Original Research Paper

# Antimicrobial Susceptibility of Community Acquired *Escherichia coli* in Urinary Tract Infections (UTI) in Benin for Eleven Years (2005-2015)

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Corresponding Author: Tchiakpe Edmond National Laboratory Fighting Against AIDS, National Direction of Health, Health Ministry, 04PO Box 378 Cotonou, Benin E-mail: edtchi@yahoo.fr Abstract: Antimicrobial resistance became a growing public health problem in the world and Escherichia coli (E. coli) appeared as one of nine bacteria commonly causing infections in community and hospitals. It prevalence and it resistance to antibiotics were evaluated in Benin throughout an eleven year period. In this retrospective study, routine urine samples from patients collected at the National Laboratory (NL) of Health Ministry during the period 2005-2015 were analyzed. Samples higher than 103 CFU/mL bacterial growth were considered positive and for these cases, the bacteria were identified and Antimicrobial Susceptibility Test (AST) was performed. From the 4467 samples analyzed, 1455 (32.6%) were positive with E. coli preponderance (38.3%) of all isolated germs and (58.5%) of enterobacteriaceae. Most of the isolates were susceptibility to netilmicin (80%). gentamycin (93%), chloramphenicol (70%), pipemidic acid (60%), nalidixic acid (75%), ciprofloxacin (75%), amoxicillin/clavulanic acid (80%) and nitrofurantoin (100%). Susceptibility rates increased for cefotaxim (78-100%), ceftriaxon (71-100%) and aztreonam (67-100%). Resistances were observed for minocyclin (70%), Trimethoprim/Sulfamethoxazole (TMP-SMX) (60%), ampicillin (67%), amoxicillin (75%), carbenicillin (86%), cephalothin and cephalexin (50 and 80% respectively). The use of drugs such as minocyclin, carbenicillin. ampicillin. amoxicillin. cephalothin. cephalexin and trimethoprim/sulfamethoxazole does not seem appropriate for empirical treatment of UTI in Benin.

Keywords: Community, Urinary Tract Infections, *Escherichia coli*, Antimicrobial Effectiveness, Benin

## Introduction

Antimicrobial resistance had become a growing public health problem in the world and *E. coli* appeared as one of nine bacteria commonly causing infections in

community and hospitals (WHO, 2014). Worldwide, *E. coli* remains the most encountered pathogen among those implicated in community acquired urinary tract infection (Jean-Marie *et al.*, 2007). With (46.4-74.2%) of global representation (Inês *et al.*, 2013), it's the first



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strains isolated in children between 2 to 5 years of age followed by Klebsiella, Proteus and Pseudomonas (Rajiv et al., 2013). In a recent study done in 2012 at Ngaoundere in Cameroon focused on community acquired infection, Carine et al. (2012) found that 66.7% of E. coli was mostly Extended-spectrum betalactamase producers in the samples of stools. It is well known the proximity between anal cavity and urinary tract in women. Urinary Tract Infections (UTI) is the most commonly diagnosed infections in communities which are often treated with different antibiotics. Some studies have reported an increasing resistance rate to antibiotics such as penicillin, amoxicillin amino trimetropimreducing sulfamethoxazole others and therapeutic possibilities (Gupta et al., 1999; Kahlmeter, 2003a).

A study conducted in Benin have shown, that the beta lactam antibiotics were the most prescribed and represented 44.4% of prescriptions, followed, in order quinolones, of imidazole, aminoglycosides, sulfonamides, macrolides and related, cyclins and phenicol (Dissou et al., 2009). However, in Benin, few documentations exist in antimicrobial resistance of uropathogen isolated in UTI. The aim of this study was to describe the bacterial profile of UTI and antimicrobial susceptibility of E. coli strains isolated from the urine cultures of Benin patients in National Laboratory (NL) of Heath Ministry over 11 year period.

## Methods

Routine urine samples data were collected from patients in Cotonou between 1<sup>st</sup> January 2005 and 31<sup>st</sup> December 2015 period. This study was approved by the Research Ethics Committee for Applied Biomedical Sciences (CER-ISBA) of Abomey Calavi University of Benin. A retrospective cross sectional study was conducted in NL where urine samples were analyzed. For each patient we extracted from the laboratory records, year, age, sex, early urine culture results, identification of the bacterial strain responsible of UTI and the corresponding Antimicrobial Susceptibility Test (AST) results.

The statistical test EpiData 3.1 was used to check-in results and EpiData Analysis V2.2.2.182 for data analysis. Data were expressed an absolute values and percentage. The chi-2 test was used to compare proportions. To be significant, the p value should be less than 0.05.

# **Results**

Overall 4467 urine cultures were performed during 2005-2015 period. The rate of positive cultures was 32.6% (1455/4467) of which 1470 bacteria have been isolated. Characteristics of patients included in the study are shown in Table 1. Mean age and sex ratio (M/F) of study population were 30.5 years [95% Confidence Interval (CI) 30.01-30.98] and 0.5. Frequencies and distribution of isolated uropathogenic bacteria during the eleven study years were summarized in Table 2. The enterobacteriaceae proportion was 65.5% (963/1470) with E coli preponderance which represented 38.3% (563/1470) of all isolated bacteria and 58.5% (563/963)of enterobacteriaceae. The others most represented isolated bacteria were Klebsiella pneumoniae 23.4% (344/1470), Staphylococcus aureus 16.0% (235/1470), Staphylococcus spp 7.6% (111/1470), Staphylococcus dore 3.7% (54/1470).

The Fig. 1 shows the susceptibility rates of E. coli for aminoglycosides, tetracycline, phenicol, quinolones, sulfonamides+associations, penicillin, cephalosporin, aztreonam and nitrofurantoin.

Regarding aminoglycosides, we observed susceptibility increase of E. coli to netilmicin (10 µg) (80 to 100%) on three use years. During the 11 study years, the maximum susceptibility rate to gentamycin (30 µg) was 93.3%.

Since 2007, E. coli susceptibility to tetracyclin (30 µg) decrease to 12% at 2015. Concerning minocyclin (30 μg), on three use years, his susceptibility rate was 30%.

Among phenicol, chloramphenicol (30 µg) was one used during the first ten years. E. coli susceptibility to this drug varied between (38 to 70%) until 2014.

Among quinolones, pipemidic acid (20 ug) was tested during three years (2005-2007) with E. coli susceptibility rate not exceeded 60%. The others same class antibiotics used (2005-2015) were nalidixic acid (30 ug) while the maximum E. coli susceptibility rate was 75% and ciprofloxacin (10 ug) while susceptibility rate increased and reached 100% at 2010 and decreased to 75% at 2015.

For sulfonamides+associations (25 ug) class, E. coli susceptibility to Trimethoprim/Sulfamethoxazole (TMP-SMX) not exceeded 40% (2005-2011). This susceptibility decrease from 2011 to 2015 (at 0%). Four penicillin class of antibiotics have been tested. Ampicillin (30 ug) has been used during two years (2005 and 2014) and E. coli susceptibility rate around 33%. During the 11 study years and the same class, amoxicillin (25 ug) was less active to E. coli while susceptibility rate not exceeded 25%. As for Amoxicillin/Clavulanic acid (AMC) (30 ug), the most tested antibiotic in this class, effectiveness was observed until 2009 (80%). But from this year, a loss effectiveness was observed until 2015 (62%). On three use years, carbenicillin (100 ug) was less active to E. coli while susceptibility rate not exceeded 14%.

Table 1. Characteristics of patient	ts included in the study

Characteristic	All patients ( $N = 4467$ )
Sex	
Male	1576 (35.28%)
Female	2891 (64.72%)
Age (years)	
Minimum	1
Maximum	95
Mean	30.50
Cultures	
Positive	1455 (32.57%)
Negative	3012 (67.43%)

Tchiakpe Edmond *et al.* / American Journal of Infectious Diseases 2017, 13 (2): 21.27 DOI: 10.3844/ajidsp.2017.21.27 . .

Table 2. Percentage and distribution of isolated uropathogenic germs during the eleven study years												
Bacteria	n (%)	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015
Escherichia coli	563 (38.3)	119	118	67	28	34	35	16	45	53	40	8
Klebsiella pneumoniae	344 (23.4)	49	40	29	29	32	25	25	48	33	22	12
Staphylococcus aureus	235 (16.0)	38	47	28	10	26	10	13	22	21	17	3
Staphylococcus spp	136 (9.7)	0	8	8	7	10	7	11	24	23	29	9
Staphylococcus dore	54 (3.7)	50	2	1	0	0	0	0	1	0	0	0
Streptococcus $\beta$ hemolytique	36 (2.4)	2	5	4	5	2	4	2	3	2	7	0
Pseudomonas spp	23 (1,2)	8	5	2	1	1	2	0	2	1	1	0
Staphylococcus citrin	13 (0.9)	12	0	1	0	0	0	0	0	0	0	0
Pseudomonas aeruginosa	13 (0.9)	0	3	3	0	1	2	0	0	1	2	1
Klebsiella oxytoca	13 (0.9)	0	0	2	1	3	2	1	3	1	0	0
Citrobacter spp	10 (0.7)	1	3	2	0	0	1	0	2	1	0	0
Proteus mirabilis	7 (0.5)	1	1	2	1	0	0	1	0	1	0	0
Candida albicans	4 (0.3)	0	1	2	1	0	0	0	0	0	0	0
Proteus vulgaris	4 (0.3)	2	0	0	1	0	0	0	0	1	0	0
Levinea	4 (0.3)	3	0	0	1	0	0	0	0	0	0	0
Proteus morganii	3 (0.2)	2	0	0	1	0	0	0	0	0	0	0
Bacille pyocyanique	2 (0.1)	2	0	0	0	0	0	0	0	0	0	0
Edwardsiella	1 (0.1)	0	0	0	0	0	0	0	1	0	0	0
Arizona rhinochromatis	1 (0.1)	0	0	1	0	0	0	0	0	0	0	0
Listeria monocystogene	1 (0.1)	0	1	0	0	0	0	0	0	0	0	0
Proteus rettgeu	1 (0.1)	0	0	0	0	1	0	0	0	0	0	0
Levinea malonalica	1 (0.1)	0	0	0	0	1	0	0	0	0	0	0
Streptococcus faecalis	1 (0.1)	0	0	0	0	1	0	0	0	0	0	0
Total	1470 (100,0)	289	234	152	86	112	88	69	151	138	118	33



Tchiakpe Edmond *et al.* / American Journal of Infectious Diseases 2017, 13 (2): 21.27 DOI: 10.3844/ajidsp.2017.21.27



Fig. 1. Susceptibility rates of *E. coli* for aminoglycosides, tetracyclin, phenicol, quinolones, sulfonamides+associations, penicillin, cephalosporin, aztreonam and nitrofurantoin

About cephalosporin, four antibiotics were tested: cephalothin (30 ug), cephalexin (30 ug), cefotaxim (30 ug), ceftriaxone (30 ug). Cephalothin and cephalexin have been tested three years each. *E. coli* susceptibility rate to these drugs was 50% and 20% respectively. Cefotaxim and ceftriaxon were very effectiveness and were mainly tested of antibiotics. Their effectiveness rate to *E. coli* reached 100% to 2010. From this year, these effectiveness have loss until 2015 with 33.3% and 37.5% rate respectively. *E. coli* developed resistance to cephalothin and cephalexin (50% and 80% respectively).

Aztreonam (30 ug) tested during six years showed 100% rate effectiveness to *E. coli*. Following a decrease in 2008, a continuous increase in susceptibility rates was observed until 2010 (100%).

Among nitrofurans tested, nitrofurantoin (300 ug) using till 2014, showed a good effectiveness with 100% rate.

#### Discussion

*E. coli* appeared as the most frequent bacteria in UTI representing 38.3% (563/1470) of all isolated bacteria and 58.5% (563/963) of enterobacteriaceae. The second major finding in our study was *E. coli* susceptibility to antibiotics.

In Benin, UTI were major causes of antibiotics prescribing among adults and adolescences (Dissou *et al.*, 2009). The lack of national recommandations involve their bad utilization (Dissou *et al.*, 2009). In fact, Benin is West Africa country where a lot of counterfeit drug

are circulating. A Pharmaceutical desposit drug is in Cotonou at Dantokpa market. Thus, the population supply drug and taking care by self-medication. In addition, due to their results quality, the NL received widely patients for uroculture analyzes. This should explain the 32.6% rate positive uroculture observed in this study. This rate was two fold higher that observed in Senegal (Diop-Ndiaye et al., 2014) and Portugal (Inês et al., 2013). The major bacteria isolated in those studies was E. coli as in our study (38.3%) which the virulence factors are specific properties. Those specificity confer on the bacterium the ability to adhere to the urinary tract and invade the host tissues causing injury (Moura et al., 2009). This study shows a proportion of *E. coli* (38.3%, 563/1470), significantly lower than that observed by Nisel et al. (2016) (67%, 8975/13281).

Antibiotics resistance has increased in alarming proportions (WHO, 2001a; 2001b). Thus, their effectiveness for UTI treatment became quite limited (Temesgen *et al.*, 2015). In our study, among amino glycosides, netilmicin showed susceptibility above 80%. This rate was similar those observed in Portugal and Bangui (98.7 and 96% respectively) (Hadiza *et al.*, 2003; Inês *et al.*, 2013). As for gentamycin, frontline antibiotic in the treatment of gram negative bacteria, its efficacy was 93% equal almost those observed in India, Senegal and Bangui (83.6, 93.8, 94% respectively) (Hadiza *et al.*, 2003; Jean-Marie *et al.*, 2007; Rajiv *et al.*, 2013).

As for tetracycline, from 2007 year, the resistance rate around 76%. The loss effectiveness was very

alarming and this antibiotic should not be prescribed against UTI. At Pakistan, a similar resistance rate of *E. coli* to tetracyclin was observed in 70% (Ahmad *et al.*, 2015). Minocyclin in the first three years showed effectiveness around 30% while a study conducted in China reported 92.1% of susceptibility by (Haihong *et al.*, 2015).

The maximum efficacy rate of chloramphenicol to *E. coli* was 60%. Behailu *et al.* (2016) reported in Dil Chora Referral Hospital, Dire Dawa, Eastern Ethiopia an effectiveness rate of 77.8% in study conducted among pregnant women attending at antenatal clinic (Behailu *et al.*, 2016).

Over 11 year, nalidixic acid had maximum efficacy rate at 75%. This rate was lower than observed at Bangui (90%) (Hadiza et al., 2003). However, none difference has been observed between the rate that of Dakar (76.1%) (p = 0.889) (Jean-Marie *et al.*, 2007). This antibiotic was the first among quinolone used only for the therapy of gram-negative UTI, because it has ability to penetrate the tissues (Moura et al., 2009). Of high susceptibility until 2010, the loss effectiveness of ciprofloxacin should be explain by its large use. In fact, ciprofloxacin can be used in UTI patients with allergies to others drug or in old age patients with recurrent infections and in diabetics (Eom et al., 2002; Schilling et al., 2002; Killgore et al., 2004). Nevertheless, the 75% susceptibility rate of E. coli observed in 2015 was not significant (p = 0.811) that observed in Dakar (Jean-Marie et al., 2007).

Until 2011, forty percent of *E. coli* were susceptibility to TMP-SMX. This antibiotic was also called co-trimoxazole. Some variable susceptibility rates were observed in several studies: (70.9-71.8%) in Northern Israel (Wasseem *et al.*, 2007), 31.9 and 40% at Dakar (Jean-Marie *et al.*, 2007; Diop-Ndiaye *et al.*, 2014), 15% in Bangui (Hadiza *et al.*, 2003), 35% to Pakistan (Ahmad *et al.*, 2015). TMP-SMX resistance rates of 25-68% were also reported in others countries (Urbina *et al.*, 1989; Wylie and Koornhof, 1989). After 2011 year, a total loss of effectiveness was observed with this antibiotic, that should be explained by its no medically in use through its supply by the people in the market.

The effectiveness rates of ampicillin and amoxicillin in our study were 33% and 25% respectively. Primarily, resistance to these antibiotics by  $\beta$ -lactamase producing *E. coli* come from their sub-inhibitory concentrations (Moura *et al.*, 2009). Concerning amoxicillin, some studies conducted at Dakar have shown similar rates as in our study (Jean-Marie *et al.*, 2007; Diop-Ndiaye *et al.*, 2014). As for ampicillin, the study investigated the prevalence of antibiotic resistance in urinary tract infections caused by *E. coli* in children showed 53.4% of resistance rate (Ashley *et al.*, 2016) while 67% has been reported in our study and Turkey (Nisel *et al.*, 2016). Of all cases observed in 16 countries in Europe plus Canada, the *E. coli* resistance to ampicillin was 30% (Kahlmeter, 2003b).

The amoxicillin/clavulanic acid activity depends on the level of  $\beta$ -lactamase production by *E. coli*. In our study, it

efficacy rate was 82% during all study period, more one fold higher that reported (59.3%) by (Alemu *et al.*, 2012). An opposite, Sharan reported 88% of resistance among children less of 5 years in India (Rajiv *et al.*, 2013).

A resistance rate of 86% of *E. coli* to carbenicillin was observed in our study, similar that reported (100%) on children diarrhea and swim areas on Tigris River in Baghdad city (Israa *et al.*, 2014).

Cephalosporins activity against E. coli has increased from first to fourth generation and the new classes were too much. Recently, the number of resistant strains to cephalosporins has increased (Ribeiro et al., 2002). In our study, four antibiotics of this class were tested. Cefotaxim and ceftriaxon showed effectiveness to 100% during use years. Our results were similar those observed at Dakar (97%) and Northern Israel (98%) for each antibiotic respectively (Jean-Marie et al., 2007; Urbina et al., 1989). In contrary, Sharan reported a high resistance for both antibiotics (73.5% and 73.5%) among children less of 5 years in India (Rajiv et al., 2013). For cephalosporin for three years, a 50% susceptibility rate was reported in our study. Wasseem et al (2007) reported effective rate of cephalothin to E. coli ranging between (60%-74%) in Northern Israel. About cephalexin, a high resistance rate of 80% of this antibiotic to E. coli was observed as reported by Datta in India (Datta et al., 2004).

Severe UTI requires appropriate antimicrobial treatment preferably cephalosporins (third generation). But, in patients with an allergy to cephalosporins, aztreonam may be used (Grabe *et al.*, 2015). Our study showed effectiveness of aztreonam to *E. coli* at 100%. This effectiveness was already seen since 2005 with 67% rate. In India, Iraj *et al.* (2015) reported 56.7% of susceptibility rate among children less than 12 years.

As for nitrofurantoin, a high effectiveness (100%) observed in our study was similar that observed in Portugal (94%) (Inês et al., 2013) and india (100%) (Rajiv et al., 2013). In several studies, nitrofurantoin resistance rates remained at lower levels (Arman et al., 2012; Kahlmeter and Poulsen, 2012; Nisel et al., 2016). The lower rate (1.3%) was reported equally in study performed among children used antibiotics in primary care by (Ashley et al., 2016). Because their therapy efficacy has been demonstrated clinically and microbioligically (69% and 68%) success respectively (Isikgöz et al., 2012), nitrofuratoïn should be recommended for ambulantory treatment and the cystitis and pyelonephritis treatment in women even pregnancy (Kalpana et al., 2011). It is a good alternative for uncomplicated UTI treatment when it is dispensed at appropriate dosages and suitable time intervals (Nisel et al., 2016).

Based on our above results mentioned, *E. coli* susceptibility decreased until 33% for ampicillin, 30% for minocyclin, 24% for tetracyclin and 25% for

amoxicillin and fallen to 14% for carbenicillin, 0% in 2015 for TMP-SMX, remains around 75% for ciprofloxacin, while cephalosporin third generation, aztreonam and nitrofurantoin were maintained their effectiveness rate until 100%.

These results are important in medical practice in Benin. We have confirmed now that ampicillin, minocyclin, tetracyclin, amoxicillin, carbenicillin and TMP-SMX couldn't be appropriated for treatment of UTI and that use of quinolone could exposed to resistance in 25% of cases and that third-generation cephalosporin, aztreonam and nitrofurantoïn are those that offers the highest effectiveness of UTI treatment.

One limitation of this study is the low number of patients per year that did not allow a comparative analysis of the resistance of *E. coli* during the 11 years of study.

# Conclusion

The use of drugs such as minocyclin, ampicillin, amoxicillin, carbenicillin, cephalothin, cephalexin and trimethoprim/sulfamethoxazole does not seem appropriate for empirical treatment of UTI. These findings highlighted the importance in Benin to implement national guideline of antibiotics use and done toward medic and nurse training.

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### **Author Contributions**

**Tchiakpe Edmond:** Study design, data collection, data analysis and writing.

Laurence Carine Yehouenou and Zahra Fall Malick: Study design, writing and reviewing.

Kpangon Amadohoué Arsène and Abou Abdallah Malick Diouara: Study design writing.

Keke Kpemahouton René: Final approval of the version to be submitted.

Issa Tondé: Contributed to the writing.

Honoré Sourou Bankolé and Halimatou Diop Ndiaye: Study design, data collection and writing.

Bekou Kossi Wilfried and Esse Atchéni Marius: Statistical analysis.

**Coumba Touré Kane:** Study design and data collection.

### **Conflict of Interest**

There are no conflicts of interest to declare.

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