

MULTICENTER STUDY OF THE BURDEN OF MULTIDRUG-RESISTANT BACTERIA IN THE ETIOLOGY OF INFECTED DIABETIC FOOT ULCERS

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Abstract

Background: Infected diabetic foot ulcer (IDFU) is a public health issue and a leading cause of non-traumatic limb amputation. Very few published data on IDFU is available in most West African countries. The objective of this study was to investigate the etiological agents of IDFU and the challenge of antibacterial drug resistance in the management of infections.

Methods: This was a prospective cross-sectional hospital-based study involving three tertiary healthcare facilities. Consecutive eligible patients presenting in the facilities were recruited. Tissue biopsies and/or aspirates were collected and cultured on a set of selective and non-selective media and incubated in appropriate atmospheric conditions for 24 to 72 hours. Isolates were identified by established standard methods. Antibiotic susceptibility testing was performed using modified Kirby-Bauer disc diffusion method. Specific resistance determinants were investigated by polymerase chain reaction-based protocols. Data analysis was done with SPSS version 20.

Results: Ninety patients with clinical diagnosis of DFI were studied between July 2016 and April 2017. A total of 218 microorganisms were isolated, comprising 129 (59.2%) Gram-negative bacilli (GNB), 59 (27.1%) Gram-positive cocci (GPC) and 29 (13.2%) anaerobic bacteria. The top five facultative/aerobic bacteria encountered were: *Staphylococcus aureus*, *Escherichia coli*, *Pseudomonas aeruginosa*, *Klebsiella pneumoniae* and *Citrobacter* spp. representing 41 (18.8%), 23 (10.5%), 20 (9.2%), 19 (8.7%) and 19 (8.7%) isolates, in that order, respectively. The commonest anaerobes were *Bacteroides* spp., and *Peptostreptococcus anaerobius* which accounted for 7 (24.1%) and 6 (20.7%), respectively. Of the 93 IDFU cases, 74 (80%) were infected by multidrug-resistant (MDR) bacteria predominantly methicillin-resistant *S. aureus*, extended-spectrum β -lactamase-producing GNB, mainly of the CTX-M variety. Only 4 (3.1%) GNB were carbapenemase-producers

encoded by *bla*_{VIM}. Factors associated with presence of MDR bacteria were peripheral neuropathy ($r= 4.05$, $P= 0.042$) and duration of foot infection >1 month($r= 7.63$, $P= 0.015$).

Conclusions: MDR facultative/aerobic bacteria are overrepresented amongst agents causing IDFU. A relatively low proportion of the etiological agents were anaerobic bacteria. This finding should help formulate empirical therapeutic options for managing IDFU. Furthermore, drastic reduction in inappropriate use of cocktail of antibiotics for IDFUs is advocated to combat infection by MDR bacteria in these patients.

Keywords: Infection, diabetic foot, ulcers, multidrug resistance.

Introduction:

Infected diabetic foot ulcer (IDFU) is associated with inflammation or purulence occurring in a site below the ankle in a person with diabetes mellitus (DM).¹ It is a major global public health with substantial medical, socio-economic and psychological burden. IDFU is one of the most common diabetes-related infections in clinical practice, and a common indication for hospital admission.² At 7.2% (95%CI: 5.1-9.3%) and higher than the global prevalence of 6.3% (95%CI: 5.4-7.3%), Africa has the second highest global prevalence of diabetic foot ulcer, a precursor of IDFU.³ Paradoxically, foot infections are most common and lethal in Africa than elsewhere globally.⁴ Between 25-60% of diabetic patients with background foot ulcer will develop IDFU which remains a major reason for non-traumatic amputation of the lower limbs.⁵

Wide varieties of organisms, including anaerobic bacteria, have been implicated in the etiology of IDFU depending on severity of infection and time from onset to presentation at the healthcare facility. Advanced IDFUs with features of sepsis at admission usually harbor anaerobic pathogens.² Emergence and current global threat of antimicrobial resistance in the face of dwindling antibiotics in the pipeline has added a new twist to the burden of IDFU.⁶ Increasing involvement of multi-drug resistant organisms in diabetic patients with infected foot ulcers has significantly reduced antibiotic treatment options, thus posing a serious challenge in resource-constrained low- and middle-income countries where access to antimicrobial drugs is of grave concern.⁷ A recent systematic review and meta-analysis on the global burden of diabetic foot ulceration in Cameroon, a West African country, has concluded that paucity of data is a bane on the strategy for treatment and prevention of foot infections in diabetic patients.³ Thus, our study was designed to determine the prevalent bacteria involved in IDFUs, assess the burden of MDR bacteria among the isolates and evaluate the associated risk factors.

Materials and Methods:

Patients:

This prospective cross-sectional hospital-based multicenter study was carried out at the Obafemi Awolowo University Teaching Hospitals Complex, Ile-Ife (Ife Hospital Unit, Wesley Guild Hospital, Ilesa and Ladoke Akintola University of Technology Teaching Hospital, Osogbo between July 2016 and April 2017. All consecutive diabetic patients with foot infections meeting the criteria for the diagnosis of IDFU, seen and managed at these hospitals, were recruited into the study. They were clinically assessed and foot lesions graded according to diabetic foot infection severity classification system issued by the Infectious Disease Society of America.² Only non-duplicate patients and samples were studied. Ethical approvals for this study were granted by the Ethics and Research Committees of the hospitals (protocol numbers: ERC/2015/11/02 and LTH/ER/2016/01/254). Relevant biodata were recorded for each patient.

Samples collection and bacterial identification:

Aspirates were obtained from deep-seated abscesses while tissue samples were collected after washing the wound vigorously with sterile saline and debridement of the slough to exclude mere colonizers. Necrotic tissues were curetted into Anaerobic Basal Broth (Oxoid, Basingstoke, Hants, UK) for anaerobic culture. The samples were immediately transported to the laboratories and processed within 2 hours of sample collection by inoculating them onto a set of selective and non-selective media which were: 5% (v/v) sheep blood agar (BA: Oxoid, Basingstoke, Hants, UK), MacConkey agar (Oxoid, Basingstoke, Hants, UK), chocolate agar and Anaerobic Basal Agar (Oxoid, Basingstoke, Hants, UK) supplemented with 5% (v/v) laked sheep blood, Vit K1 (1 µg/ml), L-cysteine hydrochloride (5 µg/ml) and gentamicin (100 µg/ml) (GBA).

A set of inoculated plain Blood agar and MacConkey agar were incubated in air at 37°C for 24 h, chocolate agar in CO₂ at 37°C for 24 h. Another set of inoculated plain GBA, as well as GBA with kanamycin (75 µg/l) and vancomycin (5 µg/l) supplements, was incubated under atmospheric condition made up of 80% H₂, 10% CO₂, 10% N₂ for 48 h and extended for 5 days if necessary; anaerobiosis was achieved using a Bactron Anaerobic Chamber (SHEL LAB, Cornelius, USA). Representative colonies were identified by colonial morphology, Gram staining characteristics and conventional biochemical tests using Microbact™ GNB 24E (Oxoid, Basingstoke, Hants, UK) and RapID™ STR (Remel, Lexena, USA) for facultative/aerobic Gram-negative bacilli and *Streptococcus* spp. respectively. *Staphylococcus* spp. was identified with catalase and coagulase tests. The obligate anaerobes were identified by RapID™ ANA II (Remel, Lexena, USA). Quality control strains, *Staphylococcus aureus* ATCC 25923, *Escherichia coli* ATCC 25922, *Bacteroides fragilis* ATCC 25285 and *Peptostreptococcus anaerobius* ATCC 27337, were used to assess the quality of the media and identification systems.

Antibiotic susceptibility test:

Antibiotic susceptibility testing for aerobic and facultative anaerobes was performed by modified Kirby-Bauer disk diffusion technique as recommended by Clinical and Laboratory Standards Institute (CLSI)⁸ using *Escherichia coli* ATCC 25922 and *S. aureus* ATCC 25923 as control strains. Discrete colonies were emulsified in sterile saline to match 0.5 McFaland turbidity standard from where confluence inocula were made on Mueller-Hinton agar (MHA) with sterile cotton swab. The swabbed MHA were allowed to dry at room temperature and a set of six antibiotics discs were placed evenly on each of them. After 18 - 24 hours of incubation, the diameter of the zone of inhibition around each antibiotic disc was measured, recorded and interpreted as “sensitive”, “intermediate” or “resistant” in accordance with CLSI guidelines.⁸ Isolates with intermediate sensitivity were regarded as “resistant”.

Extended-spectrum β -lactamase production was determined among Enterobacteriaceae and other GNB that have shown reduced susceptibility to at least one third generation cephalosporin or aztreonam by combination disc method according to CLSI guidelines.⁸ Gram-negative bacilli with intermediate sensitivity or resistance to one or more carbapenems were tested for production of carbapenemases by the Modified Hodge test (MHT) and interpreted by CLSI guidelines.⁸ Methicillin-resistant *S. aureus* (MRSA) was detected by disc diffusion test, using cefoxitin disc (30 μ g) on Mueller-Hinton agar according to CLSI guidelines.⁸

Organisms that were phenotypically multidrug-resistant (MDR), including ESBL-producing GNB, carbapenem-resistant GNB and MRSA, were further tested for resistance determining genes using PCR-based protocols with specific oligonucleotide primers (Table 1) and template DNA of the bacteria extracted by boiling method.⁹ Electrophoresis of each PCR product (5 μ l) was carried out in 1.5% (w/v) Agarose gel (Biomatik, Ontario, Canada) in 1X Tris-Acetate-EDTA (TAE) buffer for 45 minutes. The size of amplified products was estimated using 100bp molecular weight marker (100 – 1200bp).

Statistical analysis:

Data analysis was performed with Statistical Package for Social Sciences (SPSS) version 20. Comparison of mean values was done using the Student's t test for continuous and chi-square test for categorical variables. Risk factors for infection of diabetic foot by MDR organisms among were identified by logistic regression analysis. A p-value of 0.05 was considered to be statistically significant.

Table 1: Oligonucleotides Primers and Amplification Reactions for Targeted Resistance Genes

Target gene	Name	Primer Sequence	Amplicon Size	Amplification reactions	References
<i>bla_{CTX-M}</i>	CTX-M-F	TTGCGATGTGCAGTACCAGTAA	754bp	Initial denaturation at 94°C for 3	10,11
	CTX-M-R	CGAATATCGTTGGTGGTGCCATA		mins, followed by 35 cycle of	
<i>bla_{SHV}</i>	SHV-F	ATTTGTGCGCTTCTTTACTCGC	294bp	denaturation at 94°C for 45 secs,	
	SHV-R	TTTATGGCGTTACCTTTGACC		annealing at 60°C for 30 secs and	
<i>bla_{TEM}</i>	TEM-F	ATGAGTATTCAACATTTCCGTG	404bp	extension at 72°C for 1 min, and	
	TEM-R	TTACCAATGCTTAATCAGTGAG		a final extension at 72°C for 3 mins	
<i>bla_{KPC}</i>	KPC-F	ATGTCACTGTATCGCCGTCT	893bp	Initial denaturation at 95°C for 15	12,13
	KPC-R	TTTTTCAGAGCCTTACTGCCC		minutes, followed by 35 cycles of dena-	
<i>bla_{NDM}</i>	NDM-F	GACAACGCATTGGCATAAG	447bp	turation at 94°C for 30 secs, annealing at	
	NDM-R	AAAGGAAAACCTTGATGGAATTG		60°C for 30 secs, extension at 72°C for	
<i>bla_{VIM}</i>	VIM-F	ATTCCGGTCGGMGAGGTCCG	633bp	10 mins and final extension at 72°C for	
	VIM-R	GAGCAAGTCTAGACCGCCCG		10 mins	
<i>mecA</i>	<i>mecA</i> -F	ATCGATGGTAAAGGTTGGC	530bp	Initial denaturation at 94°C for 4 mins for	14
	<i>mecA</i> -R	AGTTCTGCAGTACCGGATTTCG		followed by 30 cycles of denaturation	
				at 94°C for 30 secs, annealing at 53°C for 30	
				secs and extension at 72°C for 1 min	

Results:

Ninety patients (53 males and 37 females) presented with 93 IDFUs during the 11-month study period. The patients ranged between 18 and 85 years (mean, 54.7 ± 12.8 years) of age. Of the 93 cases of foot infections, 70 (75.3%) were hospitalized patients and 56 (60%) had lasted for a duration of at least one month. Sixty-six (70.1%) of the ulcers were categorized as at least Wagner's grade 3. The vast majority ($n=74$; 82.2%) of the patients used antibiotics in the month before presentation at the facilities and over 90% ($n=84$) had taken antibiotics before wound samples were obtained (Table 2).

Results further showed a total of 218 organisms were isolated from the 93 specimens examined with an average of 2.34 organisms per sample. Of the organisms, 129 (59.2%) were Gram-negative aerobic bacilli, 59 (27.1%) Gram-positive aerobic cocci and 29 (13.2%) anaerobic bacteria; only one (0.5%) was yeast. *S. aureus*, 41 (18.8%) was the single most common organism followed by *E. coli*, 23 (10.6%) and *Pseudomonas aeruginosa*, 20 (9.2%). Others included *Klebsiella* spp. (19; 8.7%), *Citrobacter* spp. (19; 8.7%), *Enterococcus* spp. (13; 6%), *Enterobacter* spp. (11; 5.1%), *Proteus mirabilis* (10; 4.6%) and *Acinetobacter* spp. (9; 4.1%). On the other hand, the predominant anaerobic bacteria were *Bacteroides* spp. (7; 3.2%) and *Peptostreptococcus anaerobius* (6; 2.3%) as shown in Table 3.

Gram-positive bacteria were highly resistant to co-trimoxazole (69.5%), penicillin G (66.1%) and gentamicin (40.1%) but minimally resistant to piperacillin/tazobactam (6.8%) and amikacin (10.2%). On the other hand, Gram-negative bacteria were highly resistant to the tested third generation cephalosporins which included ceftriaxone (56%), cefotaxime (55%) and ceftazidime (48.1%); cotrimoxazole (89%), gentamicin (54.3%) and ciprofloxacin (54.3%). Low rates of

resistance were found to ertapenem (6.4%), piperacillin/tazobactam (9.3%) and amikacin (12.4%).

Analysis of antibacterial resistance profiles of the organisms showed that of the 188 aerobic isolates, 121 (64.4%) were multidrug-resistant (MDR), being resistant to one or more agents in at least three antibiotic classes (Table 4). Further analysis of specific MDR phenotypes showed that 13 (31.7%) of *S. aureus* were methicillin-resistant (MRSA), while 43 (33.3%) and 10 (7.8%) of Gram-negative bacteria were ESBL-producing and carbapenem-resistant respectively (Table 5). Ten (76.9%) of the methicillin-resistant *S. aureus* isolates harbored *mecA* gene and 37 (86.1%) of the 43 ESBL-producing phenotypes harbored at least one of the ESBL-determining genes sought with the most common being *bla*_{CTX-M} (30; 81.1%). Others were *bla*_{TEM} (27; 72.8%) and *bla*_{SHV} (8; 21.6%). Multiple genes were detected in 72.1% of ESBL-producing GNB (n= 31). Carbapenemase determinants were detected in four (40%) of the 10 carbapenem-resistant organisms; they were *bla*_{VIM} (3; 30%), *bla*_{KPC} (2; 20%) and *bla*_{NDM} (1; 10%).

Significant factors associated with presence of MDR organisms in diabetic foot infections included peripheral sensory neuropathy, foot infection duration >1 month and admission duration >1 month (Table 2). Further analysis with logistic regression however identified only peripheral neuropathy (r= 4.05, p= 0.042) and foot infection duration >1 month (r= 7.63, p= 0.015) as the predisposing factors for acquisition of multidrug-resistant bacteria among patients with diabetic foot infection.

Table 2: Association between Clinical/Socio-Demographic Variables of IDFU and MDR Bacteria (N= 90 unless otherwise stated)

Variables		Multi-drug Resistant (MDR) Bacteria		χ^2	P value
		Present	Absent		
Type of Patients	In-patient	58 (82.9)	12 (17.1)	2.978	0.084
	Out-patient	13 (65.0)	7 (35.0)		
Age group (yrs)	<30	4 (80.0)	1 (20.0)	1.101	0.692
	30-49	16 (84.2)	3 (15.8)		
	50-69	44 (78.9)	13 (21.1)		
	70-79	5 (71.4)	2 (28.6)		
	>80	2 (100.0)	0 (0.0)		
Gender	Male	41 (77.4)	12 (22.6)	0.181	0.670
	Female	30 (81.1)	7 (18.9)		
Level of Education	None	14 (82.4)	3 (17.6)	0.292	0.589
	Primary	13 (72.2)	5 (27.8)		
	Secondary	25 (78.1)	7 (21.9)		
	Tertiary	19 (82.6)	4 (17.4)		
Peripheral Sensory Neuropathy (N=93)	Yes	68 (82.9)	14 (17.1)	4.806	0.028
	No	6 (54.5)	5 (45.5)		
Peripheral Vascular Disease (N=93)	Yes	47 (79.7)	12 (20.3)	0.001	0.977
	No	27 (79.4)	7 (20.6)		
Evidence of systemic infection	Yes	37 (82.2)	8 (17.8)	0.600	0.438
	No	34 (75.6)	11 (24.4)		
Ulcer Grade (N=93)	II	21 (77.8)	6 (22.2)	3.916	0.271
	III	32 (82.1)	7 (17.9)		
	IV	18 (85.7)	3 (14.3)		
	V	3 (50.0)	3 (50.0)		
Foot Infection Duration (N=93)	≤1month	35 (94.6)	2 (5.4)	8.533	0.003
	>1month	39 (69.6)	17 (30.4)		
Previous Admission for the Same Ulcer	Yes	14 (82.4)	3 (17.6)	0.151	0.698
	No	57 (78.1)	16 (21.9)		
Glycated Haemoglobin at Presentation (N= 51)	Poor	23 (76.7)	7 (23.3)	0.275	0.872
	Fair	11 (78.6)	3 (21.4)		
	Good	6 (85.7)	1 (14.3)		
Admission Duration (N= 70)	>1month	30 (73.2)	11 (26.8)	6.537	0.011
	≤1month	28 (96.6)	1 (3.4)		
Antibiotic Use in the Last 1 Month before Presentation	Yes	59 (79.7)	15 (20.3)	0.177	0.674
	No	12 (75.0)	4 (25.0)		
Antibiotic Use Before Sample Collection	Yes	67 (79.8)	17 (20.2)	0.577	0.448
	No	4 (66.7)	2 (33.3)		

Table 3: Bacterial etiological agents of infected diabetic foot ulcers

	Names of isolates	Proportion within category (%)	Total (N=218) n (%)
GPC (n=59; 27.1%)	<i>S. aureus</i>	69.5	41 (18.8)
	<i>Enterococcus faecalis</i>	15.3	9 (4.1)
	CoNS	5.1	3 (1.4)
	<i>Enterococcus mundtii</i>	3.4	2 (0.9)
	<i>Enterococcus faecium</i>	1.7	1 (0.5)
	<i>Enterococcus avium</i>	1.7	1 (0.5)
	<i>Streptococcus bovis</i>	1.7	1 (0.5)
	<i>Aerococcus</i> specie	1.7	1 (0.5)
GNB (n=129; 59.2%)	<i>Escherichia coli</i>	17.8	23 (10.6)
	<i>Pseudomonas aeruginosa</i>	15.5	20 (9.2)
	<i>Klebsiella</i> species ^a	14.7	19(8.7)
	<i>Citrobacter</i> species ^b	14.7	19 (8.7)
	<i>Enterobacter</i> species ^c	8.5	11 (5.1)
	<i>Proteus mirabilis</i>	7.8	10 (4.6)
	<i>Acinetobacter</i> species ^d	7.0	9(4.1)
	<i>Morganella morganii</i>	5.4	7 (3.2)
	<i>Hafnia alvei</i>	3.9	5 (2.3)
	<i>Providencia</i> species ^e	2.3	3 (1.4)
	<i>Salmonella</i> Arizonae	1.6	2 (0.9)
	<i>Stenotrophomonas maltophilia</i>	0.8	1 (0.5)
Anaerobes (n=29; 13.3%)	<i>Bacteroides</i> species ^f	24.1	7 (3.2)
	<i>Peptostreptococcus anaerobius</i>	20.7	6(2.8)
	<i>Staphylococcus saccharolyticus</i>	13.8	4(1.8)
	<i>Micromonas micros</i>	10.3	3(1.4)
	<i>Prevotella melaninogenicus</i>	10.3	3(1.4)
	<i>Lactobacillus acidophilus</i> *	6.9	2(0.9)
	<i>Streptococcus intermedius</i> *	3.4	1(0.5)
	<i>Fusobacterium varium</i>	3.4	1(0.5)
	<i>Anaerococcus hydrogenalis</i>	3.4	1(0.5)
	<i>Porphyromonas asaccharolyticus</i>	3.4	1(0.5)
Yeast (n=1; 0.5%)	<i>Candida albicans</i>		1(0.5)

^a*Klebsiella* spp. (*K. pneumoniae*-12; *K. oxytoca*- 7), ^b*Citrobacter* species (*C. freundii*- 8; *C. koserii*- 8; *C. sedlakii*- 3), ^c*Enterobacter* species (*E. aerogenes*- 7; *E. cloaca*- 4), ^d*Acinetobacter* species (*A. baumannii*- 6; *A. johnsonii*- 3), ^e*Providencia* spp. (*P. alcalifaciens*- 2; *P. stuartii*- 1), ^f*Bacteroides* spp. (*B. fragilis* - 6; *B. vulgatus*- 1), CoNS- coagulase negative *Staphylococcus*

*Some of the strains are not strict anaerobes.

Table 4: Prevalence of Multidrug-resistant bacterial isolates in IDFU

BACTERIA ISOLATES (N)	MDR (n)	PREVALENCE (%)
Gram positive cocci (59)	33	59.9
<i>Staphylococcus aureus</i> (41)	20	48.8
<i>Enterococcus</i> spp. (13)	11	84.6
CoNS (3)	0	0.0
<i>Streptococcus bovis</i> (1)	1	100.0
<i>Aerococcus</i> spp. (1)	1	100.0
Gram negative bacilli (129)	88	68.2
<i>Escherichia coli</i> (23)	16	69.6
<i>Pseudomonas aeruginosa</i> (20)	10	50.0
<i>Klebsiella</i> species (19)	13	68.4
<i>Citrobacter</i> species (19)	14	73.7
<i>Enterobacter</i> species (11)	9	81.8
<i>Proteus mirabilis</i> (10)	6	60.0
<i>Acinetobacter</i> species (9)	8	88.9
<i>Morganella morganii</i> (7)	4	57.1
<i>Hafnia alvei</i> (5)	4	80.0
<i>Providencia</i> species (3)	2	66.7
<i>Salmonella</i> Arizonae (2)	1	50.0
<i>Stenotrophomonas maltophilia</i> (1)	1	100.0
TOTAL (188)	121	64.4

MDR – multidrug-resistant, CoNS- coagulase negative *Staphylococcus*

Table 5: Types of multidrug-resistant bacteria in IDFU cases

BACTERIA ISOLATES (N=188)	MRSA/MRCoNS n (%)	VRSA/VRE n (%)	ESBL n (%)	AmpC n (%)	Carbapenem Resistance n (%)
GPC (N=59)					
<i>S. aureus</i> (41)	13 (31.7)	0 (0.0)			
<i>Enterococcus</i> spp (13)		0 (0.0)			
<i>Streptococcus bovis</i> (1)					
<i>Aerococcus</i> sp (1)					
CoNS (3)	0 (0.0)				
GNB (N=129)			43 (33.3)	4 (3.1)	10 (7.8)
<i>Escherichia coli</i> (23)			10 (43.5)	2 (8.7)	
<i>Klebsiella</i> spp (19)			10 (52.6)		
<i>Citrobacter</i> spp (19)			10 (52.6)	1 (5.3)	
<i>Enterobacter</i> spp (11)			6 (54.6)		
<i>P. aeruginosa</i> (20)					3 (15.0)
<i>Acinetobacter</i> spp (9)					4 (44.4)*
<i>M. morganii</i> (7)			2 (28.8)	1 (14.3)	1 (14.3)
<i>P. mirabilis</i> (10)			2 (20.0)		
<i>Hafnia alvei</i> (5)			1 (20.0)		2 (40.0)***
<i>S. maltophilia</i> (1)			1 (100.0)		
<i>Providencia</i> spp (3)			1 (33.3)**		

*Specifically *A. baumannii*, **Specifically *P. alcalifaciens*, ***Carbapenemase-producing (Modified Hodge Test Positive),

MRSA – methicillin-resistant *Staphylococcus aureus*, MRCoNS – methicillin-resistant coagulase negative *Staphylococcus*, VRSA – vancomycin-resistant *Staphylococcus aureus*, VRE – vancomycin-resistant *Enterococcus*, ESBL – extended-spectrum β -lactamase, AmpC – AmpC β -lactamase. GPC – Gram-positive cocci, GNB – Gram-negative bacilli

Discussion:

Infected diabetic ulcers continue to be a polymicrobial infection involving aerobic as well as obligate anaerobic organisms. All IDFUs studied in this series have at least an average of two different bacteria implicated in the disease. Gram-negative bacteria including *E. coli*, *Pseudomonas aeruginosa*, *Klebsiella* species and *Enterobacter* species predominate, which reflects the long-standing nature of the infections, itself a consequence of poor health-seeking behavior in this part of this world.² Furthermore, a wide range of anaerobic bacteria primarily *Bacteroides* species and *Peptostreptococcus anaerobius* are important agents of the infections and were isolated from a third of the cases. This suggests infections that are chronic and below the superficial layers of the skin.¹⁵

Antibiotic resistance remains a huge problem among diabetic foot ulcer infections; it worsens prognosis and makes treatment outcomes poor.¹⁶ Seventy four (80%) of the 93 IDFU cases in this series harbour one or more MDR bacteria, largely attributable to inappropriate antibiotics use and unrestricted access to antimicrobial drugs in many low- and middle-income countries.¹⁷ This contrasts with several studies in high-income countries including France with low prevalence of MDR bacteria among patients with IDFU.^{16,18} A wide spectrum of aerobic and facultative anaerobic bacteria are found to be multidrug-resistant in this study, comparable to findings elsewhere in Africa and Asia.^{19,20}

A third of the *S. aureus* isolates were methicillin-resistant (MRSA). Though prevalence of MRSA appears to be rising in Africa, most of the countries have rates lower than 50%.²¹ This study also revealed that *mecA* was detected in 77% of the MRSA and this is similar to the observation of Chaudhry *et al.* who detected the gene in 20 (84%) of the 25 phenotypically

confirmed MRSA isolates.²² That there are MRSA that lack *mecA* gene may be on account of *mecC*, a variant of *mecA* discovered in 2011 as well as other mutations of penicillin-binding proteins as alternate mechanisms of penicillin resistance.²³

Extended-spectrum β -lactamases were produced by 33.3% of all Gram-negative bacilli isolated and all the organisms except two were members of the family *Enterobacteriaceae*. The leading bacterial hosts producing ESBLs were *E. coli*, *Klebsiella* and *Citrobacter* species. Published rates of ESBL-producing bacteria vary widely across countries in Africa; as high as 96% in Mali to 0.3% in South Africa.²⁴ have all been reported. The most prevalent ESBL type was the CTX-M which has been reported as the most predominant variant worldwide.²⁵ In this study, only 10 (7.8%) of the Gram-negative bacteria were resistant to the carbapenems. Carbapenem resistance determining genes were present in *Acinetobacter baumannii*, *Hafnia alvei* and *Morganella morganii*. Carbapenems are drugs of last resort in the treatment of resistant Gram-negative bacilli infections and with variable and increasing rates of resistance being reported.^{26,27}

Independent risk factors for acquisition of MDR bacteria found in our study are peripheral sensory neuropathy and foot infection duration > 1 month. This is similar to reports among IDFU cases from India.^{28,29} Other authors also documented prolonged duration of wound infection as a predictor of infection of diabetic foot ulcers with MDR bacteria.^{30,31} Contrary findings have however been documented from other studies in China, Iran and Portugal.^{18,20,32} Our findings is also discordant with the report of Noor *et al.* which established that ulcer size is a risk factor for infection by multidrug-resistant organisms.³¹ This study also observed a significant association between presence of multidrug-resistant bacteria in IDFU and long duration of hospitalization (>1 months) similar to previously documented reports by another author in Turkey.³³ We did not find any socio-demographic factors that were significantly associated with occurrence of MDR

IDFU in our study as with other reports.²⁸⁻³⁰ However, on the contrary to our finding, Trivedi *et al.* in the United States noted smoking as an independent risk factor for multidrug-resistant foot wound infection.³⁴

Conclusion:

The spectrum of agents causing IDFU is wide and includes numerous species of aerobic and anaerobic bacteria. There is a high prevalence of MDR aerobic bacteria among them which poses a great limitation to effective treatment of cases. Improved health seeking attitude and timely antibiotic coverage for MDR bacteria will therefore mitigate IDFU in our environment.

Abbreviations

IDFU: Infected diabetic foot ulcer

SPSS: Statistical Package for Social Sciences

GNB: Gram-negative bacilli

GPC: Gram-positive cocci

MDR: multidrug resistant

CLSI: Clinical and Laboratory Standards Institute

MHA: Mueller Hinton agar

MRSA: methicillin-resistant *Staphylococcus aureus*

ESBL: extended-spectrum β -lactamase

Declarations

Ethics approval and consent to participate

Ethical approvals for this study were granted by the Ethics and Research Committees of the Obafemi Awolowo University Teaching Hospitals Complex and Ladoke Akintola University College of Technology with protocol numbers ERC/2015/11/02 and LTH/ER/2016/01/254 respectively.

Consent for publication

Not applicable.

Availability of data and material

The datasets used and analysed during the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests.

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Authors' contributions

ATA and AOA conceived and designed the study. BAK and VOR contributed to the design of the study. ATA and AOA conducted laboratory experiments. ATA and AOA analyzed the data. ATA, AOA and VOR wrote the final report. All authors reviewed and approved the final report.

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