Control group	Urban	No.	0	22	22
		%	0.00	100.00	100.00
	Rural	No.	1	12	13
		%	7.7	92.3	100.00
Total			1	34	35

Table -6-:The Parasitic infections among patients and control groups:

Parasites	Patients		Control	
	No.	%	No.	%
G.lamblia	9	15.5%	2	5.7%
B.hominis	5	8.6%	2	5.7%
E.histolytica	6	10.3%	-	
G + B	4	6.9%	-	
Total	24	41.3%	4	11.4%

Table -7-:Type of diarrhea in patient with rotavirus infection:

Patient	No.	%
Continues diarrhea (chronic diarrhea)	4	66.7 %
Discontinues diarrhea (acute diarrhea)	2	33.3%
Total	6	100.00

## REFERENCES

- 1- Chakravarti A . ,Chauhan MS., Sharma A ,Verma A. Distribution of human rotavirus G and P genotypes in a hospital from northern India .Southeast Asian J . Trop. Med. Public Health 2010 sep.,41 (5).1145-52 .
- 2- Parashar UD , Hummelman EG , Bresee JS , Miller MA , Glass RI . Global illness and deaths caused by rotavirus disease in children . Emerg Infect Dis . 2003 ;9:565 -72 .
- 3- Butz AM, Fosarelli P, Dick J, Cusack T, Yolken R. Prevalence of rotavirus on high-risk fomites in day-care facilities. *Pediatrics* 1993;92:202-5.
- 4- Leen –Jan van Doorn ,Bernhard Kleter ,Evert Hoefnagel , Isabelle Stainier , Annick Poliszczak , Brigitte Colan,and Wim Quint . Detection and genotyping of human Rotavirus VP4 and VP7 genes by reverse transcriptase PCR and reverse hybridization . J. Clin microbial . 2009 sep . 47 (9) : 2704 -2712 .
- 5- Centers for Disease Control and Prevention (CDC). Prevention of rotavirus gastroenteritis among infants and children. Recommendations of the Advisory Committee on Immunization Practices (ACIP). *MMWR - Morbidity & Mortality Weekly Report* 2006;55(RR-12):1-13.
- 6- Dennehy PH, Nelson SM, Crowley BA, Saracen CL. Detection of rotavirus RNA in hospital air samples by polymerase chain reaction (PCR). [1998 abstract The American Pediatric Society and The Society for Pediatric Research]. *Pediatric Research* 1998;43(Suppl 2):143A.
- 7- Rodriguez WJ, Kim HW, Brandt CD, et al. Longitudinal study of rotavirus infection and gastroenteritis in families served by a pediatric medical practice: clinical and epidemiologic observations. *Pediatr Infect Dis J* 1987;6:170-6.
- 8- Glass RI, Kilgore PE, Holman RC, et al. The epidemiology of rotavirus diarrhea in the United States: surveillance and estimates of disease burden. *J Infect Dis* 1996;174(suppl 1):S5-11.

- 9- Saulsbury FT, Winkelstein JA, Yolken RH. Chronic rotavirus infection immunodeficiency. *J Pediatr.* 1980 Jul;97(1):61-5.
- 10- Oishi I; Kimura T; Murakami T; Haruki K; Yamazaki K; Seto Y; Minekawa Y; Funamoto H; Serial observations of chronic rotavirus infection in an immunodeficient child. *Microbiol Immunol.* 1991;35(11):953-61.
- 11- Yolken RH, Bishop CA, Townsend TR, et al. Infectious gastroenteritis in bone-marrow-transplant recipients. *New England Journal of Medicine* 1982;306:1009-12.
- 12- Kapikian AZ, Hoshino Y, Chanock RM. Rotaviruses. In: Knipe DM, Howley PM, Griffin DE, et al, eds. *Fields Virology*. 4th ed. Philadelphia, PA: Lippincott Williams & Wilkins, 2001 .
- 13- Gleizes O, Desselberger U, Tatochenko V, et al. Nosocomial rotavirus infection in European countries: a review of the epidemiology, severity and economic burden of hospital-acquired rotavirus disease. *Pediatric Infectious Disease Journal* 2006;25(1 Suppl):S12-21
- 14- Marshall J, Botes J, Gorrie G, et al. Rotavirus detection and characterisation in outbreaks of gastroenteritis in aged-care facilities. *Journal of Clinical Virology* 2003;28:331-40
- 15- Al – Megrin WA .Intestinal parasites infection among immunocompromised patients in Riyadh, Saudi Arabia . *Pak J Biol Sci.* 2010 Apr 15;13(8):390-4..
- 16- Snelling T, Cripps T, Macartney K, et al. Nosocomial rotavirus infection in an Australian children's h
- 17- ROTTERDAM, H. & TSANG, P. - Gastrointestinal disease in the immunocompromised patient. *Hum. Path.*, **25**: 1123-1140, 1994 .
- 18- SNELLER, M. & CLIFFORD, H. - Infections in the immunocompromised host. In: RICH, R. *Clinical immunology principles and practices*. St Louis, Mosby, 1996. p. 579-593.
- 19- Baron E.J. , Peterson L.R. , Finegold S.M. , *Diagnostic microbiology* , 9<sup>th</sup> Edition , United States of America ,Mosby Co. , 1994; 786 : 791 -2 .
- 20- Garcia LS. And Bruckner D. A. ,*Diagnostic medical parasitology* ,2<sup>nd</sup> edition , Washington , D.C ,United States of America ,1993 ;49 -57 .
- 21- [http://www.curesearch.org/our\\_research/index\\_sub.aspx?id=1469](http://www.curesearch.org/our_research/index_sub.aspx?id=1469).
- 22- Malkin D., *Cancers of Childhood* , In :Devita V. T. ,Hellman S. , Rosenberg S .A .(Eds.) , *Cancer principles and practice of oncology* , Part 3 , 5<sup>th</sup> Edition Philadelphia , New York , Lippincott – Raven , 1997: p . 2083 – 91 .
- 23- Yacoub A. ,AL – Sadoon I. , Hassan G . , et al , Depleted Uranium and health of people in Basrah : An epidemiological evidence , *Med . J. Basrah Uni .* , 1999 ;17 : 17 -25
- 24- Campbell K ., *Chronic Myeloid Leukemia* , *Leukemia Research found* , 1997; p.7 .
- 25- Karimi M, Mehrabani D, Yarmohammadi H, Jahromi FS. The prevalence of signs and symptoms of childhood leukemia and lymphoma in Fars Province, Southern Iran . *Cancer Detect Prev.* 2008;32(2):178-83. Epub 2008 Jul 16.
- 26- Al-falahi RF (2002) .Role of Rotavirus gastroenteritis in children under three year ,of age hospitalized in AL-Ramadi Matirnty and children Hospital. Diploma thesis. The Collage of Medicine .University of AL-Anbar.
- 27- Ahmed HM , Coulter JB ,Nakagomi O., Hart CA ,Zaki JM . ,AL-Rabaty AA., Dove W. ,Cunliffe NA. Molecular characterization of Rotavirus gastroenteritis strains Iraqi Kurdistan .*Emerg. Infect . Dis.* 2006 May , 12(5) :824- 6 .
- 28- Ali M. Kheyami, Toyoko Nakagomi, Osamu Nakagomi, Winifred Dove, C. Anthony Hart, and Nigel A. Cunliffe . Molecular Epidemiology of Rotavirus Diarrhea among Children in Saudi Arabia: First Detection of G9 and G12 Strains. *J Clin Microbiol.* 2008 April; 46(4): 1185–1191.

- 29- Vesikari T., MakiM.,sarkkinen H.K., Arstila P.P and Halonen P.E .Rotavirus , Adenovirus ,and non- Viral enteropathogens in diarrhea . Arch. Dis. Child (1981); 56 : 264-270.
- 30- Pabst WL., M. Altwegg , C . Kind , S. Mirjanic ,D. Hardegger and D .Nadal .(2003) .Prevalence of enteroaggregative Escherichia coli among children with and without diarrhea in Switzerland .J .Clin .Microbiol . 41: 2289-2293 .
- 31- Branth P.V. ,Steinsland H., Fischer T.K., Perch M ., Scheutz F., Dias F., Aaby P., Molbak K., and Sommerfelt H. (2003). Cohort Study of Guinean Children: Incidence, Pathogenicity, Conferred Protection, and Attributable Risk for Enteropathogens during the First 2 Years of Life . J .Clin . Microbiol. 2003 September; 41(9): 4238–4245.
- 32- AL-Ani R.T.M. (2004) . Microbial causes of diarrhea in Neonates and Infants in Ramady city . M.Sc. Thesis .The collage of medicine University of AL-Anbar .
- 33- Azab ME , Mohamed NH , Salem SA , Safan EH , Bebars MA, Sabry NH , et al . Parasitic infections associated with malignancy and leprosy . J .Egypt soc parasitol 1992 , 22 : 59 -70.
- 34- Gilger MA, Matson DO, Conner ME, Rosenblatt HM, Finegold MJ, Estes MK Extraintestinal rotavirus infections in children with immunodeficiency. J Pediatr. 1992 Jun;120(6):912-7.

## الخلاصة

على مدى فترة خمسة أشهر ، تمت دراسة فايروس الروتا وإصابات الطفيلية في المرضى الذين يعانون و يستلمون علاج للأمراض السرطانية في محافظة البصرة.

العدد الكلي للمرضى في هذه الدراسة هو ٥٨ مريض [٣٨ ذكر و ٢٠ أنثى] استخدمت لتشخيص كل عينات البراز التي جمعت من المرضى طريقة التركيز ( فورمالين – أيثر ) وطريقة اللطخة المباشرة.

أعلى نسبة للإصابة في هذه الدراسة وجدت بين الذكور منها في الإناث وكذلك

وجد ارتفاع في نسبة الإصابة في المناطق الريفية منها في المدينة واغلبهم

مع إصابات طفيلية مختلطة. كلهم بأعراض مرضية وأكثر المرضى إيجابيا للروتا فايروس يعانون من الإسهال المزمن (٦٦,٧ %) أعلى من الإسهال المتقطع (٣٣,٣ %).