Research Article

Catheter-Associated Urinary Tract Infection and Obstinate Biofilm Producers

Govinda Maharjan,1 Priyatam Khadka,2 Gomik Siddhi Shilpakar,1 Ganesh Chapagain,3 and Guna Raj Dhungana1

1Janamaitri Foundation Institute of Health Science (JFIHS), Kathmandu, Nepal
2Department of Microbiology, Tribhuvan University Teaching Hospital, Kathmandu, Nepal
3Department of Pathology, Tribhuvan University Teaching Hospital, Kathmandu, Nepal

Correspondence should be addressed to Priyatam Khadka; khadka.priyatam@gmail.com

Received 23 May 2018; Accepted 29 July 2018; Published 26 August 2018

Academic Editor: Leticia Reyes

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Background. Biofilms, or colonies of uropathogen growing on the surface of indwelling medical devices, can inflict obstinate or recurring infection, thought-provoking antimicrobial therapy. Methods. This prospective analysis included 105 urine samples from catheterized patients receiving intensive care. Ensuing phenotypic identification, antibiotic sensitivity test was performed by modified Kirby–Bauer disc diffusion method following CLSI guidelines; MDR isolates were identified according to the combined guidelines of the European Centre for Disease Prevention and Control (ECDC) and the Centers for Disease Control and Prevention (CDC). Biofilm-forming uropathogens were detected by the tissue culture plate (TCA) method. Results. The predominant uropathogen in catheter-associated UTIs (CAUTIs) was Escherichia coli 57%, followed by Klebsiella pneumoniae 15%, Pseudomonas aeruginosa 12%, Staphylococcus aureus 8%, Enterobacter spp. 3%, Enterococcus faecalis, Acinetobacter spp., and Proteus mirabilis 1.5%, of which 46% isolates were biofilm producers. Prime biofilm producers were Escherichia coli 33%, followed by Klebsiella pneumoniae 30%, Pseudomonas aeruginosa 20%, Staphylococcus aureus 10%, Acinetobacter, and Enterobacter 3.33%. Multidrug resistance associated with biofilm producers were greater than biofilm nonproducers. The Gram-negative biofilm producers found 96.15%, 80.76%, 73.07%, 53.84%, 53.84%, 46.15%, 19.23%, and 11.5% resistant to amoxyclave, ceftazidime, tetracycline, gentamicin, meropenem, nitrofurantoin, amikacin, imipenem, and fosfomycin, respectively. Gram-positive biofilm producers, however, were found 100% resistant to tetracycline, cloxacillin, and amoxyclave: 66.67% resistant to ampicillin while 33.33% resistant to gentamicin, ciprofloxacin, and nitrofurantoin. Conclusion. High antimicrobial resistance was observed in biofilm producers than non-biofilm producers. Of recommended antimicrobial therapies for CAUTIs, ampicillin and amoxicillin-clavulanate were the least active antibiotics, whereas piperacillin/tazobactam and imipenem were found as the most effectual for gram-negative biofilm producer. Likewise, amoxicillin-clavulanate and tetracycline were the least active antibiotics, whereas vancomycin, fosfomycin, piperacillin-tazobactam, and meropenem were found as the most effective antibiotic for Gram-positive biofilm producer. In the limelight, the activity fosfomycin was commendable against both Gram-positive and Gram-negative biofilm producers.

1. Background

Of nearly 40 percentile of all health care-associated infections, urinary tract infections (UTIs) are the foremost cause of infections; out of these, a bulky proportion, 80%, involve catheter-associated urinary tract infections (CAUTIs) [1]. The urinary catheters are routinely used in urology practice; albeit, advances in design and materials used, UTIs persist as the major snags, owing to the contamination of such indwelling devices [2]. Approximately, prior admission, 12 to 16% of adult hospital inpatients have an indwelling urinary catheter, however, known to be associated with high morbidity, high mortality, increased the length of hospital stay, and increased the cost of treatment [1–3].
Furthermore, the catheter-associated biofilm producers, preceding drug resistivity, and their thought-provoking infection control procedures have been reported in aforementioned studies, which raises our concern on CAUTIs and biofilm producers in our settings [4, 5].

Biofilms are the sessile polymicrobial communities attached to the substrate of biotic and abiotic surfaces and are sheathed within a self-produced extracellular polymeric matrix, that is, polysaccharides intercellular adhesin [2, 5, 6]. The extracellular matrix facilitates communications among the cells through biochemical signals—acyl-homoserine lactone in Gram-negative bacteria and oligopeptides in Gram-positive bacteria—in a phenomenon called as quorum sensing [7]. Not only the matrix precludes the pathogen against host defense but also attributes antimicrobial resistance, by subordinating antibiotic penetration, horizontal transmission of plasmid-associated drug-resistant gene, and altered microenvironment [6, 7]. In this standpoint, early detection of biofilm producers is crucial, to reduce the irrational antimicrobial burden preceding antimicrobial resistance in the patient; hence, it would be an auxiliary in controlling device-associated infections in medical centers.

The rationale of our study was to explicate bacterial etiologies, illuminate biofilm-associated resistivity patterns, and to endorse suitable antimicrobial therapy against biofilm producers in CAUTIs.

2. Material and Methods

2.1. Study Design and Setup. The cross-sectional study was conducted at the Department of Microbiology, Janamaitri Foundation Institute of Health Science (JFIHS), Nepal, over a period of six months. The study hospital is a referral center with medical, surgical, gynecological, pediatric, geriatric, and other specialties.

2.2. Inclusion and Exclusion Criteria. The urine sample of all catheterized patients irrespective of gender and age between 12 and 70 years who met the criteria of CAUTI were included in the study. Nevertheless, noncatheterized patients, either nursed in ward or formerly under antimicrobial therapy at least 48 h prior catheter insertion and no more than two types of organism grown from the clinical sample, were considered as contaminated and consequently, excluded from the study.

2.3. Laboratory Methods. CAUTI was defined using a combination of clinical signs and symptoms and laboratory criteria as described by Stamm [2]. A total of 105 urine sample from the catheterized patient, admitted in intensive care units, were processed semiquantitatively by inoculating 0.001 ml of the specimen (by using a calibrated wire loop) onto the Cystine Lactose Electrolyte Deficient (CLED) agar for the isolation and identification of significant uropathogens [8]. Following the inoculation, the plates were incubated for 24 hours at 37°C in an aerobic atmosphere. The growth of a single organism with a count of ≥10^5 colony-forming units (CFU)/ml was considered to represent as CAUTIs and was identified using appropriate routine identification methods including colony morphology, Gram stain, and an in-house set of biochemical tests [8].

2.4. Antimicrobial Susceptibility Testing. The susceptibility of bacterial isolates against recommended antibiotics was tested by the modified Kirby–Bauer disk diffusion method on Mueller Hinton agar (HiMedia, India) following guidelines of Clinical and Laboratory Standards Institute (CLSI), Wayne, USA [9]. Antibiotics that were tested in our study include amoxycillin clavulanan (amc 20/10 μg), ampicillin (amp 10 μg), amikacin (ak 30 μg), ceftazidime (caz 30 μg), ceftazolin (30 μg), cefuroxime (cfm 30 μg), ciprofloxacin (cip 5 μg), claxacillin (cox 30 μg), crotimoxazol (cot 25 μg), fosfomycin (fo 200 μg), gentamicin (gen 10 μg), imipenem (imp 10 μg), meropenem (mrp 10 μg), nitromarurant (300 μg), piperacillin-tazobactam (pit 100/10 μg), teteracycline (te 30 μg), and vancomycin (VAN 30 μg) (HiMedia Laboratories, India). Further, elucidations of antibiotic susceptibility results were made referring to the zone size interpretative standards of CLSI [9]. MDR isolates, resistant to at least one antimicrobial from three different groups of first-line drugs, tested were identified according to the combined guidelines of the European Centre for Disease Prevention and Control (ECDC) and the Centers for Disease Control and Prevention (CDC) [10]. Escherichia coli ATCC 25922, Staphylococcus aureus (ATCC 25923), and Pseudomonas aeruginosa (ATCC 27853) were used as a control organism for antibiotic susceptibility testing.

2.5. Detection of Biofilm Production. The detection of biofilm was done by tissue culture method/microtiter plate method (TCA), the gold standard method, as described as Christensen et al. [11]. In brief, the bacterial isolates from fresh agar plates were inoculated in 2 ml of BHI broth and incubated for 24 h at 37°C. The cultures were then diluted 1:40 with fresh medium (BHI broth supplemented with 1% glucose); 200 μl of the sample was dispensed in the individual microtitration plate (AD Touch, apDianv) and incubated further 24 h at 37°C. With a gentle tapping, the content was removed further with a subsequent washing with phosphate buffer saline (pH 7.2) three times to remove free floating sessile bacteria. The adherent bacteria, biofilm producer, were fixed with sodium acetate (2%) and stained with crystal violet (0.1% w/v) for 10–15 min. The unbound crystal violet solution was removed with a triplicate washing with PBS, and the plate, then, was kept for drying. Finally, all wells were filled with 200 μl ethanol (95%) to release dye from the well and Optical Density (OD) was taken at the wavelength of 630 nm. For a precision, the experiment was performed in triplicate two times. Average OD values of each test strain and negative control were calculated, and OD cut-off values (ODc) were assessed as described by Stepanovie et al. [12].

2.6. Data Analysis. The information regarding patients’ profile and the results were entered into a computer
program. Data analysis was carried out using the Statistical Package for Social Sciences (SPSS™) version 20.0 (IBM, Armonk, NY, USA) and presented in percentage base distribution.

2.7. Ethical Consideration. Written approval was taken from the Institutional Review Committee (IRC) of Janamaitri Foundation Institute of Health Science (JFIHS) after submitting and presenting the research proposal. Written informed consent was taken from every patient or their guardians before enrollment in the study.

3. Results

3.1. Patient Demographics. During the study period, a total of 105 urine specimens from the patients suspected with catheter-associated UTIs were processed. Among total clinical specimens, 61.9% (65/105) were found with a growth of at least one significant pathogen confirming the urinary tract infection (UTI). Female (43, 66.2%) was the most affected group, with predominant etiologies as *Escherichia coli* 56.9% (37/65) followed by *Klebsiella pneumoniae* 10 (15%), *Pseudomonas aeruginosa* 8 (12%), *Staphylococcus aureus* 5 (8%), *Enterobacter* spp. 2 (3%), and *Enterococcus faecalis*, *Acinetobacter* spp., and *Proteus mirabilis* 1 (1.5% each) (Table 1).

3.2. Detection of Biofilm Producers. In the current study, 30 (46%) strains were in vitro positive for the biofilm production and 35 (54%) were negative for the biofilm production. Out of which (n = 7, 9, and 14) were strong, moderate, and weak biofilm producer. *Escherichia coli* (33.33%) was found to be more biofilm producer followed by *Klebsiella pneumoniae* (30%), *Pseudomonas aeruginosa* (20%), *Staphylococcus aureus* (10%), and *Acinetobacter* spp. and *Enterobacter* spp. (3.33% each) (Table 2).

Microtitration plate shows the biofilm production by TCA method as shown in Figure 1.

3.3. The Antimicrobial Resistant Pattern in Biofilm Producers and Nonproducers. Gram-negative biofilm producers, more than 90%, were resistant to ampicillin and amoxicillin-clavulanate, and nearly 94% of biofilm producer and nonproducer were found to be sensitive to fosfomycin. Besides, the greater percentile of antimicrobial resistivity was found to be associated with biofilm producers than biofilm nonproducers. The Gram-positive biofilm producers, 3 of 3 bacterial isolates, were resistance to amoxicillin-clavulanate, tetracyclins, and cloxacillin; nonetheless, the antibiotics—vancomycin, fosfomycin, meropenem, and piperacillin/tazobactam—were found to be effective even for the biofilm producers (Table 3).

The multidrug combination, in Gram-positive isolates and Gram-negative isolates, was found to be resistant to ciprofloxacin/cotrimoxazole/ampicillin (75%) and ampicillin/cloxacillin/teicoplanin (66.67%) (Table 4).

4. Discussion

Most aspects of the diagnosis, treatment, and prevention of CAUTI are influenced by the tenacity of biofilm-associated uropathogens [13]. Meanwhile, in patients with underlying diseases or under intensive care, the relevance of detection of biofilm producers is crucial since CAUTIs is a common nosocomial infection [1]. The prevalence of catheter-associated urinary tract infection was found to be 61.9%, with predominance in female sufferers (43, 66.2%). Similar prevalence was reported in a review of Nicolle from 15 developing countries like ours [14]. This might be due to the anatomical differences of urogenital organs—anal proximity and shorter urethra in female [15].

Our study revealed that the most common bacterial etiology contributing CAUTIs was *Escherichia coli* 57%, followed by *Klebsiella pneumonia* 15%, *Pseudomonas aeruginosa* 12%, *S. aureus* 8%, *Enterobacter* spp. 3%, and *Enterococcus faecalis*, and *Acinetobacter* spp. and *Proteus mirabilis* (1.5% each) which was nearly similar to the findings Seif Elden Salwa et al. who found that *Escherichia coli* (50%) was predominantly acquired infection followed by *Klebsiella* spp. (30%), *Pseudomonas* spp. (8%), *Staphylococcus aureus* (8.6%), and *Proteus* spp. (4.6%) [16]. In a study, Niveditha et al. and Pramodhini reported 70% of *Escherichia coli* were isolated [17, 18]; in contrast, Ahmed Abdallah et al. reported 31% *Escherichia coli* in their study [19]. The recent prevalence study, however, depicted the predominance of the *Escherichia coli* attaining 79.1% leading UTIs [20]. Similarly, 46% of the bacterial isolates were detected as biofilm producers in the CAUTIs, ensuing with the TCA method. Similar findings were reported by Ghanwat et al. and Ahmed Abdallah et al. where the biofilm producers were 47.5% and 43.5%, respectively [19, 21]. Niveditha et al. and Reid et al., however, found the associated biofilm producers 60% and 73%, respectively, which matched with our study [17, 22].

In our study, 46% isolates were biofilm producers using the tissue culture plate method. Similar findings were observed by Safia Maqbool et al. and Ahmed Abdallah et al. where 47.5% and 43.5% uropathogenic isolates were biofilm producers, respectively; Niveditha et al. and Reid et al., however, found 60% and 73% biofilm producers as CAUTIs [17, 19, 22, 23]. The uropathogen, *Escherichia coli* 33%, emerges as a predominant biofilm producer followed by *Klebsiella* 30%, *Pseudomonas* spp. 20%, *S. aureus* 10%, and

| Organism                | Frequency | Percent |
|-------------------------|-----------|---------|
| *Escherichia coli*      | 37        | 56.9    |
| *Klebsiella pneumoniae* | 10        | 15.4    |
| *Pseudomonas aeruginosa*| 8         | 12.3    |
| *Staphylococcus aureus* | 5         | 7.7     |
| *Enterobacter* spp.     | 2         | 3.1     |
| *Enterococcus faecalis* | 1         | 1.5     |
| *Proteus mirabilis*     | 1         | 1.5     |
| *Acinetobacter* spp.    | 1         | 1.5     |
| Total                   | 65        | 100.0   |
Acinetobacter and Enterobacter (3.33% each) in our settings. The findings were comparable with Sharma et al. 67.5% and Nivenditha et al. 60% for the predominant isolate—biofilm producing Escherichia coli [17, 24]. However, Ponnusamy et al. in their study revealed that of 100 Escherichia coli strains, 72 were biofilm-positive phenotype [25]. Conferring a degree of biofilm producers, in our study, 11%, 14%, and 21% were strong, moderate, and weak biofilm producers, whereas Mishra et al. disclosed, 1.5%, 10.5%, and 34.3% as strong, moderate, and weak biofilm producers in their study [4].

Antibiotic resistance patterns of biofilm producer and nonproducer were observed; sequentially, high resistance against tested antibiotics was attributed in biofilm producers. In Gram-negative bacterial isolates, the resistance against four groups of antibiotics, ampicillin, nitrofurantoin, tetracycline, and meropenem, was equated in biofilm producer versus non-biofilm producer isolates; the consecutive antibiotics were 92.3% versus 84.84%, 46.15% versus 15.15%, 73.07% versus 51.51%, and 53.84% versus 36.31%, respectively. Similarly, the antibiotics ampicillin, ciprofloxacin, nitrofurantion, and tetracycline were 66.67% versus 33.33%, 33.33% versus 0%, 33.33% versus 0%, and 100% versus 66.67%, respectively. Pramodhini et al., nonetheless, disclosed ampicillin (83.3% versus 60%), cephoxaxime (73.3% versus 35%), norfloxacin (80% versus 60%), and nalidixic acid (93% versus 70%) from his study [18]. Furthermore, fosfomycin has shown promising in vitro activity against both Gram-positive and Gram-negative biofilm producers, as earlier experienced by Neuner et al., Mihaiescu et al. and Marquès et al. in treating urinary tract infections [26–28].

Hence, the antimicrobial resistivity in the isolate possibly attributed with the biofilm productions. In Gram-positive isolates, the resistance against 3 groups of antibiotics amoxicillin-clavulanate, tetracyclin, and cloxacillin was equated in biofilm producer versus non-biofilm producer isolates; the consecutive antibiotics were 100% resistant in biofilm producers while 33.3%, 66.7%, and 66.77% were respectively.
found to be resistant in non-biofilm producers. Similarly, MRSA, resistant pattern linked with the tenacity of biofilm producers, was observed by Ando et al. [29]. In biofilms, poor antibiotic penetration, nutrient limitation and slow growth, adaptive stress responses, and formation of persister cells are hypothesized to constitute a multilayered defense [30].

The backdrop of antimicrobial resistant in biofilm producers urges seeking of naive therapeutic alternatives despite conventional antibiotic therapies. In the meantime,
to prevent or to combat these obstinate biofilm producers, small molecules (N-acetylcycteine, Ca2+ and Mg2+ chelators), and matrix-targeting enzymes (dispersin B, DNase I, proteinase K and trypsin), bactericidal and antiadhesion coatings (trimethylsilane, carboxybetaine methacrylate, organoselenium, heparin, and hyaluronic acid; polymer brush coatings; and furanones) were commenced successfully in developed nations [31]. Ironically, the indorsed therapeutic solutions are farther than reach in the developing countries like ours.

5. Conclusion
Biofilm producing bacteria are responsible for many recalcitrant infections and are notoriously hard to eradicate, owing to the possible acquisition of multidrug status. High antimicrobial resistance was observed in biofilm producing pathogens than nonproducers. Of recommended antimicrobial therapies for CAUTIs, ampicillin and amoxicillin-clavulanate were the least active antibiotics, whereas piperacillin/tazobactam, fosfomycin, and imipenem were found as the most effective antibiotics among Gram-negative biofilm producers. Likewise, among Gram-positive biofilm producer isolates; amoxicillin-clavulanate and tetracycline were the least active, whereas vancomycin, fosfomycin, piperacillin/tazobactam, and meropenem were found as the most effective antibiotic. In the limelight, the activity of fosfomycin was laudable against both Gram-positive and Gram-negative biofilm producers. Hence, an early identification of biofilm producers with subsequent detection of antibiotic resistivity pattern is mandatory, to improve the clinical management of CAUTIs when the patient requires an intensive care.

Data Availability
The data used to support the findings of this study are available from the corresponding author upon request.

Ethical Approval
Written informed consent was obtained from every patient for granting participation in an interview and to extract pertinent sociodemographic and clinical data from their respective clinical files, respecting confidentiality.

Consent
Written informed consent was obtained from the patients for relevant investigations, and publication of the findings was taken from every patient and coauthors.

Conflicts of Interest
The authors declare that they have no conflicts of interest.

Authors’ Contributions
Govinda Maharjan and Priyatham Khadka conceived the study, designed the manuscript, and reviewed the literature. Ganesh Chapagain, Gomik Siddhi Shilpakar, and Guna Raj Dhungana reviewed the manuscript and gave the concept of the research paper and critically reviewed the manuscript. All authors read the final version of the manuscript and approved for publication.

Acknowledgments
The authors would like to thank Asst. Prof. Shyam Kumar Mishra (Institute of Medicine, TUTH) for his tremendous technical support.

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