PRESCRIBING AND SENSITIVITY PATTERNS OF ANTIMICROBIALS IN UNCOMPLICATED URINARY TRACT INFECTIONS IN FEMALES

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ABSTRACT

The overuse and misuse of antibiotics have been related to growing emergence of bacterial resistance worldwide. The aim of this present study was to detect the causative agents of urinary tract infection, assess the pattern of antimicrobial prescription along with the antimicrobial sensitivity pattern. Patient information was obtained by interviewing the patients and through their medical record files. Antibiotic sensitivity testing was performed by disc diffusion. The study showed that UTI was mostly prevalent in females of age group 20-30. *Escherichia coli* were the predominant (64.4%) bacterial pathogen followed by *Klebsiella* species (13.3%), *Pseudomonas* species (3.8%) and others. Most of the strains of *E. coli* were resistant to cephalexin whereas sensitive to cefpodoxime, amikacin, gentamycin and nitrofurantoin. Most of the urinary isolates showed high degree of resistance to cephalexin, norfloxacine and nalidixic acid. A single antibiotic was most commonly prescribed in both the hospitals, however, more than three antibiotics were also found to be prescribed. This study revealed that *E. coli* was the predominant bacterial pathogen of uncomplicated UTIs in both hospitals. It also demonstrated an increasing resistance to cephalexin, norfloxacine and nalidixic acid. Thus, formulation of a policy for hospital antibiotic use is a must for proper use of antibiotics and to ensure safe and efficient treatment of UTIs.

Keywords: UTI, *Escherichia coli*, cephalexin, prescribing, sensitivity

GENERAL INTRODUCTION

Urinary Tract Infections (UTI) is a heterogeneous disease, which can be divided into several types of infection, such as acute, uncomplicated bacterial pyelonephritis, complicated UTI, recurrent cystitis and asymptomatic bacteruria (Akortha and Ibadin, 2010). Urinary tract infection (UTI) is defined as significant bacteriuria in the presence of symptoms.

The urinary tract is normally sterile. Uncomplicated UTI involves the urinary bladder in a host without underlying renal or neurologic disease. The clinical entity is termed cystitis and represents bladder mucosal invasion, most often by enteric coliform bacteria (*e.g.*Escherichia coli) that inhabit the perirethral vaginal introitus and ascend into the bladder via the urethra.

Sexual intercourse may promote this migration, and cystitis is common in otherwise healthy young women. Urine is generally a good culture medium; factors unfavorable for bacterial growth include a low pH (5.5 or less), a high concentration of urea, and the presence of organic acids derived from a diet that includes fruits and protein.

It is the most common infection experienced by both male and female particularly responsible for discomfort in elderly patients, thus representing a risk of bacteremia, septic shock, respiratory distress syndrome and death (Gradvohl et al., 2005). Infection of adjacent structures such as prostate and epididymis is also included in this entity. Infection of urinary tract is amongst the most common bacterial infections that prompt patients to seek medical advice second only to infection of respiratory tract (Jha and Bapat, 2005). The geographical distribution of UTI amongst the Nepalese population is 0.57% in the mountains and 0.45% is estimated to be in planes.

Prescribing patterns

Prescribing is the act of determining which drug for which patient at an appropriate dosage regimen and with optimal duration of treatment. This is dynamic, highly individualized and involved medical, social and marketing forces. Inappropriate prescribing including supraoptimal or suboptimal choice of medication, higher or lower drug dosage, inappropriate duration, therapeutic duplication, potentially dangerous drug-drug interaction, prescribe an expensive drug when a cheaper and equally effective agent is available, are important health care problem.

Factors influencing prescribing patterns:

System factors: System or exogenous or structural environment factors which affect prescribing pattern are drug policies, hospital formularies, practice organization, influence of pharmaceutical companies, fragmentation of care, overabundance of drug therapy option and information quality.

Prescriber factors: Differences in characteristics of prescriber knowledge ability and experience in the field of specialty would affect the choice of drug to be used. Internal factors that would affect prescribing patterns include an inadequate training or practices, outdated medical knowledge, lack of continuing education, inadequate drug information especially its effectiveness or adverse reaction.
and drug cost, forgetfulness and temptation to the offer from pharmaceutical industries.

Patients or family factors: Patients or families who have a different backgrounds or cultural belief demand for drug or treatment which may be inappropriate. For example, patients demand an injected drug instead of an oral drug because they believed that it is more potent than the oral one and the disease will be cured more rapidly.

UTI are often treated with different broad-spectrum antimicrobials suspecting infection with resistant organisms though a narrow spectrum of activity may be appropriate. In general, β-lactam antibiotics, trimethoprim-sulphamethoxazole (TMP-SMX) and fluoroquinolones are used most often in uncomplicated UTI. In the case of a high-risk patient profile due to complicated factors, quinolones might be added to the armamentarium of treatment. However, the antibiotic of choice is currently highly dependent on the local situation with regard to the antibiotic resistance of the pathogenic bacteria (Matute AJ et al., 2004). However, the extensive use of antimicrobial agents has invariably resulted in the development of antimicrobial resistance, which, in recent years, has become a major problem worldwide (Dimitrov et al., 2004). In UTI, antimicrobial therapy is initiated even before the result of urine culture is available. Hence, there exists a great need for antimicrobial resistance surveillance at local, national and international level from time to time (Khan and Zaman, 2006).

METHODS

Collection of suspected prescription
The prescription of female patient visiting the hospital with sign and symptoms of UTI were enrolled in our study for the prescribing and sensitivity pattern (Huang and Stafford, 2002).

Inclusion criteria
The female patient visiting the hospital with the following sign and symptoms were enrolled in our study.
- Flank pain, Urine urgency, Urine frequency, Fever, Rigors, Burning micturition, Itching vulva, History of Discharge, Painful intercourse, Foul odor.

Exclusion criteria
Severe anemia (hemoglobin <7 gm/dl), Presence of heart disease with signs of cardiac failure, severe renal and hepatic diseases, documented tuberculosis with ongoing treatment, Associated other severe diseases that require special care or surgical intervention, Pregnancy and Diabetes.

Total number of enrollment
Total no of patients: 175
Minimum 50 patients suffering from UTI from each hospital.

Noting of patient history and empirical treatment therapy
Performa was filled up with patient name, age, address. Along with them, patient’s history of illness with empirical treatment and investigative suggestions were also noted.

Collection of urine sample
Freshly voided mid-stream urine with labia separated of suspected females was collected into a wide mouthed sterile container. Once collected, the sample was transferred to the laboratory without delay within 30 min. If a delay of more than 1-2 hours was to be made then it was stored in a refrigerator at 4°C or by collection and transportation in a container with boric acid at bacteriostatic concentration of 1.8% (Collee et al., 1996).

Preparation of culture media
Different culture media were prepared which were used according to standard method.

Culture and isolation of pathogens
Clean catch midstream collected urine samples was inoculated on Mac-Conkey and Blood Agar media using calibrated platinum loop following standard bacteriological technique. The primary inoculum was made by a loop and the material was spread into four quadrants of the plate. A platinum or nichrome wire (24 S.W.G. size) loop of 2-4 mm diameter with 2 to 3 inches long wire was first sterilized in the Bunsen flame and cooled by touching an uninoculated part of the medium. Then, a loopful of specimen was gently smeared onto the surface of a well dried plate of medium near the peripheral area. The inoculum was then thinly spread in parallel lines in different segments of the plate. The loop was sterilized between different sets of streaks. Finally, the plate was incubated at 37°C for overnight. Pure bacteriocolonies counting 100,000 or more was considered as significant and were subjected to identification based on colony characters and biochemical tests (Uwaezuoke and Ogbulie, 2006).

Bacterial identification
Colonial Morphology: The appearance of bacterial colony on the surface of a solid medium was used for the primary identification.

Smears prepared from the bacterial colony were examined by staining methods. Gram's staining divide bacteria into Gram-positive and Gram-negative. It is the most widely used stain in medical bacteriology.

Gram-positive bacteria resist decolourisation and stain violet while Gram-negative bacteria and other cells (pus cells) are decolourised and stain pink with counterstain.

Biochemical test
Indole Test
Indole is one of the degradation products of amino acid metabolism. It is useful in identifying E. coli. This test was used to determine whether bacteria possess the enzyme tryptophanase which will produce the byproduct indole from the catabolism of tryptophan. Kovac’s reagent was used after inoculation and incubation to read the results Kovac’s reagent contains HCl and dimethylaninobenzaldehyde (DMABA) dissolved in amyl alcohol, which forms a layer on top of the inoculating medium which causes the red color to be very easily visible and distinguishable. The alcohol in Kovac’s reagent floats on the media and another chemical in the reagent react with indole to form a red color in the alcohol layer. A positive test produced pink/purple colour at interface and the organism was E. coli and a negative test remained colorless.
Methyl Red/Voges Proskauer Broth

Voges-Proskauer Test

Some microbes do not produce stable acids from glucose fermentation but instead produce 2,3 butanediol from glucose breakdown and in the process an intermediate chemical acetoin is produced. Two reagents Barritt’s a (alpha-naphthol) and Barritt’s B (KOH) was added to a 48 hour culture of the MRVP broth. A pipette was taken to aliquot out a small amount of the broth to do the VP test and the broth was returned for further incubation for the MR test if necessary. A wine red color change with the addition of Barritt’s reagents A and B is a positive test detecting the presence of acetoin and therefore 2, 3 butanediol; a brown or copper color is negative.

Methyl Red Test

The methyl red test is used to determine organisms that ferment glucose to a stable acid end product in a great degree, lowering the pH of the system despite the presence of buffer. Media for the methyl red test (MRVP media) was prepared and contained peptone, glucose, and a phosphate buffer. Broth was inoculated and incubated at 30°C for five days to allow stable acids to be produced. At the end of the fifth day, methyl red indicator was added. Methyl red indicator is red at pH less than 4.4 and yellow at pH above 6.0, so a red result was labeled positive, and a yellow result negative.

Citrate Utilization Test

Simmon’s Citrate agar is used to determine if an organism can use citrate as its only carbon source using the enzyme citrase (also contains ammonia as the only nitrogen source). Citrate utilization is an aerobic process and a slant tube is used to increase exposure of bacterial growth to the air; inoculate the slant, pH indicator is Bromthymol blue, which is green at neutral pH but turns Prussian blue at pH levels above 7.6

Triple Sugar Iron Agar

TSI Agar slants contain 2% polypeptone, 1% lactose, 1% sucrose, 1% glucose, phenol red pH indicator, and ferric ammonium citrate. Utilization of sugars proceeds in much the same way as in sugar broth tests, with acid production changing a pH indicator. Specifically, fermentation of lactose/sucrose and glucose causes the entire tube to be yellow. Fermentation of glucose alone causes the butt of the culture to be yellow, but the shallow slant portion turns red as glucose is oxidatively exhausted and peptone is metabolized, producing NH₃ resulting in an alkaline pH. Gas production during the utilization of sugar is indicated by fissures or pockets in the slant. Some bacteria also produce hydrogen sulfide (H₂S) by reducing thiosulfate in the medium or breaking down cysteine in the peptone. Ferric ammonium citrate reacts with hydrogen sulfide to produce a black precipitate in the butt of the agar. The slants were made from commercially prepared TSI mix and poured to allow the butt of the agar to be four centimeters deep, and the slants were inoculated. The slants were incubated for twenty-four hours at 37°C and tests were read for all three sugars as well as gas and H₂S production.

Preparation of MacFarland turbidity standard

0.5 ml of 0.048 M Barium chloride was added in 66.5 ml of 0.36 N sulfuric acid and 5 ml of it was aliquoted into screw-capped tubes of same size and was stored in dark at room temperature (Shaikh et al., 2005)¹⁰.

Preparation of inoculum

Four or five well isolated colonies of same morphological type were touched with a sterile wire loop, suspended into tubes containing 5 ml of MHB. The medium was then incubated at 35°C for 2-8 hours until the turbidity reached or exceeded that of 0.5 MacFarland standards (already prepared). If the suspension was exceeded, it was diluted with broth and was visually comparable to the 0.5 MacFarland standards (Shaikh et al., 2005)¹⁰.

Inoculation of MHA plates

A sterile swab was dipped into the broth suspension of the organisms within 15 minutes of standardization. Excess inoculum was removed by rotating the swab several times against the wall of the tube above the fluid level. The entire surface of an MHA plate was then streaked evenly in three/two directions approximately 60 degrees from each other (Shaikh et al., 2005)¹⁰.

Antimicrobial sensitivity test

Antimicrobial sensitivity test was performed by disc diffusion method (Kirby-Bauer’s technique) using commercially available disc. The antimicrobial impregnated discs was placed with sterile forceps on the Muller Hinton Agar surface such that each disc was at least 24 mm from the other disc avoiding overlap during incubation. Then, it was incubated at 37°C for 18-24 hours. These test discs will be of the antimicrobials observed in prescription pattern (Khan and Zaman, 2006)⁹. At the end of incubation period, the diameter of zones of inhibition around each disc was measured with a Vernier caliper on the back of the plate, with reflected light against a dark non-reflective background. The zone of diameter for each antimicrobial agent was then interpreted as resistant/intermittent/sensitive by comparing with the standard given by the HI Media Laboratories Pvt. Ltd. (Shaikh et al., 2005)¹⁰.

Noting of isolated bacteria and sensitivity pattern

The bacteria responsible for the pathogenesis were noted after isolation of bacteria with culture and biochemical tests. Along with it, the sensitivity pattern of the antimicrobials was also noted down.

Noting of final diagnosis and prescribing pattern

The final diagnosis made after seeing the report by the physicians and the drug prescribed after it was noted down.

Data analysis

Statistical Package for Social Sciences for windows (SPSS), version 11.5 was used for statistical analysis.

RESULTS

A total of 175 urine sample were collected, out of these 104 patients’ urine were found to have significant bacterial growth. Total six species of bacteria were isolated viz. E.coli 67(64.4%), Proteus species 3(2.9%), Klebsiella species 14(13.3%), Staphylococcus aureus 11(10.6%), Citrobacter species 5(4.8%) and Pseudomonas aeruginosa 4(3.8%).
Table 1: Respondence of UTI in relation to age distribution of female patients

<table>
<thead>
<tr>
<th>Age group</th>
<th>Hospital A</th>
<th>Hospital B</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 10</td>
<td>20.0%</td>
<td>30.9%</td>
<td>26.9%</td>
</tr>
<tr>
<td>10 - 20</td>
<td>13.8%</td>
<td>18.2%</td>
<td>16.6%</td>
</tr>
<tr>
<td>20 - 30</td>
<td>36.9%</td>
<td>24.5%</td>
<td>29.1%</td>
</tr>
<tr>
<td>30 - 40</td>
<td>10.8%</td>
<td>15.5%</td>
<td>13.7%</td>
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<tr>
<td>40 - 50</td>
<td>9.2%</td>
<td>5.5%</td>
<td>6.9%</td>
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<tr>
<td>50 - 60</td>
<td>3.1%</td>
<td>2.7%</td>
<td>2.9%</td>
</tr>
<tr>
<td>60 - 70</td>
<td>3.1%</td>
<td>2.7%</td>
<td>2.9%</td>
</tr>
<tr>
<td>&gt; 70</td>
<td>3.1%</td>
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<tr>
<td>Total</td>
<td>100.0%</td>
<td>100.0%</td>
<td>100.0%</td>
</tr>
</tbody>
</table>

Figure 1: Shows the incidence of UTI by occupational groups

Figure 2: Bacterial growth positive rate in urine samples collected.
Figure 3. Frequency of different pathogens isolated from urine samples of females with uncomplicated UTI (N = 104) in both hospitals.

Table 2: Overall sensitivity Pattern of antimicrobials in hospital A and hospital B

<table>
<thead>
<tr>
<th>Hospital A</th>
<th>Cefpodoxime (100%) &gt; Alizarin (98.0%) &gt; Nitrofurantoin (95.9%) &gt; Gentamycin (94.1%) &gt; Azithromycin (88.1%) &gt; Cotrimoxazole (60.9%) &gt; Ciprofloxacin (58.1%) &gt; Ofloxacin (53.3%) &gt; Cefixime (44.7%) = Norfloxacin (44.7%) &gt; Nalidixic Acid (35.7%) &gt; Amoxicillin (33.3%) &gt; Cephalexim (25.5%).</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hospital B</td>
<td>Azithromycin (100%) &gt; Amikacin (80.3%) &gt; Gentamycin (72.3%) &gt; Ciprofloxacin (56.8%) &gt; Nitrofurantoin (54.2%) &gt; Levofloxacin (53.3%) &gt; Amoxicillin (53.1%) &gt; Cotrimoxazole (48.5%) &gt; Cefixime (47.1%) &gt; Ofloxacin (45.7%) &gt; Cefpodoxime (45.5%) &gt; Nalidixic Acid (40.4%) &gt; Norfloxacin (31.0%) &gt; Cephalexime (20.8%) &gt; Other antibiotics (11.1%).</td>
</tr>
</tbody>
</table>
Table 3: Frequency of antibiotics prescribed in respective hospitals.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>HOSPITAL</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>A</td>
<td>B</td>
</tr>
<tr>
<td>No. of antibiotics</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 antibiotic</td>
<td>75.0%</td>
<td>77.2%</td>
</tr>
<tr>
<td>2 antibiotics</td>
<td>25.0%</td>
<td>18.8%</td>
</tr>
<tr>
<td>3 antibiotics</td>
<td>3.0%</td>
<td>2.0%</td>
</tr>
<tr>
<td>&gt; 3 antibiotics</td>
<td>1.0%</td>
<td>0.7%</td>
</tr>
<tr>
<td>Total</td>
<td>100.0%</td>
<td>100.0%</td>
</tr>
</tbody>
</table>

Table 4: Frequency of antibiotics prescribed in hospital A and hospital B.

<table>
<thead>
<tr>
<th>No. of antibiotics</th>
<th>HOSPITAL</th>
<th>Total</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>A</td>
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<td>1 antibiotic</td>
<td>75.0%</td>
<td>77.2%</td>
</tr>
<tr>
<td>2 antibiotics</td>
<td>25.0%</td>
<td>18.8%</td>
</tr>
<tr>
<td>3 antibiotics</td>
<td>3.0%</td>
<td>2.0%</td>
</tr>
<tr>
<td>&gt; 3 antibiotics</td>
<td>1.0%</td>
<td>0.7%</td>
</tr>
<tr>
<td>Total</td>
<td>100.0%</td>
<td>100.0%</td>
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</tbody>
</table>
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Conflicts of interest:

It’s me myself corresponding author of this article and assure you that nobody have conflicts of interest if we publish this article in your journal: JOURNAL OF DRUG DELIVERY AND THERAPEUTICS (JDDT).

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