

Original Research Article

<https://doi.org/10.20546/ijcmas.2017.605.251>

Urinary Tract Infection in Cancer Patients at Government Cancer Hospital Aurangabad, India

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ABSTRACT

Keywords

Urinary Tract Infection, Antimicrobial susceptibility pattern

Article Info

Accepted:
25 April 2017
Available Online:
10 May 2017

Urinary tract infection (UTI) is one of the major causes of morbidity in cancer patients. The purpose of the present study is to analyse the cases of urinary tract infection and to study their antimicrobial susceptibility pattern. A prospective analysis of cancer patients in year 2016-2017 was carried out at Government cancer hospital to identify the common uropathogens and to know their susceptibility pattern. Out of 121 samples processed, 42 were culture positive. Overall *E. coli* 16 (38.09%) was the predominant pathogen followed by *pseudomonas aeruginosa* 08 (19.04%), *Klebsiella pneumonia* 05 (11.9%), *Candida* species 05 (11.9%), *Acinetobacter species* 04 (9.52%), *Staphylococcus aureus* 02 (4.76%), *Enterobacter species* 01 (2.3%) and *Enterococcus species* 01 (2.3%). Susceptibility to polymyxin B and colistin was highest (100%) followed by the carbapenems (90%) and piperacillin + Tazobactam (73%). Resistance to fluoroquinolones (96%), cephalosporins (80%) and aminoglycosides (50%) was noted.

Introduction

Urinary tract infection (UTI) remains the most common hospital associated infection despite the increasing awareness of the dangers of urethral catheterization (Patwardhan, 2006). According to an estimation, about 150 million reports of UTIs per annum were recorded worldwide (Gonzalez *et al.*, 1999). About 35% of all the UTIs are of nosocomial origin (Mishra *et al.*, 2013). Urinary tract infections are the major cause of morbidity and mortality in patients with malignant diseases due to their several impairments of host defense. General factors in the etiology of UTIs are *E. coli* and other organisms of the family *Enterobacteriaceae*. In epidemiological studies, aimed at finding

the sources of nosocomial infections, it is important to identify the bacterial species involved. Most of the UTIs are treated empirically, the criteria for the selection of antimicrobial agent should be determined on the basis of the most likely pathogen and its expected resistance pattern in a geographical area (Alabi *et al.*, 2013). Knowledge of infection epidemiology and the institutional resistance pattern can help physicians to select the optimal empirical treatment in critically ill patients (Jeena *et al.*, 2013). The purpose of the present study is to analyse the cases of urinary tract infection in cancer patients and to study their antimicrobial susceptibility pattern. This study will help to

know the current resistance pattern of uropathogens and to suggest an appropriate empirical treatment.

Materials and Methods

A prospective analysis of cancer patients suspected to have UTI in period from Jan 2016 to March 2017 was carried out at government cancer hospital Aurangabad. A total of 121 midstream urine samples were received from cancer patients suspected to have UTI for routine and culture examination.

Sample collection

Precautions were taken in the collection of specimen (mid-stream urine) from each patient, in sterile screw cap wide-neck and leak proof disposable plastic container.

Gross examination: were done for color and turbidity of samples.

Microscopy

1-2 drops of urine were placed on a clean dry slide and covered with cover slip. The slide was examined by light microscope using high power field for presence of pus cells, red blood cells, epithelial cells, and bacteria.

Isolation method for optimal isolation of causative agent of UTI, CLED and SDA medium were used. Using sterile calibrated loop of 0.001, urine specimens were inoculated on culture media. The plates were incubated at 37° C for 18-24 hours. After incubation, the plates were examined for the presence of significant growth. A bacterial count of 10⁵/ml or (100 colonies or more in medium) indicates urinary tract infection. Colony morphology of grown microorganisms was studied. More than two colonies of microorganisms were not processed.

Gram stain Gram staining was performed on smear prepared from significant colonies and were examined for gram positive cocci/gram negative bacilli/ yeast cells.

Further identification was done by catalase test, coagulase test, oxidase, IMViC(indole, Methyl Red, Voges Proskauer, Citrate), Triple sugar iron agar, Nitrate reduction and urease. Yeast identification was performed by using germ tube test, corn meal agar and HiChrome candida agar (Koneman *et al.*, 2006).

Antimicrobial susceptibility testing was done by Kirby Bauer disc diffusion test on Muller Hinton agar. 0.5 MacFarland turbidity standard was used for inoculums preparation. Antimicrobial disk were placed on MH agar and incubated aerobically at 37° C overnight. *Staphylococcus aureus* (ATCC 25923), *E. coli* (ATCC 25922) and *P. aeruginosa* (ATCC 27853) were used as quality control throughout the study for culture and antimicrobial susceptibility testing. Results of reading were interpreted according to the diameters given in the most recent CLSI (Clinical Laboratory Standard Institute) documents (CLSI, 2016).

For Gram-positive organisms, the antibiotics to be tested and reported were chosen from the following (depending on the organism isolated): Penicillin (10 units), Gentamicin (10mcg) and high level (120mcg), Cefoxitin (30µg), Norfloxacin (5 µg), Vancomycin (30µg/MIC), Linezolid (30 µg), and Co-trimoxazole (1.25/23.75 µg), Nitrofurantoin (300µg). For gram negative organisms the antibiotics for respective organisms were chosen from the following: Gentamicin (10 µg), Piperacillin+Tazobactam (100/10µg), Polymyxin B (300 units), Colistin (10 µg), Norfloxacin (10 µg), Nitrofurantoin(300 µg), Ceftazidime (30 µg), imipenem(10 µg) and Cotrimoxazole (1.25/23.75 µg).

Results and Discussion

Out of the 121 urine samples received in microbiology laboratory, 42(34.71%) yielded positive culture while 20(16.52%) urine samples showed insignificant growth, 49(40.49%) cases had no growth and 10(8.26%) urine samples showed collection contaminants. Out of 42 bacterial isolates, 39 were Gram negative bacilli (GNB) and 03 Were Gram positive cocci (GPC)

Overall *E. coli* 16 (38.09%) was the predominant pathogen followed by *Pseudomonas aeruginosa* 08 (19.04%),

Klebsiella pneumoniae 05(11.9%), *Candida* species 05(11.9 %), *Acinetobacter species* 04 (9.52%), *Staphylococcus aureus* 02 (4.76%), *Enterobacter species* 01 (2.3%) and *Enterococcus species* 01 (2.3%) which correlates with the results of study conducted by Ramdhan Eldomany *et al.*, (2011).

Amongst the cancer patients 38% patient showed *E. coli* as commonest isolate. Out of 16 *E. coli* isolates, 12 (75%) were Extended Spectrum β lactamase (ESBL) producer which were 91% sensitive to imipenem and remaining four were non ESBLs.

Table.1 Bacterial profile of urine culture

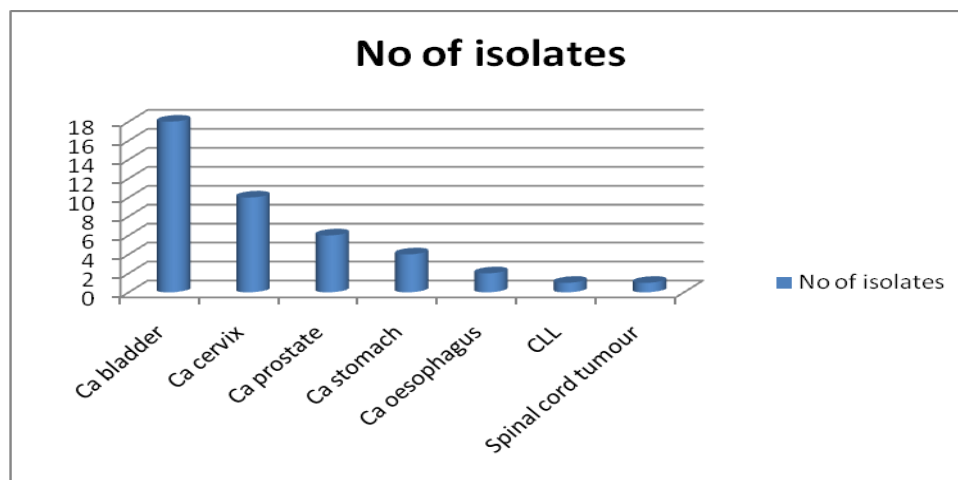
Sr. No	Organism	Number	Percentage
1.	<i>E.coli</i>	16	38.09
2.	<i>Pseudomonas aeruginosa</i>	08	19.04
3.	<i>Klebsiella pneumonia</i>	05	11.9
4.	<i>Candida species</i>	05	11.9
5.	<i>Acinetobacter species</i>	04	9.52
6.	<i>Staphylococcus aureus</i>	02	4.76
7.	<i>Enterobacter species</i>	01	2.3
8.	<i>Enterococcus species</i>	01	2.3

Table.2 Percentage of antimicrobial resistance in Gram negative microorganisms

Antibiotics	<i>E. coli</i>	<i>Pseudomonas aeruginosa</i>	<i>Klebsiella pneumonia</i>	<i>Acinetobacter species</i>	<i>Enterobacter species</i>
Ceftazidime	80	75	82	92	71
Colistin	0	0	0	0	0
Cotrimoxazole	83	NA	76	NA	80
Gentamicin	50	70	75	71	55
Imipenem	09	10	08	12	07
Nitrofurantoin	15	NA	40	40	12
Norfloxacin	90	91	95	96	94
Piperacillin +Tazobactam	20	25	30	34	25
Polymyxin B	0	0	0	0	0

NA: Not Applied

Fig.1 Cancer type in patients (N=42)



Susceptibility to carbapenem and Piperacillin+Tazobactam of *E. coli* was 90% and 73% respectively. While newer polypeptides like Polymyxin B and colistin were found to be effective against all gram negative bacteria. Refer Table 2 for detailed antibiogram of GNB.

Urinary tract infection (UTI) is the most common bacterial infection. The role of urinary tract infection in cancer patient's etiology is not well established. To gain more insight on the role of urinary tract infection in cancer patient's risk, this study analyzed data from one hundred and twenty one cancer patients.

In this study the culture of urine taken from cancer patients, 34.7% were positive. This result agrees with the study conducted by Parsad *et al.*, 1995 which revealed an incidence 33.3%. The same finding was reported by (Kouskouni *et al.*, 2005) who showed that the urinary tract infections in patients with malignancies were 33.4%

In our study, *E. coli* 16 (38.09%) was the predominant pathogen followed by *pseudomonas aeruginosa* 08(19.04%), *Klebsiella pneumoniae* 05 (11.9%), *Candida* species 05 (11.9%), *Acinetobacter* species 04 (9.52%), *Staphylococcus aureus* 02 (4.76%), *Enterobacter* species 01 (2.3%) and

Enterococcus species 01 (2.3%). This study correlates with Parikh and Bhat (2015) who showed that *E. coli* 40% followed by *Klebsiella pneumoniae* 25%, *Pseudomonas aeruginosa* 11%, *Enterococcus* species 11% and *Proteus mirabilis* 5%.

Antibiotic resistance is a major clinical problem in treating UTI in cancer patients caused by different bacterial pathogens. In our present study most of *E. coli* were ESBL producers and multidrug resistant which correlates with Monali Mishra *et al.*, (2013). In our study, Polymyxin B and colistin were found to be effective for MDR (Multidrug resistant) gram negative infections in patients with limited therapeutic options by considering its toxicity. These findings matched with other studies (Pastewski *et al.*, 2008; Ray *et al.*, 2007). In our study, UTI due to *Acinetobacter* showed high degree of resistance to commonly used antibiotics suggesting testing for newer drugs which correlates with study conducted by Gupta *et al.*, (2002). In conclusion, urinary tract infection in cancer patients is a common problem. This study suggests urine culture is very important for cancer patients specially those receiving cancer treatments and to control drug resistance, the use of antibiotics should be restricted and to be given only after doing culture and sensitivity test.

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How to cite this article:

Mukta Khaparkuntikar, Nazneen Siddiqui and Prasanna Bhirud. 2017. Urinary Tract Infection in Cancer Patients at Government Cancer Hospital Aurangabad, India. *Int.J.Curr.Microbiol.App.Sci.* 6(5): 2259-2263. doi: <https://doi.org/10.20546/ijcmas.2017.605.251>