

# MULTIDRUG RESISTANT PROFILE OF UROPATHOGENIC *Escherichia coli*(UPEC) ISOLATED FROM DIABETIC PATIENTS IN SOME HOSPITALS OF BAUCHI METROPOLIS, NIGERIA

## ABSTRACT

**Background:** Urinary Tract Infection (UTI) is a common pathogenic inflammatory, distressing and occasionally life-threatening condition that affects people of all ages and gender, with difficulty in treatment due to high rate of antibiotic resistance. Diabetic patients are more prone to urinary tract infection due to their immunocompromised system and hyperglycemia level compared to non-diabetic patients. *Escherichia coli* is the primary cause of UTIs in humans both in diabetic and non-diabetic patients. Antibiotics are becoming less and less effective, therefore there is an urgent need to curtail this problem in order to have good administration of antibiotics to patients for effective treatment.

**Aim:** The aim of this study was to isolate uropathogenic *Escherichia coli* from diabetic patients and determine its antibiotic-resistant pattern

**Methods:** A total of 288 study participants were enrolled in the study, (194 diabetic and 94 non-diabetic patients). Clean catch mid-stream urine samples were collected from all the participants in sterile containers. Each urine sample was streaked onto CLED (cysteine lactose electrolyte deficient agar), incubated at 37°C for 24h and the isolates were identified using standard methods. Data obtained were analyzed statistically.

**Results:** A total of 64 UPEC was isolated from diabetic patients and 35 UPEC was isolated from non-diabetic patients. Age group of 31-40 had high frequency of occurrence in both the study participants, while age group of 10-20 and  $\geq 71$  years had the least. There was no significant difference between age group and the number of isolates as  $p > 0.05$ . Highest frequency of UPEC was found within the female than their male counterparts. Type 2 patients have high frequency of isolates compared to type 1 patients in both the study participants. In the present study, 52 UPEC isolates from diabetic patients and 27 UPEC isolates from non-diabetic patients were resistant to 2 or more antimicrobial agent (multidrug resistance). The highest resistance was observed against ampicillin and piperacillin-tazobactams, while the least resistant in imipenem.

**Conclusion:** The study established that UPEC infection was more prevalent in diabetic than non-diabetic patients, and also more prevalent in the middle age group, female gender and type 2 diabetic patients. High rate of multidrug resistance was observed in both the study participants, and this signals a tremendous problem in prescription of antibiotics to patients. The emergence of multi resistant strains of UPEC has added to the need of urgent development of more control measures and policies to the use of antibiotics.

**Keywords:** UPEC (uropathogenic *E. coli*), diabetic patients, non-diabetic patients, multidrug resistance.

## 1. INTRODUCTION

Urinary Tract Infection (UTI) which is defined as the presence and active multiplication of microorganisms within the urinary tract is one of the commonest bacterial infections seeking treatment in clinical practice [1]. 250 million people globally experience urinary tract infections (UTIs), it is one of the most common diseases in humans with a variety of etiological factors [2]. As stated by [3], the infection is named after the affected urinary organ or part and is known as cystitis (bladder infection) and pyelonephritis (kidney infection). The symptoms of bladder and

kidney infections differ, with cystitis causing painful and frequent urination and pyelonephritis causing high fever and flank pain [3].

Diabetes mellitus (DM) is an ever-growing heterogenic disorder altering the metabolic abilities of the body, primarily characterized by persistently high glucose levels (hyperglycemia) resulting from defects in insulin secretion attenuating every bodily function [4,5]. It is among the most common non-communicable diseases in emerging and developed nations[6], it has a number of effects on the genitourinary system [7]. According to [8], Diabetes Mellitus (DM) has become a significant public health issue worldwide and has emerged as a significant socio-economic burden for developing nations.

UTIs are classified based on the type of infection (upper or lower UTI), the presence or absence of symptoms (symptomatic or asymptomatic), the tendency to recur (single episode or recurrent UTI), and the presence or absence of complicating factors (uncomplicated or complicated UTI) [9,10,11,12,3,2]. Similarly, risk of UTI increases with age, poor metabolic control, various impairments in the immune system and incomplete bladder emptying due to autonomic neuropathy[13].

The common symptoms of UTI include burning micturition, urgency, dysuria, cramping in the lower abdomen, mental irritability, back or flank pain, chill, nausea, fever, vomiting, fatigue, and weakness [14]. According to [15], it is very important to screen diabetic patients for UTIs for timely diagnosis, complete treatment, and prevention of progression to renal complications and ultimately severe renal failure.

Urinary tract infection (UTI) is among the most common medical condition seen in all age groups with DM [13]. Diabetic patients are highly susceptible to UTI compared to non-diabetics[8]. Evidence from various epidemiological studies showed that UTI is more common in females with diabetes than in non-diabetic females[16], it is more common in diabetes because of a combination of host and local risk factors. Modification of chemical composition of urine in diabetes mellitus can alter the ability of urine and support the growth of microorganisms[7].

Recent epidemiological studies and analytical experimentation of patients with preexisting diabetes mellitus have authenticated their plausibility of developing UTIs that are potentially perilous with fatal manifestations [4]. Various impairments in the immune system, including humoral, cellular, and innate immunity may contribute to the pathogenesis of UTI in diabetic patients[5].

According to [3], bacteria are the primary cause of UTIs in humans and the most contributed bacterial pathogen of UTIs is *Escherichia coli* in diabetic and non-diabetic patients and others are *Klebsiellapneumonia*, *Staphylococcussaprophyticus*, *Proteusmirabilis*, *Enterococcusfaecalis*, Group-B *Streptococcus*, *Pseudomonas aeruginosa*, *Candidaspp*, and *Staphylococcus aureus*[14,15,17].

However, among the bacterial species involved in UTIs, uropathogenic *Escherichia coli* strains (UPEC) are the most common. UPEC account for about 80% of uncomplicated UTIs, 95% of community-acquired infections, and 50% of hospital-acquired infections [18]. UPEC also remains the most frequent pathogen in complicated UTIs [19]. According to [2], UPEC is a heterogeneous group of extraintestinal pathogenic *E. coli* (ExPEC) that seem to originate from the gut.

Antimicrobial resistance is naturally occurring as a reaction of microbial organisms to environment[20]. The emergence of multi-drug resistant (MDR) strains is escalating, causing urinary tract infections increasing both in community and hospital settings [1,15,21]. Increasing cases of diabetes mellitus which consequently lead to more UTI cases and irrational use of

antibiotics has led to emergence of multi-drug resistant strains[22]. More so, studies in Africa have shown the need to have systematic screening of UTI in diabetic patients due to the increasing prevalence [22].

There is widespread concern about the high rates of resistance to antimicrobials used in the treatment of urinary tract infections, particularly in developing countries. Antibiotics, including broad-spectrum antibiotics, are frequently prescribed, which may lead to the development of antibiotic-resistant urinary pathogens. Patients with diabetes mellitus are more likely to have resistant pathogens, necessitating longer and more potent antimicrobial treatment. Improved glycemic control in diabetics may thus aid in the control of UTIs. Accurate screening for UTIs in diabetic patients is also critical to enable appropriate treatment and avoiding related complications [23].

Diabetes mellitus (DM) has now become a global health issue to health care professionals [8]. Frequent prescription of antibiotics, including the ones with broad-spectrum, has result in development of antibiotic-resistant urinary pathogens. Since patients with DM are more prone to have resistant pathogens, they inevitably require longer and more potent antimicrobial treatment [23]. Therefore, improved control of glycaemia, timely diagnosis, complete treatment and screening for UTIs in diabetic patients is critical to prevent progression to renal complications and ultimately severe renal failure and other related complications.

## **2. MATERIALS AND METHODS**

### **2.1 Study Area**

The study area was conducted in Bauchi State, Nigeria. The study area for the collection of samples include; Abubakar Tafawa Balewa University Teaching Hospital (ATBUTH), Bauchi State, Bauchi state specialist hospital.

### **2.2 Sample Collection**

Each patient was informed to collect approximately 20 ml of midstream urine into a crew cap sterile calibrated urine container. Proper instructions and illustration were given to the patients in order to avoid contamination. At the point of collection, samples were labeled with name, sex and age of the patients. The samples were placed in an ice box and transported to the laboratory for further analysis, but in cases of delay, the urine samples were refrigerated at 4°C to avoid multiplication of bacteria [24].

### **2.3 Data Collection**

A structured questionnaire and patient clinical sheet was used to sought for demographic data and clinical details (clinical symptoms, previous antibiotic usage, risk factors/behaviours). Verbal/informed consent was obtained from each patient to be enrolled in this study.

### **2.4 Inclusion/ Exclusion Criteria**

The study will include diabetic and non-diabetic patients regardless with presence of UTI symptoms. All other patients without these criteria are excluded.

### **2.5 Sample Size Determination**

Sample size determination is the act of choosing the number of observations or replicates to include in a statistical sample. Sample size determination is the mathematical estimation of the number of subjects/units to be included in a study.

The number of samples that was collected was determined using the formula of [25]. Prevalence of  $p=0.25(25\%)$  [26], based on previous study.

Using the formula

$$n = \frac{(Z)^2 p(1-p)}{d^2}$$

Where;

n = Desired sample size

Z = 1.96 (The standard normal deviate, corresponds to the 95% confidence level).

p = Prevalence of previous study or related (0.25)

d = Degree of accuracy (5%)

Therefore  $n = (1.96)^2 \times 0.25(1-0.25)$

$$\frac{(0.05)^2}{(0.05)^2}$$
$$n = (1.96)^2 \times 0.25 \times (0.75)$$

$$n = 3.8416 \times 0.1875$$

$$\frac{0.0025}{0.0025}$$

$$n = 0.7203$$

$$\frac{0.0025}{0.0025}$$

Sample size (n) =288.1

Approximately n=288

## 2.6 Blood Glucose Test

Plasma glucose (after an overnight fasting of eight or more hours) was determined using the glucose meter Accu-Chek Active system (Roche Diabetes Care, Basel, Switzerland), this was carried out within fractions for each participant. DM was diagnosed according to [27] criteria with symptoms of diabetes plus fasting blood glucose level equal or more than 126 mg/dl.

## 2.7 Sample Processing

### 2.7.1 Inoculation and Isolation of Bacteria from Urine Sample

Samples was observed macroscopically for colour, blood tinge and turbidity. All the urine samples were aseptically inoculated using a sterile wire loop, a loopful of well-mixed uncentrifuged urine was aseptically inoculated unto Cysteine Lactose Electrolyte Deficient (CLED) by streak method as described by [28,29,30]. The plates were incubated at 37°C for 18-24 hours.

### 2.7.2 Identification and Characterization of Isolates

The colonies were further identified based on colonial morphology and biochemical test (Indole test, Methyl Red test, Vogues Proskauer test, Citrate Utilization test, coagulase and catalase tests) as described by [28]. Colonies were observed for morphological features such as size, shape, edge consistency, margin, colour, opacity and effect on media i.e. lactose or non-lactose fermenters. In addition to these morphological features of the colonies, motility, Gram staining reaction, and biochemical tests was used in identifying the isolates. The isolates were maintained on Nutrient agar slants, until required for further use [29].

## 2.8 Antimicrobial susceptibility testing of the UPEC Isolates

The susceptibility pattern of the isolates to commonly used antimicrobial agents were determined using as Kirby-Bauer disc diffusion techniques as described by [31]. A loopful of growth of each isolate on agar medium will be suspended in sterile saline and then was diluted in steps of 1:10 to give turbidity equivalent to the 0.5 McFarland standards (a density of  $1 \times 10^8$  cells/mL) before inoculation. Muller-Hinton agar medium was prepared according to manufacturer's instructions and was poured (about 25 ml of the media) into each of sterile petri-plates, the plates was allowed to solidify. After the adjusting turbidity of the inoculum, a sterile cotton swab stick was dipped into the suspension, pressed firmly against the inside wall of the tube; the swab was streaked over the surface of the solidified Muller-Hinton agar plates 3 times

rotating the plate after each application to ensure an even distribution and allowed to stand at room temperature for 10 minutes [32].

Antibiotic discs of known concentration of antibiotics were aseptically placed using sterile forceps and then gently pressed down on the Muller- Hinton agar plates to ensure contact. The plates were then inverted and incubated at 37°C for 24 hours. The diameter of the zone of inhibition produced by each antibiotic disk was measured and interpreted according to Clinical and Laboratory Standard Institute [33] guidelines.

## 2.9 Data Analysis

The data obtained was recorded in Microsoft excel and analyzed by using Chi-square statistical analysis.

## 3. RESULT AND DISSCUSION

### 3.1 Characteristics of Participants Based on Blood Sugar Level (FBS)

In this study, patients who had  $\geq 126$  mg/dl of fasting blood sugar were considered as positive for diabetes mellitus, while those with fasting blood sugar (FBS)  $< 126$  mg/dl were considered nondiabetics. As shown in table 1.

**Table 1: Distribution of Study Participants Based on Blood Sugar Level (FBS)**

| Blood glucose level (mg/dL) | No (%) of patients Tested (n=288) | Percentage (%) |
|-----------------------------|-----------------------------------|----------------|
| High                        | 194                               | 67.4           |
| Normal                      | 94                                | 32.6           |

**Key:** high=  $\geq 126$ mg/dl, normal=  $< 126$  mg/dl (WHO criteria for diabetic diagnosis), FBS- Fasting Blood sugar.

The amount of glucose in the bloodstream is referred to as blood glucose or blood sugar. Fasting blood glucose test is a simple test, accurate and inexpensive test that can screen for diabetes [27]. Diabetics have an impaired immune system, making them more susceptible to many illnesses, among these is urinary tract infection. Based on several researches, diabetic patients are more likely to develop UTIs than non-diabetic patients. There are numerous reasons that could explain the increased occurrence of UTI in diabetic individuals. Studies indicated that high glucose levels in urine promote the formation of uropathogens [34]. Higher glucose levels in renal parenchyma create a favorable environment for bacterial colonization, resulting in complications including emphysematous pyelonephritis [35]. Increased glucose levels may impair humoral, innate, and cellular immunity. Autonomic neuropathy can cause bladder dysfunction, resulting in urine retention and stasis.

Poor metabolic regulation causes hyperglycemia, which can lead to a decreased renal threshold for glucose reabsorption and glycosuria. Glycosuria provides a rich medium for bacterial species to thrive in the presence of decreased immunity. According to [36], elevated plasma glucose levels cause glucosuria, which promotes bacterial proliferation through an increase in cell count, implying neutrophil dysfunction.

Several researchers used fasting blood sugar to screen their study participants from diabetic mellitus. This follows the suites of [37] who conducted a research on pattern of organism causing urinary tract infection in diabetic and Non diabetic patients in Bangladesh, [38] who conducted their research on Clinical profile of urinary tract infections in diabetics and non-diabetics, [39] who worked on prevalence of urinary tract infections and their antimicrobial sensitivity among diabetic and non-diabetic patients in Zakho, Iraq., [40] who conducted their research in Iraq also used fasting blood sugar to screen their study participants from diabetic mellitus, also [41] who conducted their research on diabetic patients in port Harcourt, Nigeria. All the above researchers used fasting blood sugar to screen their patients.

### 3.2 Sociodemographic Variables of UPEC infection Among Study Participants

In this study, the age range of 31-40 had the highest rate of *E. coli* infection in both the study participants, 28.1% in diabetic and 25.7% in non-diabetic patients. In this study, the female gender in both the study population has a high rate of occurrence compared with male counterparts. In this study, married participants have higher rate of infection in both diabetic and non-diabetic patients. While illiterate have high rate of infection in NDM patients than NDM patients. And also, Urban dwellers have high rate of occurrence of UTI in both DM and NDM patients than rural dwellers. As shown in table 2.

**Table 2: Distribution of UPEC Isolates in Diabetic and Non-Diabetic Patients According to Demographic Variables**

| <b>Patients</b>       | <b>No of Samples</b>       | <b>No (%) of UPEC in DM</b> | <b>No (%) of UPEC in NDM</b> | <b>P-value</b> |
|-----------------------|----------------------------|-----------------------------|------------------------------|----------------|
| <b>Details</b>        | <b>Collected<br/>N=288</b> | <b>Patients<br/>(n=64)</b>  | <b>Patients<br/>(n=35)</b>   |                |
| <b>Age (years)</b>    |                            |                             |                              |                |
| 10-20                 | 10                         | 5(7.8)                      | 2(5.7)                       | 0.99           |
| 21-30                 | 30                         | 11(17.2)                    | 7(20)                        |                |
| 31-40                 | 38                         | 18(28.1)                    | 9(25.7)                      |                |
| 41-50                 | 76                         | 14(21.9)                    | 8(22.9)                      |                |
| 51-60                 | 68                         | 9(12.5)                     | 6(17.1)                      |                |
| 61-70                 | 55                         | 5(7.8)                      | 3(8.6)                       |                |
| ≥71                   | 11                         | 2(3.1)                      | 1(2.9)                       |                |
| <b>Gender</b>         |                            |                             |                              |                |
| Female                | 168                        | 36(56.3)                    | 20(57.1)                     | 0.93           |
| Male                  | 120                        | 28(43.8)                    | 15(42.9)                     |                |
| <b>Marital status</b> |                            |                             |                              |                |
| Single                | 14                         | 10(15.6)                    | 5(14.3)                      | 0.62           |
| Married               | 216                        | 42(65.6)                    | 20(57.1)                     |                |
| Divorce               | 28                         | 8(12.5)                     | 8(22.9)                      |                |
| Widow                 | 30                         | 4(6.3)                      | 2(5.7)                       |                |
| <b>Education</b>      |                            |                             |                              |                |

|                  |     |          |          |      |
|------------------|-----|----------|----------|------|
| Literate         | 189 | 40(62.5) | 19(54.3) | 0.42 |
| Illiterate       | 99  | 24(37.5) | 16(45.7) |      |
| <b>Residence</b> |     |          |          |      |
| Rural            | 70  | 21(32.8) | 11(31.4) | 0.88 |
| Urban            | 218 | 43(67.2) | 24(68.6) |      |

Key: UPEC= uropathogenic *E. coli*, DM= Diabetic Mellitus, NDM= Non-Diabetic Mellitus

The age distribution of patients with UPEC isolates was examined across different age groups. Among patients aged 10-20 years, 7.8% of diabetic patients and 5.7% of non-diabetic patients were infected with UPEC. In the 21-30 age group, the prevalence was 17.2% for diabetics and 20% for non-diabetics. For the 31-40 age group, UPEC was found in 28.1% of diabetic patients and 25.7% of non-diabetic patients. In the 41-50 age range, the prevalence was 21.9% in diabetics and 22.9% in non-diabetics. For patients aged 51-60, UPEC was observed in 12.5% of diabetic patients and 17.1% of non-diabetic patients. Among those aged 61-70, 7.8% of diabetic patients and 8.6% of non-diabetic patients had UPEC isolates. For patients aged 71 and above, UPEC was present in 3.1% of diabetic patients and 2.9% of non-diabetic patients. The chi-square test for this variable resulted in  $p = 0.99$ , indicating no significant association between age and UPEC prevalence.

The gender distribution showed that among female patients, 56.3% of diabetic patients and 57.1% of non-diabetic patients were infected with UPEC. For male patients, 43.8% of diabetics and 42.9% of non-diabetics had UPEC isolates. The chi-square test for gender resulted in  $p = 0.93$ , suggesting no significant relationship between gender and UPEC prevalence.

The marital status of patients was categorized as single, married, divorced, or widowed. Among single patients, UPEC was found in 15.6% of diabetic patients and 14.3% of non-diabetic patients. In married patients, the prevalence was 65.6% in diabetics and 57.1% in non-diabetics. For divorced patients, 12.5% of diabetics and 22.9% of non-diabetics had UPEC isolates. Among widowed patients, UPEC was present in 6.3% of diabetic patients and 5.7% of non-diabetic patients. The chi-square test for marital status resulted in  $p = 0.62$ , indicating no significant association between marital status and UPEC prevalence.

The residence of patients was classified as rural or urban. Among rural residents, UPEC was found in 32.8% of diabetic patients and 31.4% of non-diabetic patients. For urban residents, the prevalence was 67.2% in diabetics and 68.6% in non-diabetics. The chi-square test for residence resulted in  $p = 0.88$ , indicating no significant association between residence and UPEC prevalence. The education level of patients was divided into literate and illiterate categories. Among literate patients, 62.5% of diabetic patients and 54.3% of non-diabetic patients were infected with UPEC. For illiterate patients, the prevalence was 37.5% in diabetics and 45.7% in non-diabetics. The chi-square test for education resulted in  $p = 0.42$ , showing no significant relationship between education level and UPEC prevalence.

UTIs are more common and severe in patients with DM. they are also frequently caused by resistant pathogens [42]. Urinary tract infection is the commonest bacterial infection with a high rate of morbidity and financial cost. Uropathogenic *E. coli* (UPEC) is the commonest etiological agent of UTI in diabetic and non-diabetic patients. UPEC associated UTIs among diabetic and non-diabetic patients have been reported in previous studies [37,35,36]. The most

frequently isolated microorganism from urine sample of this study is *E. coli*. The risk of developing urinary tract infection in diabetes is higher due to abnormalities in the host defense and high glucose in urine, the occurrence and infection with UPEC have been increased in diabetic patients because hyperglycemia suppress the level of immune system among this category of people [43].

The age distribution of patients with UPEC isolates was examined across different age groups. Among patients aged 10-20 years, 7.8% of diabetic patients and 5.7% of non-diabetic patients were infected with UPEC. In the 21-30 age group, the prevalence was 17.2% for diabetics and 20% for non-diabetics. For the 31-40 age group, UPEC was found in 28.1% of diabetic patients and 25.7% of non-diabetic patients. In the 41-50 age range, the prevalence was 21.9% in diabetics and 22.9% in non-diabetics. For patients aged 51-60, UPEC was observed in 12.5% of diabetic patients and 17.1% of non-diabetic patients. Among those aged 61-70, 7.8% of diabetic patients and 8.6% of non-diabetic patients had UPEC isolates. For patients aged 71 and above, UPEC was present in 3.1% of diabetic patients and 2.9% of non-diabetic patients

In this study, the age range of 31-40 had the highest rate of *E. coli* infection in both the study participants, 28.1% in diabetic and 25.7% in non-diabetic patients. This is in agreement with previous researches done by [39,41] who all reported the high rate of urinary tract infection in this range: reported high-rate UTI in that age range. While this study is not in agreement with the reports of [44], who conducted his research in Sudan reported high rate in the age range 44 and above, [45] reported age range of 40-49 have the highest range of infection and also [46] from Kebbi Nigeria who reported high rate of frequency in the age range of 61-65. High rate of occurrence of UTI infection in this age range (31-40) may be attributed to the fact that people are more sexually active at this age range, and also difference might be due to the competent immune system and high treatment-seeking behavior at a young age [47].

The gender distribution showed that among female patients, 56.3% of diabetic patients and 57.1% of non-diabetic patients were infected with UPEC. For male patients, 43.8% of diabetics and 42.9% of non-diabetics had UPEC isolates.

In this study, the female gender in both the study population has a high rate of occurrence compared with male counterparts. This could be attributed to the fact that the female gender anatomical structure, short urethra, close proximity of the urethra to the anus, decreases of normal vagina flora, in general lifestyle habits of women are some of the predisposing factors that can increase the occurrence of UTI in females than males regardless of diabetic status [48,43]. The result is in disagreement with the reports of [46,45] who reported high rate of infection in males than their female counterparts.

The marital status of patients was categorized as single, married, divorced, or widowed. Among single patients, UPEC was found in 15.6% of diabetic patients and 14.3% of non-diabetic patients. In married patients, the prevalence was 65.6% in diabetics and 57.1% in non-diabetics. For divorce divorced patients, 12.5% of diabetics and 22.9% of non-diabetics had UPEC isolates. Among widowed patients, UPEC was present in 6.3% of diabetic patients and 5.7% of non-diabetic patients

In this study, married participants have higher rate of infection in both diabetic and non-diabetic patients. This may be attributed to the fact that they are sexually active at this status, while the widows have the lowest rate of infection in both the study population. This is in accordance with the report of [16].

The education level of patients was divided into literate and illiterate categories. Among literate patients, 62.5% of diabetic patients and 54.3% of non-diabetic patients were infected with UPEC. For illiterate patients, the prevalence was 37.5% in diabetics and 45.7% in non-diabetics.

Literate have high rate of infection in DM compared to NDM patients. While illiterate have high rate of infection in NDM patients than NDM patients.

The residence of patients was classified as rural or urban. Among rural residents, UPEC was found in 32.8% of diabetic patients and 31.4% of non-diabetic patients. For urban residents, the prevalence was 67.2% in diabetics and 68.6% in non-diabetics. Urban dwellers have high rate of occurrence of UTI in both DM and NDM patients than rural dwellers. This may be attributed to the fact that the study was conducted in the city and most of the study participants live in urban areas.

### 3.3 Clinical Profile of UPEC Infection Among Participants Investigated in the Study

In this study, the rate of occurrence of UPEC infection base on history of UTI is higher in first time encounter of UTI in both diabetic and non-diabetic patients. In this study, history of antibiotic usage shows high rate in DM than NDM patients in those that did not use antibiotics, while slightly higher in those that use antibiotics in NDM than DM patients. As shown in table 3.

**Table 3: Distribution of UPEC Infection According to Clinical Variables of Patients**

| Clinical Variables                     | No of Samples Collected (n=288) | No (%) of UPEC in DM (n=64) | No (%) of UPEC in NDM (n=35) | <i>p</i> |
|--|---------------------------------|-----------------------------|------------------------------|----------|
| <b>History of UTI</b>                  |                                 |                             |                              |          |
| Yes                                    | 84                              | 10 (15.6)                   | 9 (25.7)                     | 0.22     |
| No                                     | 204                             | 54 (84.4)                   | 26 (74.3)                    |          |
| <b>History of Antibiotic Treatment</b> |                                 |                             |                              |          |
| Yes                                    | 23                              | 5 (7.8)                     | 3 (8.6)                      | 0.89     |
| No                                     | 265                             | 59 (92.2)                   | 32 (91.4)                    |          |
| <b>Types of Diabetes</b>               |                                 |                             |                              |          |
| Type 1                                 | 92                              | 16 (25)                     | N.A                          | -        |
| Type 2                                 | 196                             | 48 (75)                     | N.A                          |          |
| <b>Duration of Diabetes</b>            |                                 |                             |                              |          |
| < 5 years                              | 113                             | 35 (54.7)                   | N.A                          | -        |
| ≥ 5 years                              | 175                             | 39 (60.9)                   | N.A                          |          |

Key: N.A = non-applicable, DM = Diabetic Mellitus

The history of urinary tract infections (UTI) was examined in relation to UPEC prevalence. Among patients with a history of UTI, 15.6% of diabetic patients and 25.7% of non-

diabetic patients were infected with UPEC. For those without a history of UTI, the prevalence was 84.4% in diabetics and 74.3% in non-diabetics. The chi-square test for this variable resulted in  $p = 0.22$ , indicating no significant association between UTI history and UPEC prevalence.

In this study, DM patients with no previous history of UTI had higher rate of contracting UTI compared with those who had previous history of UTI

In this study, the frequency of UTI was higher among duration of DM greater than 5 years compared to those patients of DM duration less than 5 years. This is in agreement with the studies conducted by [21,44] from Sudan found high rate of infection present in  $\geq 5$  years of DM duration. It is very well known that patients with longer duration of DM have increased prevalence of diabetic chronic complications, which may lead to an increased presence of UTI [42]. In many of these patients, autonomic neuropathy results in dysfunctional voiding and urinary retention. [40]

The history of antibiotic treatment was also analyzed. Among patients with a history of antibiotic treatment, 7.8% of diabetic patients and 8.6% of non-diabetic patients had UPEC isolates. For those without such history, the prevalence was 92.2% in diabetics and 91.4% in non-diabetics. The chi-square test for this variable resulted in  $p = 0.89$ , indicating no significant association between antibiotic treatment history and UPEC prevalence.

Various studies have demonstrated that different outbreak of urinary tract infection in type 2 diabetic patients. Factors such as immune system disorders, weakening of white blood cells, poor blood supply, bladder dysfunction due to nephropathy and glucosuria can cause urinary tract infections in type 2 diabetic patients [5]. Dysuria is a complication of urinary tract infection in diabetic patients due to organ damage and even death due to the complexity of pyelonephritis. Also, these patients experience urinary retention, urgency, and incontinence during the night due to increased urination to excrete excess glucose. In this study, type 2 DM patients have high rate of infection compared to type1 DM patients, type 2 has a prevalence of 25% while type 1 has a prevalence of 25% in this study. This is similar to a study conducted by [49] where high rate of infection was observed in type 2 DM patients

### 3.4 Prevalence of UPEC isolates according to MDR pattern in diabetic and non-diabetic patients

In this study, antibiotics of known concentration were used to screen UPEC isolates that are resistant to antibiotics. Ampicillin and piperacillin-tazobactam have the highest rate of resistance in DM and NDM patients, while the resistance was observed in imipenem, ceftriaxone and ceftazidime. While the other antibiotics have moderate rate of resistance to the isolates.

**Table 4: Distribution of UPEC isolates according to MDR pattern in diabetic and non-diabetic patients**

| Antibiotics( $\mu$ g)        | No (%) of UPEC isolates and MDR pattern<br>n=64-for DM, n=35 for NDM |  |
|------------------------------|--|--|
|                              | No (%) Resistant isolates in Diabetic (n=52)                         | No (%) Resistant isolates in non diabetic (n=27) |
| Ampicillin (10)              | 48(92.3)   | 22(81.5)   |
| Amoxicillin-clavulanate (30) | 39(75)   | 11(40.7)   |

|                               |          |          |
|-------------------------------|----------|----------|
| Ceftriaxone (30)              | 16(30.8) | 08(29.6) |
| Cefuroxime (30)               | 31(60.0) | 16(59.3) |
| Ceftazidime (30)              | 17(32.7) | 08(29.6) |
| Kanamycin (30)                | 42(80.8) | 21(77.8) |
| Amikacin (30)                 | 39(75)   | 20(74.1) |
| Gentamicin (10)               | 42(80.8) | 19(70.4) |
| Streptomycin (10)             | 38(73.1) | 18(66.7) |
| Ciprofloxacin (5)             | 27(51.9) | 13(48.1) |
| Levofloxacin (5)              | 26(50)   | 11(40.7) |
| Nalidixic acid (30)           | 47(90.4) | 21(77.8) |
| Ofloxacin (5)                 | 28(53.8) | 10(37.0) |
| Piperacillin tazobactam (100) | 48(92.3) | 22(81.5) |
| Imipenem (10)                 | 10(19.2) | 03(11.1) |

Key DM- Diabetic mellitus, NDM- non diabetic mellitus, UPEC-Uropathogenic *E.coli*, MDR-multidrug resistance.

The presence of multidrug resistance in this study could be attributed to the dissemination of antibiotic resistance among UPEC isolates. UPEC can be seen as one of the commonest pathogens causing UTI in immunocompromised patients such as diabetics. In this study, high rate of MDR was encountered in UPEC isolate of both DM and NDM patients. Ampicillin has resistant rate of 92.3% and 81.5% in diabetic and non-diabetic patients respectively. This is in agreement with studies of [50,43]. Other drugs that were resistance were piperacillin-tazobactam, nalidixic acid.

Quinolone and fluoroquinolone resistance of UPEC isolates were correlated with previous studies that resistance rate of diabetic UPEC to ciprofloxacin [50]. Resistance to fluoroquinolone among diabetic urinary *E. coli* isolates were 89.5% and non-diabetic urinary *E. coli* isolates were 50% have been reported. Common and overuse of quinolones and fluoroquinolone worldwide in the treatment of UTIs led to increased resistance in UPEC. [2].

On the other hand, imipenem, ceftriaxone, ceftazidime was found to show least resistance among the drugs used. This is agreement with the reports of [51,35,43]. The isolates resistant to three or more classes of antibiotics were termed as MDR. Diabetes can be a factor associated with MDR *E. coli*. In this study, high rates of MDR *E. coli* in both diabetic and non-diabetic patients was observed as with the reports of [43,51]. The rapid development of resistance could be attributed to the irrational use of antibiotics and practices of self-medication among the general population thereby causing a problem in antibiotic therapy especially in developing countries due to lack of awareness and lack of effective implementation of the policy that regulates the use of antibiotics [52].

#### 4. CONCLUSION

In conclusion, UPEC isolates were high in diabetic than non-diabetic patients. it could be observed that most of the UPEC isolates from both the groups exhibited remarkable rate of antibiotic resistance to commonly prescribed antibiotics for UTI irrespective of diabetic status. The study revealed that a high rate of multidrug resistance from both the study participants, this reaffirms for the need of proper diagnosis and drug administration in the treatment of urinary tract infection especially in diabetic patients due to their immunological status.

### CONSENT AND ETHICAL APPROVAL

Ethical approval was granted by; Abubakar Tafawa Balewa University Teaching Hospital, Ministry of Health Bauchi State Government Ethical Steering Committee. More so, written informed consent was obtained from all patients prior to specimen collection.

### REFERENCES

1. Acharya D, Bogati B, Shrestha G, Gyawali P. Diabetes mellitus and Urinary Tract Infection: Spectrum of Uropathogens and their Antibiotic Sensitivity. JMMIHS. 2015;1(4), 24–28. <https://doi.org/10.3126/jmmihs.v1i4.11998>.
2. Kot B. Virulence Factors and Innovative Strategies for the Treatment and Control of Uropathogenic *Escherichia coli*. In A. Samie (Ed.), *Escherichia coli—Recent Advances on Physiology, Pathogenesis and Biotechnological Applications*. InTech. 2017. pp 24-36. <https://doi.org/10.5772/67778>.
3. Vasudevan R. Urinary Tract Infection: An Overview of the Infection and the Associated Risk Factors. J Microbiol Exp. 2014; 1(2). <https://doi.org/10.15406/jmen.2014.01.00008>
4. Kaur P, Karnwal AS, Devgon I. “A Review on Clinical Manifestation and Treatment Regimens of UTI in Diabetic Patients.” Iran J Med Microbiol. 2022; 16(2):98–115. doi: 10.30699/ijmm.16.2.98
5. Jagadeesan S, Tripathi BK, Patel P, Muthathal S. Urinary tract infection and diabetes mellitus—Etiological profile and antibiogram: A North Indian perspective. J Family Med Prim Care 2022; 11:1902-6.
6. Butt MD, Ong SC, Wahab MU, Rasool MF, Saleem F, Hashmi A, Sajjad A, Chaudhry FA, Babar ZUD. Cost of Illness Analysis of Type 2 Diabetes Mellitus: The Findings from a Lower-Middle Income Country. Int. J. Environ. Res. Public Health 2022, 19, 12611. <https://doi.org/10.3390/ijerph191912611>
7. Goyal SR, Chand AE, Nakura HS. Prevalence of uropathogens among diabetic patients and their antibiogram at government medical college, Kota. IPInt J MedMicrobiol Trop Dis. 2021; 7(1), 24–27. <https://doi.org/10.18231/j.ijmmt.2021.006>
8. Ahmad S, Hussain A, Khan MSA, Shakireen N, Ali I. Diabetes mellitus and urinary tract infection: Causative uropathogens, their antibiotic susceptibility pattern and the effects of glycemic status. Pak J Med Sci. 2020; 36(7). <https://doi.org/10.12669/pjms.36.7.2881>

9. Nicolle LE. Epidemiology of urinary tract infection. *Infectious medicine*. 2001;**18**(3):153 – 162.
10. Unchu Y, Unchu G, Esmer A, Bilgel N. Should asymptomatic bacteriuria be screened in pregnancy. *Clin. Exp. Obstet. Gynecol*. 2002;**29**(4):281-51.
11. Akram M, Shahid M, Khan, AU. Etiology and Antibiotic Resistance Pattern of community- acquired urinary tract infections in JNMC Hospital Aligarh, India. *AnnClin Microbiol and Antimicrob*. 2007; **6**: 4-4.
12. Gupta K, Hootan TM, Naber KG, John AD. International Clinical Practice Guidelines for the Treatment of Acute Uncomplicated Cystitis and Pyelonephritis in Women: A 2010 Update by the Infectious Disease Society of America and the European Society for Microbiology and Infectious Diseases. *Clin Infect Dis*. 2011; **52**:103-120.
13. Zubair KU, AbdulHaleem S, Asher F, Rubina S, Anum B. “Frequency of Urinary Tract Infection and Antibiotic Sensitivity of Uropathogens in Patients with Diabetes: Frequency of Urinary Tract Infection and Antibiotic Sensitivity.” *Pak J Med Sci*. 2019; 35(6). doi: 10.12669/pjms.35.6.115
14. Abu-Ashour W, Twells L, Valcour J, Randell A, Donnan J, Howse P, Gamble JM. The association between diabetes mellitus and incident infections: a systematic review and meta-analysis of observational studies. *BMJ Open Diab Res Care*. 2017; 5: 000336. doi:10.1136/ bmjdr-2016-000336
15. Kumar R, Kumar R, Perswani P, Taimur M, Shah A, Shaukat, F. Clinical and Microbiological Profile of Urinary Tract Infections in Diabetic versus Non-Diabetic Individuals. *Cureus*. 2019. <https://doi.org/10.7759/cureus.5464>
16. Chito Clare E, Oluchi JA, Chioma MO, James RA, Evelyn AKO. Urinary Tract Infections and Antimicrobial Sensitivity Patterns of Uropathogens Isolated from Diabetic and Non-diabetic Patients Attending Some Hospitals in Awka. *AJMR*. 2021; 9(3), 83–91. <https://doi.org/10.12691/ajmr-9-3-3>
17. Alrwithey FA, Alahmadi AEA, Alshehri AMF, Abalhassan A, Alhamad FMS, Khedher YZ, Mohammed A, Beig STM, Althubayani AA, Ghamdi RAA. Urinary Tract Infection in Patients with Diabetes Mellitus. *Egypt J Hosp Med*. 2017; 69 (3),2133-2136
18. Tabasi M, Karam MRA, Habibi M, Mostafavi E, Bouzari S. Genotypic Characterization of Virulence Factors in *Escherichia coli* Isolated from Patients with Acute Cystitis, Pyelonephritis and Asymptomatic Bacteriuria. *J Clin Diag Res*. 2016 ;10(12): DC01-DC07.doi: 10.7860/JCDR/2016/21379.9009.
19. Bartoletti R, Cai T, Wagenlehner FM, Naber K, Johansen TEB. Treatment of Urinary Tract Infections and Antibiotic Stewardship. *Eur Urol Suppl*. 2016; 15(4), 81-87. <https://doi.org/10.1016/j.eursup.2016.04.003>.
20. Zhu Y, Wei EH, Qiwen. “Clinical Perspective of Antimicrobial Resistance in Bacteria.” *Infection and Drug Resistance*. 2022;15:735–46. doi: 10.2147/IDR.S345574
21. [Nigussie D](#), and [Amsalu A](#). Prevalence of uropathogen and their antibiotic resistance pattern among diabetic patients. *Turk J Urol*. 2017; 43(1):85-92. Doi: 10.5152/tud.2016.86155.

22. Mogaka MV. Prevalence of urinary tract infections among type 2 diabetic patients in Kisii Teaching and Referral Hospital, Kenya. *JMed SciClin Res.* 2020; 08(08). <https://doi.org/10.18535/jmscr/v8i8.01>
23. Mwang'onde BJ, and Mchami JI. The aetiology and prevalence of urinary tract infections in Sub-Saharan Africa: A Systematic Review. *JHealth BiolSci.* 2022; 10(1), 1. <https://doi.org/10.12662/2317-3076jhbs.v10i1.4501.p1-7.2022>
24. Rushita BP, Mayur G, Pronam BC, Hiren P. Isolation and characterization of *E. coli* from urine sample and their antibiogram pattern and effect of spice extracts and natural oils against isolated *E. coli*. *Int J Curr Adv Res.* 2016; 5(4), pp 834 – 838.
25. Naing L, Winn T, Rusli BN. Practical Issues in Calculating the Sample Size for Prevalence Studies. *ArchOrofasc Sci.* 2006; **1**: 9-14.
26. Yakubu H, Muhammed B, Mukhtar MD, Kalgo ZM. Prevalence Of Extended-Spectrum Beta Lactamase and Ampc Producing Enterobacteriaceae Among Diabetic Patients in Bauchi State, Nigeria. *Bayero J Pure App Sci.* 2022; 13(1):546-553 ISSN 2006 – 6996. <http://dx.doi.org/10.4314/bajopas.v13i1.82S>.
27. World Health Organization. *Classification of diabetes mellitus.* World Health Organization. 2019. <https://apps.who.int/iris/handle/10665/325182>
28. Cheesbrough M. *District Laboratory Practice in Tropical Countries* 2ed Cambridge University press, New York. 2006; Pp: 62-70.
29. Madigan TM, John MM, Paul VP, David PC. **Biology of Microorganisms** 12<sup>th</sup> ed. Pearson International Edition 1301 Sansome Street, San Francisco. 2009; Pp. 902-911.
30. Signing AT, Marbou WJT, Beng VP, Kuete V. Antibiotic Resistance Profile of Uropathogenic Bacteria in Diabetic Patients at the Bafoussam Regional Hospital, West Cameroon Region. *Cureus.* 2020; 12(7): e9345. DOI 10.7759/cureus.9345.
31. Vineetha N, Vignesh RA, Sridhar D. Preparation, Standardization of Antibiotic Disc and Study of Resistance Pattern for First -Line Antibiotics in Isolates from Clinical Samples. *Int JApp Res.* 2015; 1(11): 624-631.
32. Tena D, Gonzalez PA, Saez-Nieto JA, Valdezate S, Bisquert J. Urinary tract infection caused by capnophilic *Escherichia coli*. *Emerg Infect Dis J.* 2008; 10: 140-150.
33. CLSI. *M100 Performance Standards for Antimicrobial Susceptibility Testing.* 29<sup>th</sup> edition. Wayne, U.S.A. 2019; Pp 33-38.
34. Wang M, Tseng C, Wu A, Lin W, Teng C, Yan J, Jiunn-Jong WJ. Bacterial characteristics and glycemic control in diabetic patients with *Escherichia coli* urinary tract infection. *J Microbiol ImmunolInfect.* 2013;46, 24e29
35. Ramrakhia S, Raja K, Dev K, Kumar A, Kumar V, Kumar, B. Comparison of Incidence of Urinary Tract Infection in Diabetic vs Non-Diabetic and Associated Pathogens. *Cureus.* 2020. <https://doi.org/10.7759/cureus.10500>
36. Taher J, Rashed STM, Shwekh FT, Mareedh TK, Ghazi HF. Bacterial Causes of Urinary Tract Infection among Diabetic and Non-Diabetic Patients in Al-Kut City, Iraq. *Borneo Epidemiol J.* 2021; 2(1). <https://doi.org/10.51200/bej.v2i1.3242>

37. Saber MH, Barai L, Haq JA, Alam JMS, Begum J. The Pattern of Organism Causing Urinary Tract Infection in Diabetic and Non Diabetic Patients in Bangladesh. *Bangladesh JMed Microbiol.* 2010; 4(1), 6–8. <https://doi.org/10.3329/bjmm.v4i1.8461>
38. Srinivas A, Chandrashekar UK, Shivashankara KN, Pruthvi BC. Clinical profile of urinary tract infections in diabetics and non-diabetics. *Aust Med J.* 2014; 7(1), 29–34. <https://doi.org/10.4066/AMJ.2014.1906>
39. Jameel AY, and Artoshi DME. Prevalence of Urinary Tract Infections and Their Antimicrobial Sensitivity Among Diabetic and Non Diabetic Patients in Zakho. *Sci JUniZakho.* 2019; 7(4), 125–131. <https://doi.org/10.25271/sjuoz.2019.7.4.629>
40. Al Qurabiy HE, Abbas IM, Hammadi ATA, Mohsen FK, Salman RI, Dilyf SH, Saja H. Urinary tract infection in patients with diabetes mellitus and the role of parental genetics in the emergence of the disease. *JMedLife.* 2022; 15(8), 955–962. <https://doi.org/10.25122/jml-2021-0331>
41. Hanson AH, Akani NP, Williams JO, Sampson T. Bacteriuria Among Diabetic and Non-diabetic Patients in some Hospitals in Rivers State, Nigeria. *Nig JMicrobiol.* 2023; 37(1): 6432 - 6441
42. Bagir GS, Haydardedeoglu FE, Colakoglu, S, Bakiner OS, Ozsahin AK, Ertorer ME. Urinary tract infection in diabetes: Susceptible organisms and antibiogram patterns in an outpatient clinic of a tertiary health care center. *Med Sci.* 2019;8(4):881-6.
43. Surendran A, Suganthi P, SudhaLakshmi SY. Prevalence of  $\beta$ -lactamase enzyme producing uropathogenic *E.coli* isolates amongst diabetic and non-diabetic patients. *Preprints.* 2024. <https://doi.org/10.22541/au.171308242.27630569/v1>
44. Abdelgader LMA, Omer SS, Mahjaf GM, Altaher, TAA, Hamad MNM. Determination of Bacteriological Profile of Antimicrobial Susceptibility Testing among Diabetic Patients with Urinary Tract Infections in Shendi, Sudan. *SAR JPathMicrobiol.* 2023; 4(03), 24–29. <https://doi.org/10.36346/sarjpm.2023.v04i03.001>
45. Rajendran J, Ramya S, Anandhalakshmi S, Kanungo R. Clinical profile, predisposing risk factors for urinary tract infections and its outcome in diabetics and nondiabetics: The significant variants. *JCurr ResSciMed.* 2023; 9(1), 67. [https://doi.org/10.4103/jcrsm.jcrsm\\_57\\_22](https://doi.org/10.4103/jcrsm.jcrsm_57_22)
46. Almustapha AA., Yari HA, Ibrahim AB, Manga SS, Tomo FA, Muhammad A, Muhammad MS. *Escherichia coli* dominance and antimicrobial resistance in urinary tract infections among diabetic patients: Insights from Birnin Kebbi Metropolis, Nigeria. *Acta Pharm Indo.* 2023; 11(1), 6231. <https://doi.org/10.20884/1.api.2023.11.1.6231>
47. Sharifi Y, Hassan A, Ghotaslou R, Naglili B, Aghazdeh M, Milani M, Bazmany A. virulence and Antimicrobial Resistance in Enterococci Isolated from Urinary tract infections. *Advanced pharmaceutical Bulletin.* 2013; 3: 197-201.
48. Okwume CC, Onyemelukwe NF, Abdullahi IN, Okoyeocha OE, Asamota, SD. Prevalence of symptomatic urinary tract infection and bacterial spectrum of diabetic and

non-diabetic patients at the two teaching hospitals in Enugu, Nigeria. *AfJ Clin Exp Micro.* 2021; 22(4), 480–488. <https://doi.org/10.4314/ajcem.v22i4.8>

49. Nabi T. Clinical characteristics and complications of symptomatic bacteriuria in patients with Type 2 diabetes. *IntJ Acad Med.* 2021; 7(4), 212. [https://doi.org/10.4103/IJAM.IJAM\\_153\\_20](https://doi.org/10.4103/IJAM.IJAM_153_20)
50. Raya S, Belbase A, Dhakal L, Govinda PK, Baidya R, Kishor BN. In-Vitro Biofilm Formation and Antimicrobial Resistance of *Escherichia coli* in Diabetic and Nondiabetic Patients. *BioMed Res Int.* 2019, 1–8. <https://doi.org/10.1155/2019/1474578>
51. Maharjan N, Thapa N, Maharjan M, Sharma VK, Shrestha P, Paudyal R. Pattern of Bacteria Causing Urinary Tract Infection and Their Antibiotic Susceptibility Profile In Diabetic And Non-Diabetic Patients In Lalitpur, Nepal- A Hospital Based Study. *Int J Inno Sci Res.* 2018; 7(8), 1248–1253.
52. Gutema G, Håkonsen H, Engidawork E, Toverud EL. Multiple challenges of antibiotic use in a large hospital in Ethiopia – a ward-specific study showing high rates of hospital-acquired infections and ineffective prophylaxis. *BMC Health Serv Res.* 2018; 18(1), 326. <https://doi.org/10.1186/s12913-018-3107-9>