A 14-Year Antibiogram Pattern of Isolated Organisms from Pressure Ulcers in Stroke Patients

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INTRODUCTION

Stroke is often complicated by pressure ulcers that are infected by microorganisms. The indiscriminate use of antibiotics, oral or parenteral for proper or improper uses has led to the emergence and spread of resistant micro-organism. Resistance of isolated microorganism to antibiotics is currently a major medical and public health problem. There is paucity of information on the antibiogram of isolated microorganism from pressure ulcer of stroke patient in Nigeria. We aimed to study the antibiotic resistance pattern of isolated organism from the pressure ulcer of stroke patients.

MATERIALS AND METHODS

This was a retrospective record review involving 679 clinical isolates from pressure ulcers of stroke patient admitted into the medical and stroke ward at University of Benin Teaching Hospital Benin City from June 2004 to June 2018.

Data extracted included demographics, stroke types, microbial isolates, and antibiogram. Antimicrobial susceptibility screening was performed using fifteen commonly used antibiotics, by the Kirby Bauer disk diffusion technique. Statistical analysis was done using the proportion of sensitive, intermediate and resistant isolates.

Antimicrobial susceptibility screening was performed using fifteen antimicrobials including, Co-

amoxiclav, gentamicin, Ceftriaxone, Chloramphenicol, Ceftazidime

Others are Ciprofloxacin, Cefuroxime, Tetracycline, Levofloxacin, Imipenem, meropenem, vancomycin, ampicillin, etapenem, nystatin. The zone size around each antimicrobial disk was interpreted as sensitive, intermediate or resistant according to the CLSI criteria.

RESULTS

The mean age of the patients was 64.2(10.8) years, with the age range from 37 to 97 years. Seventy-eight percent had ischemic stroke while 22% had hemorrhagic stroke. About fifty-four percent (363) of the sampled ulcers were from female participants. The most common pathogens isolated were Proteus mirabili (33%), Enterococcus species (24.2%) and Staphylococci species (12%). All 679 isolates were multidrug resistant. The mean microbial resistance to antibiotics during the period under review was 33.9%. In the first seven years the microbial resistance to antibiotics ranged from 25.2% to 31.1%, while the last seven years the resistance range was 34.2% to 43.9%. A high level of resistance was observed for ciprofloxacin (62%), co-amoxiclav (48.8%) and ceftriaxone (22%) while emerging resistance for imipenem (17.5%) and meropenem (20.7%) was also observed. The isolates showed the highest sensitivities for gentamicin (64.5%) and levofloxacin (77.7%).

Year	Male	Female	Total	
	N	Ν	Ν	
2004	10	9	19	
2005	21	20	41	
2006	13	23	36	
2007	14	34	48	
2008	17	22	39	
2009	23	25	48	
2010	28	21	49	
2011	19	22	41	
2012	18	20	38	
2013	22	30	52	
2014	31	20	51	
2015	20	41	61	
2016	29	35	64	
2017	31	22	53	
2018	20	19	39	

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Table 1: The isolated	micro-organism	Der vear Daseu	on dender

Table 2: Frequency and percentage resistance to the tested antibiotics

Antibiotics	Total Isolates	Sensitivity	Resistance	
	Ν	n (%)	n (%)	
AMC	682	134 (19.6)	335 (48.8)	
CN	684	442 (64.5)	56 (8.2)	
CRO	679	330 (48.6)	146 (22.0)	
CIP	615	108 (17.6)	379 (62.0)	
LEV	582	452 (77.7)	54 (9.3)	
IPM	446	298 (66.8)	78 (17.5)	
MER	217	135 (61.8)	45 (20.7)	

AMC = Co-amoxiclav, CN = Gentamicin, CRO = Ceftriaxone, CIP = Ciprofloxacin, LEV = Levofloxacin, IPM = Imipenem, MER= Meropenem

	AM				N	incy o	CR		rearr		C	5		CAZ	
	S	I	R	S	I	R	S	I	R	S	I	R	S	Ι	R
2004	11	7	1	9	3	2	12	5	2	1	10	6	7	8	4
2005	18	21	2	38	1	2	31	6	4	9	7	21	25	11	5
2006	11	20	5	30	4	2	19	12	5	-	-	-	14	13	9
2007	8	21	19	41	5	2	33	8	7	-	-	-	13	11	5
2008	9	17	13	34	9	6	23	7	9	-	-	-	8	26	5
2009	11	17	20	39	6	3	34	10	4	-	-	-	4	37	7
2010	8	13	31	35	7	7	28	11	10	-	-	-	6	34	9
2011	7	5	29	29	8	4	31	6	5	-	-	-	4	26	11
2012	3	7	28	27	9	2	29	3	6	-	-	-	7	7	24
2013	2	18	32	27	19	6	26	18	8	-	-	-	3	12	37
2014	4	16	31	26	23	2	17	6	28	-	-	-	7	37	7
2015	7	26	28	29	30	2	15	29	17	-	-	-	9	47	8
2016	14	12	38	28	27	9	12	45	8	-	-	-	5	8	51
2017	8	7	38	30	21	2	9	31	13	-	-	-	7	14	32
2018	13	9	17	19	15	5	11	6	20	-	-	-	2	10	27

Table 3a: Sensitivity and Resistance Frequency of Antibiotics Per Year in 679 Isolates

Key: I = Sensitivity, R = Resistance, I = Intermediate resistance, AMC = Co-amoxiclav, CN = gentamicin, CRO = Ceftriaxone, C = Chloramphenicol, CAZ = Ceftazidime

		CIP		C	XM			ΤE			LEV			IPM	
	S	Ι	R	S	Ι	R	S	Ι	R	S	Ι	R	S	Ι	R
2004	2	-	6	6	3	5	-	1	13	-	-	-	-	-	-
2005	19	13	9	17	12	7	-	-	25	-	-	-	-	-	-
2006	11	15	10	13	9	14	-	3	15	-	-	-	-	-	-
2007	6	12	30	21	11	16	-	-	-	38	9	1	-	-	-
2008	4	8	27	10	8	21	-	-	-	31	5	3	-	-	-
2009	9	11	28	11	7	30	-	-	-	37	8	3	-	-	-
2010	14	9	26	14	9	26	-	-	-	36	9	4	41	7	1
2011	9	3	29	13	7	21	-	-	-	35	4	2	37	3	1
2012	5	11	22	11	8	19	-	-	-	33	3	2	36	1	1
2013	4	10	38	10	20	22	-	-	-	47	2	3	48	3	1
2014	6	8	37	7	15	24	-	-	-	43	5	3	49	3	1
2015	4	12	41	8	-	-	-	-	-	46	9	6	41	10	10
2016	8	11	36	13	17	34	-	-	-	39	15	10	21	30	13
2017	7	6	40	13	10	9	-	-	-	40	2	11	16	4	31
2018	-	-	-	15	-	8	-	-	-	27	5	6	9	10	20

Table 3b: Sensitivity and Resistance Frequency of Antibiotics Per Year Amongst 679 Isolates

Key: I = Sensitivity, R = Resistance, I = Intermediate resistance, CIP = Ciprofloxacin, CXM = Cefuroxime, TE = Tetracycline, LEV = Levofloxacin, IMP = Imipenem

Table 4: A 14 -	ear Antibiotics Percentage Resistance Pattern Per Year	r

Year	Total Isolates	Resistance (%)		
2004	126	39 (30%)		
2005	298	75 (25.2%)		
2006	234	60 (25.6%)		
2007	317	80 (25.2%)		
2008	283	84 (29.3%)		
2009	336	95 (28.3%)		
2010	395	114 (28.9%)		
2011	328	102 (31.1%)		
2012	304	104 (34.2%)		
2013	416	147 (35.3%)		
2014	403	132 (32.8%)		
2015	474	112 (23.6%)		
2016	503 199 (39.6%)			
2017	401 176 (43.9%)			
2018	249	103 (41.4%)		

Isolated organisms	Male	Female	Total
	(%)	(%)	(%)
Proteus mirabilis	14	19	33
Enterococci species	7	17	24
Staphylococci species	5.8	5.4	11.2
Citrobacter freundii	2.8	6.8	9.0
Enterobacter aerogenes	3.8	3.2	7.0
Eschericha coli	1.2	4.9	6.1
Klebsiella pneumonia	2.3	2.4	4.7
Klebsiella oxytoca	0.7	0.4	1.1
Candida albicans	0.7	0.9	1.6
Mixed bacteria growth	1.1	1.2	2.3

Table 5: Isolated organisms from the pressure ulcers of the 679 stroke patients

DISCUSSION

Antimicrobial resistance is a major medical and public health problem. Gram positive and Gramnegative bacteria continuously develop resistance to antibiotics that are in regular use at different time periods. The main reason for development and spread of bacterial resistance is injudicious use of antimicrobial agents. Ciprofloxacin had the highest microbial resistance with similar findings in other studies. Co-amoxiclav is the next highest with microbial resistance from this work and it has similarly been observed in other studies. The efficacy of a new drug may change as it is used in clinical settings. And resistant bacterial clones result from the encounter of drug and organism. Newly introduced antimicrobial agents imepenem and meropenem with the levels of resistance observed are frightening with similar patterns observed elsewhere. Gentamicin and levofloxacin had the least resistance and this is explained by the very limited use of gentamicin and levofloxacin in clinical practice

CONCLUSION

Local studies meant to understand the isolated organisms and resistance pattern, may guide the clinician on the choice of appropriate empirical therapy.

It is hoped that rational use of antimicrobial agents and constant surveillance for emerging microbial resistance would help control the spread of bacterial resistance

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Conflict of interest

The authors declare that they have no conflicts of interest.

REFERENCES

- Bhattacharya S (2011). Is screening patients for antibiotic-resistant bacteria justified in the Indian context? Indian J. Med Microbiol; 29(3):213-217.
- Campbell C, Parish LC (2010). The decubitus ulcer: facts and controversies. Clin Dermatol; 28(5):527-532.
- Diekema DJ, Boots Miller BJ, Vaughn TE, et al (2004). Antimicrobial resistance trends and outbreak frequency in United States hospitals. Clin Infect Dis; 38(1):78-85.
- English BK, Gaur AH (2010). The use and abuse of antibiotics and the development of antibiotic resistance. Adv Exp Med Biol; 659:73-82.
- Gupta K (2003). Addressing antibiotic resistance. Dis Mon; 49(2):99-110.
- Ingeman A, Andersen G, Hundborg HH, Svendsen ML, Johnsen SP (2011). Processes of care and medical complications in patients with stroke. Stroke; 42:167-172.
- Krasner D (1999). The AHCPR pressure ulcer infection control recommendations revisited. Ostomy Wound Manage; 45:88 91

- Kwan J, Hand P (2007). Infection after acute stroke is associated with poor short-term outcome. Acta Neurol Scand; 115:331-338.
- Mylotte JM, Kahler L, Graham R, Young L, Goodnough S (2000). Prospective surveillance for antibiotic-resistant organisms in patients with spinal cord injury admitted to an acute rehabilitation unit. Am J Infect Control; 28(4):291-297.
- O'Meara SM, Cullum NA, Majid M, Sheldon TA (2001). Systematic review of antimicrobial agents used for chronic wounds. Br J Surg; 88:4-21.
- Rolain JM, Canton R, Cornaglia G (2012). Emergence of antibiotic resistance: need for a new paradigm. Clin Microbiol Infect; 18(7):615-616.
- Subbalaxmi MV, Lakshmi V, Lavanya V (2010). Antibiotic resistance –experience in a tertiary care hospital in South India. J. Assoc. Physicians. India; 58 Suppl:18-22.

- Tong X, Kuklina EV, Gillespie C, George MG (2010). Medical complications among hospitalizations for ischemic stroke in the United States from 1998 to 2007. Stroke; 41:980-986.
- Vandepitte J, El-Nageh MM, Tikhomirov E, Stelling JM, Estrela A (1996). Guidelines for Antimicrobial Resistance Surveillance. Alexandria, Egypt: WHO Regional Office for the Eastern Mediterranean. WHO Regional Publications, Eastern Mediterranean Series; vol 15.
- Westendorp WF, Vermeij JD, Vermeij F, et al (2012). Antibiotic therapy for preventing infections in patients with acute stroke. Cochrane Database Syst Rev; (1):CD008530
- Wright GD, Poinar H (2012). Antibiotic resistance is ancient: implications for drug discovery. Trends Microbiol; 20(4):157-159.