

Microbial spectrum and antibiotic sensitivity in cirrhotic patients with spontaneous bacterial peritonitis

IRFAN AHMAD¹, MUHAMMAD SHAHBAZ HUSSAIN ², MUHAMMAD SHOAIB AKHTAR³,¹MBBS, FCPS (Medicine), FCPS (Gastroenterology) Professor, Medical unit 1, Sheikh Zayed Medical College/Hospital, Rahim Yar Khan²MBBS, M Phil (Microbiology) Professor of Microbiology Sheikh Zayed Medical College/Hospital, Rahim Yar Khan³Muhammad Shoaib Akhtar MBBS, FCPS (Medicine) Senior Registrar Medicine Medical unit 1, Sheikh Zayed Medical College/Hospital, Rahim Yar KhanCorrespondence to Dr. Irfan Ahmad Email: uhirfan@yahoo.com +92 333 4365708

ABSTRACT

Objective: There are reports of changing microbial profile of ascitic fluid in spontaneous bacterial peritonitis (SBP) and developing resistance of these bacteria to commonly used antibiotics. This study was done to determine the micro-organism causing SBP and their sensitivity to various antibiotics.

Setting and Methods: This observational study was done in Medical department of Sheikh Zayed Medical College/Hospital, Rahim Yar Khan from March to November 2019. Thirty five cirrhotic patients with ascites admitted in ward for various reasons and having positive ascitic fluid culture were included in the study. Exclusion criteria was presence of secondary bacterial peritonitis. Ten ml of ascitic fluid was inoculated in blood culture bottle and sent immediately to hospital laboratory.

Results: The mean age of these 35 patients was 48.94 ± 13.51 years with a range of 19 to 80 years. Twenty (57.1 %) patients were male and 15 (42.9 %) were female. Bacteria that had caused SBP were E.coli (62.9 %), staph aureus (11.4 %), klebsiella (8.6 %), streptococci (8.6 %), gram positive cocci (5.7 %) and pseudomonas aeruginosa (2.8 %). Imipenem had high sensitivity rate (100 %) along with amikacin (82.9 %) and cefoperazone-sulbactam (68.6 %). Sensitivity of these organisms to other commonly used antibiotics were: ciprofloxacin 57.1 %, ofloxacin 40 %, norfloxacin 37.1 %, ceftazidime 34.3 %, ceftriaxone 31.4 % and piperacillin-tazobactam 25.7 %.

Conclusion: We found that E.coli was the commonest bacteria causing SBP, and ceftriaxone and ciprofloxacin have significantly high resistance rate in these patients.

Key words: cirrhosis, ascitic fluid, spontaneous bacterial peritonitis, antibiotic sensitivity

INTRODUCTION

Spontaneous bacterial peritonitis (SBP) usually occurs in cirrhotic patients with ascites. Higher the Model for End stage Liver Disease (MELD) score, greater is the risk of SBP.¹ If diagnosis is delayed, there is a risk of developing shock and multi-organ failure.² It is associated with higher mortality rate.³ If a patient has an attack of SBP, there is increased risk of recurrence.⁴ Total protein in ascitic fluid correlates inversely with the risk of SBP.⁵ Diagnosis of SBP is made if ascitic fluid culture is positive and/or fluid neutrophil count is more than 250/cmm and there is no evidence of secondary (surgical) peritonitis.⁶ Immediate inoculation of culture bottle increases the sensitivity of positive culture result.⁷

All cases of SBP are caused by monomicrobial infection. Polymicrobial infection favors the diagnosis of secondary peritonitis. Most common pathogen is E.coli;⁸ other common organisms are Klebsiella, enterococci and streptococci. If there is clinical suspicion of SBP, antibiotic should be started immediately after paracentesis.⁹ Third generation cephalosporin is a reasonable choice, especially cefotaxime or ceftriaxone.^{10,11} Fluoroquinolones like ciprofloxacin or ofloxacin can also be given.¹² In resistant cases piperacillin/tazobactam or carbapenem should be used.¹³

We normally give ceftriaxone or ciprofloxacin to our patients who are suspected to be suffering from SBP. As there are reports of increasing resistance to ceftriaxone and quinolones,^{14,15} we planned this study to find out types of organisms in culture positive ascitic fluid in our SBP patients and their sensitivity to various antibiotics. It would help us to choose the most appropriate antibiotic in SBP patients.

MATERIALS AND METHODS

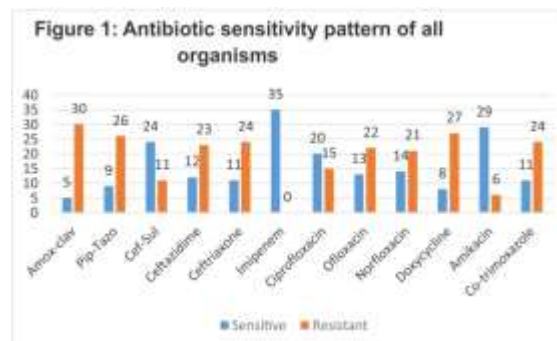
The study was conducted from March 20 to November 23, 2019 in Medical department, Sheikh Zayed Medical College/Hospital, Rahim Yar Khan. It was a descriptive study. Study protocol was approved from Institutional Review Board and Ethical Committee. Thirty five cirrhotic patients with ascites admitted in Medical departmentsuffering from spontaneous bacterial peritonitis having positive ascitic fluid culture were included in the study. Inclusion was regardless of whether SBP had been classical (culture positive and neutrophil count > 250 cells/cmm) or bacterascites (culture positive but neutrophil count < 250 cells/cmm). Patients were excluded from the study if they had intra-abdominal surgically

treatable disease, if ascitic fluid culture showed polymicrobial growth as it indicated secondary peritonitis. Peritoneal paracentesis was done using standard technique in cirrhotic patients with ascites who had been admitted due to fever, abdominal pain/tenderness, hematemesis or melena, hepatic encephalopathy or decreased urine output. Five ml fluid was sent for albumin, total leucocyte count and differential count. Ten ml fluid was inoculated immediately in 100 ml TSB tryptic soy broth and sent to our hospital's laboratory for culture/sensitivity test. Blood tests like complete blood count, liver function test, prothrombin time, albumin, sodium and creatinine were also performed. SPSS software version 25 was used for analysis of data. The qualitative variables like gender, types of organisms and antibiotic sensitivity were described as frequency and percentage, and analyzed by Chi-square test. The quantitative variables like age of patient were expressed as mean \pm SD and range, and analyzed by Student's t-test. A p value of < 0.05 was considered statistically significant.

RESULTS

Table 1: Frequency of ascitic fluid organisms in SBP

Organism	Frequency (percentage)
E.coli	22 (62.9 %)
Klebsiella	3 (8.6 %)
Gram positive cocci	2 (5.7 %)
Streptococcus	3 (8.6 %)
Staph aureus	4 (11.4 %)
Pseudomonas aeruginosa	1 (2.8 %)



The mean age of 35 included patients was 48.94 ± 13.51 years and age range was 19 to 80 years. Twenty (57.1 %) patients were male and 15 (42.9 %) were female. HCV was the cause of cirrhosis in 30 (85.7 %) patients, HBV in 3 (8.6 %) and both HCV and HBV in 2 (5.7 %). The presenting illness was hepatic encephalopathy in 22 (62.9 %), abdominal pain plus fever in 8 (22.8 %), variceal bleeding in 3 (8.6 %) and abdominal distension with fever in 2 (5.7 %) patients.

E.coli was the commonest organism causing SBP as shown in Table 1 while gram positive organisms was responsible for SBP in 9 patients. There was culture positive neutrocytic ascites in 24

(68.6 %) and culture positive non-neutrocytic (bacterascites) ascites in 11 (31.4 %) patients.

Figure 1 shows overall sensitivity/resistance pattern of commonly used antibiotics in our SBP patients. Imipenem has 100 % sensitivity and amikacin has sensitivity of 82.9 %. The most commonly used antibiotic, ceftriaxone had only 31.4 % sensitivity. The sensitivity of these antibiotics to various organisms found in our patients is shown in Table 2. E.coli which is the most common organism is sensitive to imipenem, amikacin and cefoperazone-sulbactam combination but resistant to ceftriaxone and quinolones.

Table 2: Sensitivity pattern of ascitic fluid organisms in SBP to various antibiotics

Antibiotic	Sensitivity	Organism						Total
		E.coli	Klebsiella	Gram + cocci	Streptococci	Staph aureus	P.aeruginosa	
Amoxclav	Sensitive	2	1	0	0	2	0	5
	Resistant	20	2	2	3	2	1	30
Pip-Tazo	Sensitive	2	1	2	2	1	1	9
	Resistant	20	2	0	1	3	0	26
Cef-Sul	Sensitive	16	2	1	2	2	1	24
	Resistant	6	1	1	1	2	0	11
Ceftazidme	Sensitive	4	3	2	1	1	1	12
	Resistant	18	0	0	2	3	0	23
Ceftriaxone	Sensitive	6	0	2	3	0	0	11
	Resistant	16	3	0	0	4	1	24
Imipenem	Sensitive	22	3	2	3	4	1	35
	Resistant	0	0	0	0	0	0	0
Ciprofloxacin	Sensitive	9	2	2	3	3	1	20
	Resistant	13	1	0	0	1	0	15
Ofloxacin	Sensitive	7	2	2	1	1	1	14
	Resistant	15	1	0	2	3	0	21
Norfloxacin	Sensitive	7	2	1	1	1	1	13
	Resistant	15	1	1	2	3	0	22
Doxycycline	Sensitive	5	1	1	1	0	0	8
	Resistant	17	2	1	2	4	1	27
Amikacin	Sensitive	21	2	1	1	3	1	29
	Resistant	1	1	1	2	1	0	6
Co-trimoxazo	Sensitive	4	1	2	2	2	0	11
	Resistant	18	2	0	1	2	1	24

DISCUSSION

To reduce morbidity and mortality in spontaneous bacterial peritonitis treatment should be started early.⁹As culture and sensitivity report of ascitic fluid takes few days, empirical antibiotic therapy has to be given. Gram negative organisms like E.coli are the most common cause of SBP but recent trends show that gram positive organisms are increasingly causing SBP.¹⁶For empirical treatment, cefotaxime,¹⁰ ceftriaxone¹¹ and fluoroquinolones¹² have been recommended but over last few years there is concern about resistance to these antibiotics.¹⁴

Spontaneous bacterial peritonitis is caused by a single organism and E.coli has been the most common bacteria. There are reports of increasing number of gram positive bacteria like staphylococcus and streptococcus but E.coli still remains the commonest organism. The frequency of E.coli detection in our study is 62.9 %. Similar reports have been shown by many other investigators with E.coli detection rate of 55 to 73 %^{8, 17-22} and this trend remained same over past two decades. The other common organisms were Klebsiella, staphylococcus aureus, streptococci, acinetobacter and pseudomonas.^{8, 15, 18, 19, 22, 23-25}

Iqbal S, et al (2004) found that all organisms involved in SBP were sensitive to third generation cephalosporin and fluoroquinolones.¹⁸ It had been recommended that cefotaxime, ceftriaxone and fluoroquinolones should be given empirically when there is suspicion of SBP.^{10,11, 12}Over the last few years, resistance to these antibiotics have been described. Bibi S, et al (2015) found that resistance to third generation cephalosporin was 78 % and to fluoroquinolone was 69.6 %.²⁰Other studies found 35 %, 76 % and 62 % sensitivity to ceftriaxone,^{17, 26, 27} 65 % to cefotaxime,²⁷ and 31 % and 35 % to ciprofloxacin.^{17, 26}Our study revealed 31 % sensitivity with ceftriaxone and 57 % with ciprofloxacin. During

recent years, sensitivity of ascitic fluid bacteria to imipenem, amikacin and cefoperazone-sulbactam has been found high.^{15, 25, 26, 28}Our results were sensitivity of 100 % to imipenem, 83 % to amikacin and 69 % to cefoperazone-sulbactam.

The reason for increasing gram positive bacteria in ascitic fluid in patients with SBP and increasing resistance previously commonly used antibiotics may be longer survival of cirrhotic patients with repeated use of antibiotics, multiple hospital admissions and emergence of nosocomial infection. Now there is enough evidence that recommendations for empirical antibiotic treatment of SBP patients be changed.

CONCLUSION

In our study E.coli was found to be the most common organism causing SBP. Gram positive bacteria were responsible for about one quarter of cases. Organisms causing SBP had shown significant resistant to ceftriaxone and ciprofloxacin which are commonly used for this purpose while imipenem, amikacin and cefoperazone-sulbactam had high sensitivity. In the light of these findings empirical antibiotic treatment for SBP should be changed accordingly.

REFERENCES

1. Obstein KL, Campbell MS, Reddy KR, Yang YX. Association between model for end-stage liver disease and spontaneous bacterial peritonitis. *Am J Gastroenterol* 2007; 102 (12):2732-6.
2. Hoefs JC, Runyon BA. Spontaneous bacterial peritonitis. *Dis Mon* 1985; 31 (9):1-48.
3. Kim JJ, Tsukamoto MM, Mathur AK, Ghomri YM, Hou LA, Sheibani S, et al. Delayed paracentesis is associated with increased in-hospital mortality in patients with spontaneous bacterial peritonitis. *Am J Gastroenterol* 2014; 109 (9):1436-42.

4. Chang CS, Chen GH, Lien HC, Yeh HZ. Small intestine dysmotility and bacterial overgrowth in cirrhotic patients with spontaneous bacterial peritonitis. *Hepatology* 1998; 28 (5):1187-90.
5. Runyon BA. Patients with deficient ascitic fluid opsonic activity are predisposed to spontaneous bacterial peritonitis. *Hepatology* 1988; 8 (3):632-5.
6. Dever JB, Sheikh MY. Spontaneous bacterial peritonitis – bacteriology, diagnosis, treatment, risk factors and prevention. *Aliment Pharmacol Ther* 2015; 41 (11):1116-31.
7. Wong CL, Holroyd-Leduc J, Thorpe KE, Straus SE. Does this patient have bacterial peritonitis or portal hypertension? How do I perform a paracentesis and analyze the results? *JAMA* 2008; 299 (10):1166-78.
8. Waseem Sarwar Malghani, Farooq Mohyud Din Chaudhary, Muhammad Ilyas, Asma Tameezud Din, Asim Tameezud Din. Spontaneous bacterial peritonitis; spectrum of bacterial flora causing spontaneous bacterial peritonitis in patients with liver cirrhosis. *Professional Med J* 2018; 25 (5):749-52.
9. Runyon BA, AASLD. Introduction to the revised American Association for the Study of Liver Diseases Practice Guideline management of adult patients with ascites due to cirrhosis 2012. *Hepatology* 2013; 57 (4):1651-3.
10. Chavez-Tapia NC, Soares-Weiser K, Brezis M, Leibovici L. Antibiotics for spontaneous bacterial peritonitis in cirrhotic patients. *Cochrane Database Syst Rev* 2009; 21;(1):CD002232.
11. Fernández J, Ruiz del Arbol L, Gómez C, Durandez R, Serradilla R, Guarner C, et al. Norfloxacin vs ceftriaxone in the prophylaxis of infections in patients with advanced cirrhosis and hemorrhage. *Gastroenterology* 2006; 131 (4):1049-56.
12. Terg R, Cobas S, Fassio E, Landeira G, Rios B, Vasen W, et al. Oral ciprofloxacin after a short course of intravenous ciprofloxacin in the treatment of spontaneous bacterial peritonitis: results of a multicenter, randomized study. *J Hepatol* 2000; 33 (4):564-9.
13. Fernández J, Prado V, Trebicka J, Amoros A, Gustot T, Wiest R, et al. Multidrug-resistant bacterial infections in patients with decompensated cirrhosis and with acute-on-chronic liver failure in Europe. *J Hepatol* 2019; 70 (3):398-411.
14. Fernández J, Acevedo J, Castro M, Garcia O, de Lope CR, Roca D, et al. Prevalence and risk factors of infections by multiresistant bacteria in cirrhosis: a prospective study. *Hepatology* 2012; 55 (5):1551-61.
15. Khurram Baqai, Nasir Laique, Faisal Ziauddin. Ascitic fluid cultivated organisms and their antimicrobial resilience pattern in patients with liver cirrhosis. *Pak J Med Dentistry* 2018; 7 (4):37-44.
16. Ryan G, Arslan K, Sharif M, Suril P, Aparna S, Stephen A. Changing trends of microbial organisms causing spontaneous bacterial peritonitis (SBP) in a large tertiary care centre in the Tennessee River Valley. *Am J Gastroenterol* 2017; 112 (Suppl): S508-S509.
17. Kadla SA, Mir MM, Rather MK, Shah NA, Wani ZA. Study of etiological profile and resistance pattern of spontaneous bacterial peritonitis in chronic liver disease. *IJHG* 2019; 5 (1):10-5.
18. Iqbal S, Iman NU, Alam N, Rahman S. Incidence of spontaneous bacterial peritonitis in liver cirrhosis, the causative organisms and antibiotic sensitivity. *J Postgrad Med Inst* 2004; 18 (4): 614-9.
19. Imran M, Hashmi SN, Altaf A, Rashid H, Hussain T. Spontaneous bacterial peritonitis. *Professional Med J* 2006; 13 (2):201-5.
20. Bibi S, Ahmed WD, Arif A, Khan F, Alam SE. Clinical, laboratory and bacterial profile of spontaneous bacterial peritonitis in chronic liver disease patients. *JCPSP* 2015; 25 (2): 95-9.
21. Memon AQ, Memon G, Khaskheli A. Spontaneous bacterial peritonitis in cirrhosis with ascites – an experience at PMCH Nawab Shah. *Med Channel* 1999. 5 (1): 31-4.
22. Zaman A, Kareem R, Mahmood R, Hameed K, Khan EM. Frequency of microbial spectrum of spontaneous bacterial peritonitis in established cirrhosis liver. *J Ayub Med Coll* 2011; 23 (4):15-7.
23. Ali A, Ullah Zia, Majeed M. Identification of causative organisms and role of ceftriaxone in spontaneous bacterial peritonitis (SBP) secondary to decompensated liver disease. *Int J Endorsing Health Sci Res* 2016. 4 (3):41-7.
24. Husain A, Qayyum A, Muhammad D, Javed M, Iqbal MN. Bacteriology of spontaneous bacterial peritonitis. *Professional Med J* 2007; 14 (4):551-5.
25. Prasad R, Renu K, Anand AK, Kumar A. Study on ascitic fluid culture and antimicrobial sensitivity profile in cirrhotic patients admitted in a medical college hospital. *IAIM* 2019; 6 (2):53-7.
26. Ali A, Abbasi AS. Changing antimicrobial sensitivity patterns in patients with spontaneous bacterial peritonitis: Can it be prevented? *Rawal Med J* 2018; 43 (4):598-602.
27. El-Bedewy, Tamer A, El-Sebaey, Mohamed A, Okda, Hanaa I, et al. Microbial study of spontaneous bacterial peritonitis in Tanta University Hospitals. *Egyptian Liver Journal* 2017; 7 (1-2):5-8.
28. Wieser A, Li H, Zhang J, Liss I, Markwardt D, Hornung R, et al. Evaluating the best empirical antibiotic therapy in patients with acute on chronic liver failure and spontaneous bacterial peritonitis. *Dig Liver Dis* 2019; 51 (9):1300-7.