Bacterial Pneumonia: Comparison between Diabetics and Non-Diabetics

Dr. M. Vamsi Krishna¹, Dr. G. Swarna Latha Devi.², Dr. J. Dattu Raj³

²Professor and HOD, ASRAM Medical College, Hospital Eluru

Abstract: To determine the causative organisms, anti microbial susceptibility, and outcome of community- and hospital acquired pneumonia in diabetics and to compare this with non-diabetics, sputum cultures done at Asram Medical College, Hospital, ELURU in the period between October 2016 and December 2018 were reviewed. A total of 354 cases were studied, of which 125 (35%) were diabetics. Diabetic patients were older with a male predominance compared to non-diabetics. H. influenza was the commonest pathogen in community-acquired pneumonia (CAP) in both diabetics and non-diabetics, but there was a predominance of Staphylococcusaureus in diabetics compared to non-diabetics. Gram-negative bacilli were the commonest pathogens in hospital-acquired pneumonia (HAP) in both diabetics and non-diabetics and non-diabetics and erythromycin were used empirically in CAP while aminoglycosides, fluoro- quinolones and imipenem were used in HAP in both diabetics and non-diabetics. No significant difference in mortality was found between diabetics and non-diabetics, for either CAP or HAP

Keywords: Pneumonia, Diabetics, Microorganism, Mortality, Treatment

1. Introduction

Diabetes mellitus (DM) is often identified as an independent risk factor for developing respiratory tract infections. Diabetic patients are predisposed to colonization and pneu- monia because of disease-associated impairment in host defensive functions ^[1, 2]. Also, they are more liable to develop complications such as bacteremia, delayed resolution, and recurrent pneumonia^[3]. Pneumonia is the leading cause of hospitalization and mortality [4]. Several studies have shown that the use of appropriate antimicrobial therapy can improve outcome with survival rate reaching 70%-80%^[2]. The aim of this study was to determine the causative organisms, antimicrobial susceptibility of community- and hospital-acquired pneumonia in diabetics, and to report on any difference between them and nondiabetics.

2. Patients and Methods

For this study, we reviewed sputum cultures of patients above the age of 18 years, performed in the period between OCTOBER 2016 and December 2018. Sputum cultures positive for bacteria were analyzed, those positive for acid-fast bacilli were excluded.

Sputum samples were processed by gram stain and culture. Cultures were performed on 5% sheep blood agar (oxoid) and chocolate agar. Bacteria isolated from sputum culture were considered presumptive etiologic pathogens if they were compatible with the predominant organisms present on gram stain and if cultured in abundant growth or in pure growth.

Pneumonia was diagnosed according to the American Thoracic Society criteria [7]. Cases were classified into hospital-acquired pneumonia (HAP) if the sputum culture was first positive more than 72h after admission, excluding any infection that was incubating at the time of admission. Community-acquired pneumonia (CAP) was defined by a positive sputum culture with in 72 h after admission or by a positive culture performed as outpatient.

For each patient with pneumonia, we recorded age, gender, out- come, type of organisms isolated and their antimicrobial susceptibility, empiric use of antimicrobial agents, presence of DM ,treatment regimen for DM, and degree of control (good control was defined as a glycated hemoglobin (HbA1c) <7%).

The in vivo antibacterial susceptibility of the isolated bacteria was determined by the disk diffusion method. Patients were divided into two groups according to the presence or absence of DM. Statistical analysis was done using Statistical Package for Social Sciences (SPSS) computer soft- ware and p values less than 0.05 were considered significant.

3. Results

From a total of 605 sputum cultures done during the study period, 354 cases with a positive culture were included in the study. Of these, 125 (35%) were diabetics, havingameanageof59.4 \pm 14.0yearsvs53.7 \pm 20.6years for the non-diabetics (*p*=0.006). Male predominance was noticed in the diabetic group: the male female ratio was 3:1 vs 1.2:1 for non-diabetics (*p*<0.001). Most of the diabetic patients were using oral hypoglycemia agents for blood glucose control (n=75, 60%); of the remainder, 38 (30%) were on insulin, 7 (6%) on diet, and 5 (4%) on combination therapy. There were 86 (69%) diabetics who were poorly controlled.

Volume 8 Issue 2, February 2019 <u>www.ijsr.net</u> Licensed Under Creative Commons Attribution CC BY

International Journal of Science and Research (IJSR) ISSN: 2319-7064 Impact Factor (2018): 7.426

and cares with commany acquired provincing									
Pathogen	Diabetics	Non-diabetics	p value						
	n (%)	n (%)							
Streptococcus pneumoniae	1 (4)	3 (5)	0.3						
Staphylococcus aureus	6 (23)	6 (10)	0.02						
Haemophilus influenzae	13 (50)	31 (53)	0.8						
Moraxella catarrhalis	2 (8)	8 (14)	0.6						
Pseudomonas spp.	3 (12)	5 (8)	0.7						
Klebsiella spp.	1 (4)	3 (5)	0.6						
Enterobacter spp.	-	1 (2)	0.5						
Acinetobacter spp.	-	1 (2)	0.5						
Streptococcus viridans	-	1 (2)	0.5						

 Table 1: Pathogens isolated 26 diabetics and 59 nondiabetics with community-acquired pneumonia

 Table 2: Pathogens isolated from 99 diabetics and 170 nondiabetics with hospital-acquired pneumonia

diabeties with hospital-acquired pheumonia									
Pathogen	Dia	betics	Non-	p value					
	n	(%)	n						
Streptococcus pneumoniae	3	(3)	1	(1)	0.1				
Staphylococcus aureus	14	(14)	27	(16)	0.6				
Haemophilus influenzae	13	(13)	26	(15)	0.9				
Moraxella catarrhalis	6	(6)	9	(5)	0.4				
Pseudomonas spp.	30	(30)	43	(25)	0.5				
Klebsiella spp.	6	(6)	15	(9)	0.5				
Enterobacter spp.	9	(9)	14	(8)	0.4				
Proteus spp.	2	(2)	3	(2)	0.7				
Escherichia coli	2	(2)	8	(5)	0.4				
Enterococci	1	(1)		-	0.1				
Citrobacter spp.	1	(1)	3	(2)	0.7				
Acinetobacter spp.	6	(6)	7	(4)	0.2				
Others ^a	6	(6)	14	(8)	0.3				

^aStentrophomonas maltophilia, Serratia spp.

 Table 3: Antibiotic sensitivity of some isolates in community and hospital acquired pneumonia

Antimicrobial	Pathogens, n(%)												
agent	Community acquired							Hospital acquired					
	S. pneu	imoniae	H. infli	uenzae	Moraxella	S. at	ureus	Pseudom	onasspp. Enterobacterspp.		acterspp.		
	D	ND	D	ND	D	ND	D	ND	D	ND	D	ND	
	n=1	n=3	n=13	n=31	n=2	n=8	n=14	n=27	n=30	n=43	n=9	n=14	
Penicillin	-	1 (33)	-	-	1 (50)	1 (13)	-	4(15)	-	-	6 (67)	2 (18)	
Ampicillin	1(100)	3 (100)	9 (70)	23(74)	1 (50)	8 (100)	2(14)	-	2 (7)	-	-	1 (7)	
Co-amoxyclav	1(100)	3 (100)	9 (70)	23(74)	2 (100)	8 (100)	2(14)	-	2 (7)	1 (2)	2 (22)	1 (7)	
Ciprofloxacin	-	-	10 (77)	26(84)	1 (50)	8 (100)	2(14)	-	24 (79)	30 (70)	7 (78)	11 (79)	
Cefuroxim	1(100)	3 (100)	12 (92)	30(97)	1 (50)	8 (100)	2(14)	-	1 (4)	-	2 (22)	2 (14)	
Erythromycin	1(100)	3 (100)	6 (46)	19(60)	2 (100)	7 (86)	5(36)	18 (67)	-	1 (2)	4 (44)	3 (21)	
Oxacillin	1(100)	3 (100)	-	-	-	-	9(64)	22 (81)	-	-	-	-	
Vancomycin	-	3 (100)	-	1 (3)	-	-	12(86)	18 (67)	-	-	-	2 (14)	
Azterionam	-	1 (33)	4 (31)	17(55)	-	-	-	-	20 (67)	25 (59)	2 (22)	7 (50)	
Ceftazidim	-	-	1 (8)	-	-	-	-	-	21 (71)	29 (68)	2 (22)	2 (14)	
Ceftriaxon	-	3 (100)	10 (77)	27(87)	-	1 (13)	-	-	8 (27)	9 (21)	2 (22)	7 (50)	
Gentamycin	-	1 (33)	-	1 (3)	-	-	2(14)	8(30)	27 (79)	36 (84)	7 (78)	7 (50)	
Amikacin	_	_	_	-	_	_	_	-	25 (82)	36 (85)	9 (100)	14 (100)	
Imepenum	-	-	1 (8)	2 (6)	-	_	-	-	15 (50)	29 (68)	9 (100)	13 (93)	
Pipracillin	_	_	_	-	-	-	-	-	22 (73)	30 (70)	2 (22)	7 (50)	

D, Diabetics; ND, Non-diabetics; co-amoxyclav, a combination of amoxycillin and clavulanica acid

Of the 354 patients with a positive sputum culture, 85 (24%) were diagnosed as having CAP, while the remaining 269 patients (76%) had HAP. Among the patients diagnosed with CAP, 26 (31%) were diabetics while among those with HAP there were 99 diabetics (37%). Empiric antimicrobial treatment was in use at the time of specimen collection in 81 (95%) of patients with CAP vs. 231 (86%) of patients with HAP (p=0.2). Most of the patients were started on two empiric antimicrobial agents: 72 of 81 (85%) in CAP and 212 of 231 (92%) in HAP (p=0.09). Haemophilus influenzae was the commonest cause of CAP in both diabetics and nondiabetics (Table 1). There was a predominance of infections by Staphylococcus aureus among diabetics with CAP compared to non-diabetics. Gram-negative bacilli were the commonest cause of HAP in both diabetics and nondiabetics (Table 2). Ampicillin, co-amoxyclav (a combination of amoxycillin and clavulanica acid), flouroquinolone, second-generation cephalosporins and erythromicin were used empirically in CAP, while aminoglycosides, flouroquinolones and impenem were used in HAP in both diabetics and non-diabetics (Table3).

4. Discussion

Pneumonia is one of the most common infectious in India, It is clear from our study that almost one-third of the cases admitted with bacterial pneumonia were diabetics. Diabetics have alterations of pulmonary host defenses [11] which make them more susceptible to infection. Advanced age is also associated with immune changes that increase the risk of pneumonia ^[12]. In this study, diabetics were older than non-diabetics; therefore they were at increased risk for pneumonia also for their age.Several studies have shown that S. pneumoniae is the most common pathogen isolated in CAP [13-15]. Other organisms isolated in CAP include H. influenzae, atypical bacteria, Moraxella catarrhalis, S. aureus, and gram-negative bacilli [16-18]. Interestingly, this study showed that H. influenzae was the commonest pathogen isolated in CAP in both diabetics and non-diabetics, while S. pneumoniae was isolated in a smaller percentage. Some studies have found that sputum cultures were negative in about 50% of patients with pneumococcal bacteremia, and that the rate of isolation increases when more invasive methods are used for obtaining specimens, such as trans- tracheal aspiration

Volume 8 Issue 2, February 2019 www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

International Journal of Science and Research (IJSR) ISSN: 2319-7064 Impact Factor (2018): 7.426

which eliminates contaminating oropharyngeal flora [16, ^{19].} Due to the retrospective design of this study, invasive methods for obtaining sputum specimens were not used for all the cases. Another possible reason for the low isolation rate of S. Pneumoniae is the use of antimicrobial agents at the time of specimen collection ^{[20].} The majority of these patients were started on empiric antimicrobial agents. S. aureus is a major pathogen of CAP in diabetics compared to non-diabetics. This observation can be attributed to the high nasal carriage rate of S. aureus in diabetics where it reached 30% compared to 11% inhealthy individuals^[21], The rate of nasal carriage of S.aureus is directly related to the glycosylated hemoglobin (HbA1_c)level^[21] The ATS recommends to use empiric treatment for pneumonia as pathogen identification can be difficult ^[26]. We found that co-amoxyclav, ampicillin, flouroquinolones, second-generation cephalosporins, and erythromycin were used empirically to treat CAP in both diabetics and non- diabetics, while in severe cases of CAP (especially in poorly controlled diabetics Staphylococcus can be combatted with cloxacillin or vancomycin. In HAP, aminoglycosides, flouroquinolones, and imipenem were used in both diabetics and non-diabetics, which is in agreement with what has been recommended by others [10, 27-31].One of the limitations of microbiological diagnosis of pneumonia is the lower prevalence of positive sputum cultures due to either the use of empiric antimicrobial agents at the time of specimen collection or the failure to use of more invasive methods for obtaining sputum specimens. Due to the retrospective design of this study, these limitations could not be avoided.

References

- [1] Celis R, Torres A, Gatell JM, Almela M, Rodringnez-Roisin R, Agusti-Vidal A (1988) Nosocomial pneumonia: A multivariate analysis of risk and prognosis. Chest 93(2):318–324
- [2] Carvin DE, Steger KA, Barber TW (1991) Preventing noso- comial pneumonia: a state of the art and perspectives for the 1990s. Am J Med91(3B):44S–53S
- [3] KirtlandS, Winterbauer R, Dreis D, Pardee NE, Springmeyer SC (1994) A clinical profile of chronic bacterial pneumonia. Report of 115 cases. Chest106:15–22
- [4] Freeman C, Nicolan DP (1999) Community acquired pneu- monia in the long-term care setting: the other community. Consult Pharm11:1259–1273
- [5] Murry PR, Washington JA (1975) Microscopic and bacterio- logic analysis of expected sputum. Mayo Clin Proc 50:339–344
- [6] Murry PR, Baron EJ, Pfaller MA, Tenover FC, Yolker RH (eds) (1995) Manual clinical microbiology. American Society for Microbiology, Washington DC
- [7] Factors associated with mortality in severe community-**acquired pneumonia**: A multicenter cohort <u>study</u>R Espinoza, JRL e Silva, <u>A</u> <u>Bergmann</u>of **critical care**, 2019 – Elsevier.
- [8] (1985) Diabetes mellitus: Report of a WHO Study Group. World Health Organization, Geneva, pp1–113
- [9] (1995) National Committee for Clinical Laboratory Standards. Performance standards for antimicrobial suscepti- bility testing (M100-S6). NCCLS, Villanova

(15, n.14)

- [10] Garibaldi RA (1985) Epidemiology of community acquired respiratory infections in adults: incidence, etiology and impact india Med78:321–327
- [11] Moutschen MP, Scheen AJ, Lefebvre PJ (1992) Impaired immune responses in diabetes mellitus. Analysis of the fac- tors and mechanisms involved. Relevance to the increased susceptibility of diabetic patients tosepticinfections.Diabete Metab18:187–201
- [12] Gyetko MR, Toews GB (1993) Immunology of the aging lung. Clin Chest Med14:379–391
- [13] Ishida T, Hashimoto T, Arita M, Ito I, Osawa M (1998) Etiology of community acquired pneumonia in hospitalized patients: a 3-year prospective study in Japan. Chest 114(6):1588–1593
- [14] Bernstein M (1999) Treatment of community acquired pneu- monia. IDSA guidelines. Infectious Disease Society of America. Chest115:9S–13S
- [15] Lange M (2000) Community acquired pneumonia: an approach to antimicrobial therapy. Allergy Asthma Proc 21:33–38
- [16] Bartlett JG, Mundy LM (1995) Community acquired pneu- monia. N Engl J Med333:1618–1624
- [17] Pozzi E (1999) Community acquired pneumonia. The ORI- ONE Board. Monaldi Arch Chest Dis54(4):337– 344
- [18] Sopena N, Sabria M, Pedro Botet ML, Manterola JM, Matas L, Dominguez J, Modol JM et al (1999) Prospective study of community acquired pneumonia of bacterial etiology in adults. Eur J Clin Microbiol Infect Dis 18(12):852–858
- [19] Barrett Connor E (1971) The non value of sputum culture in the diagnosis of pneumococcal pneumonia. Am Rev Respir Dis103(6):845–848
- [20] Al-Hadramy MS, Altahawi AT, Shafi M (1988) Acute lower respiratory tract infections in Jeddah. Saudi Med J9(1): 34–39 Lipsky BA, Pecoraro RE, Chen MS, Koepsell TD (1987) Factors affecting staphylococcal colonization among NIDD outpatients. Diabetes Care 10:483–486
- [21] Niederman MS, Bass JB, Campbell GD (1993) Guidelines for the initial empiric therapy of community acquired pneumonia: proceedings of the American Thoracic Society consensus on- ference. Am Rev Respir Dis148:1418–1426
- [22] Schleupner CJ, Cobb DK (1992) A study of the etiology and treatment of nosocomial pneumonia in a community-based teaching hospital. Infect Control Hosp Epidemiol 13:515–525
- [23] Rouby JJ, Martin De Lassale E, Poete P, Nicolas MH, Bodin L, Jarlier V et al (1992) Nosocomial bronchopneumonia in the critically ill: histologic and bacteriologic aspects. AmRev Respir Dis146(4):1059– 1066
- [24] (1994) Guidelines for prevention of nosocomial pneumo- nia. Centers for Disease Control and Prevention. Respir Care 39(12):1191–1236
- [25] (2000) Community acquired pneumonia. Outpatient treat- mentofpatients16yearsand older. Institute of Clinical System Improvement. Post Grad Med107:246– 253
- [26] (1993) The British Thoracic Society Guidelines for the management of community acquired pneumonia in

Volume 8 Issue 2, February 2019

<u>www.ijsr.net</u>

Licensed Under Creative Commons Attribution CC BY

adults admitted to hospital. Br J Hosp Med49(5):346-350

- [27] Bartlett JG, Breiman RF, Mandell LA, File TM Jr (1998) Community-acquired pneumonia in adults: guidelines for management. The Infectious Diseases Society of America. Clin Infect Dis26(4):811–838
- [28] Quinn JP (1998) Clinical strategies for serious infection: a North American perspective. Diagn Microbiol Infect Dis 31(2):389–395
- [29] Kashuba AD, Nafzieger AN, Drusano GL, Bertino JS Jr (1999) Optimizing aminoglycoside therapy for nosocomial pneumonia caused by gram-negative bacteria. Antimicrob Agents Chemother43(3):623–629
- [30] Jones RN, Croco MA, Kugler KC, Pfaller MA, Beach MI (2000) Respiratory tract pathogens isolated from patients hospitalized with suspected pneumonia: frequency of occur- rences and antimicrobial susceptibility patterns from the SENTRY antimicrobial surveillance program. Diagn Microbiol Infect Dis37(2):115–125