

# Research Journal of Pharmaceutical, Biological and Chemical Sciences

## Evaluation of Parenteral Antibiotic Utilization in Medical Inpatients

NA Aleykutty<sup>1</sup>, Remya Raju<sup>2</sup>, Santhosh M Mathews<sup>2\*</sup>

<sup>1</sup>Pushpagiri College of Pharmacy, Thiruvall, Kerala, India.

<sup>2</sup>Nazareth College of Pharmacy, Othara, Thiruvalla, Kerala, India.

### ABSTRACT

The study was conducted to evaluate the effectiveness of regulations on the utilization of antibiotics administered by the parenteral route. It was performed by evaluating the total number of doses of antibiotics consumed before and after imposing regulations. All the hospitalized patients in the medical ward were subjected for evaluation. The uses of all antibiotics were regulated using drug formulary and restrictions in the prescribing pattern. It was observed that the doses of antibiotics consumed by the parenteral route were altered significantly after the regulation.

**Keywords:** Antibiotics, Parenteral administration, Regulations, Doses consumed.

*\*Corresponding author*

Email: santhoshmm@hotmail.com

## INTRODUCTION

Antibiotics are most widely prescribed in the drug therapy and the complexity of their overuse is a global phenomenon [1-3]. Various studies showed that the percentage of prescriptions containing antimicrobials varies from 22% to 67 % [4-7].

It was reported that the injection was the most frequently used method to administer antibiotics. It was reported that 33% of intravenous antibiotics were prescribed empirically without documented evidence of infection [8]. Such a wide range of use have lead to an increased risk of side effects, drug toxicity, and makes the therapy more expensive and in addition, increases antimicrobial resistances [9]. To ensure that antibiotics are used appropriately and rationally antibiotic utilization has to be reviewed regularly [10].

The optimization of antibiotic utilization at its most basic level is the appropriate use of antibiotics. Various antibiotic utilization strategies have been implemented to improve antibiotic prescribing in many hospitals [11-13]. These include formalized antibiotic guidelines, antibiotic control programs, restrictive antimicrobial prescribing forms, provision of a computerized information system to guide antimicrobial selection, short-listed antibiotics in hospital formularies, etc [14, 15]. Antibiotic guidelines have prove to be simple, yet effective intervention while encouraging appropriate choice of antibiotic therapies and recommending a timely switch from intravenous to oral therapy. But it is reported a single intervention alone is not sufficient enough to make the antibiotic therapy more effective. Therefore a multidisciplinary approach is more appropriated [16-18].

## MATERIALS AND METHODS

The study was conducted in the medical ward of a superspeciality multidisciplinary teaching hospital for a period of two years. The individual medical records of all in patients admitted in this ward were studied. Regulations were imposed on the antibiotic prescription pattern and adherence to hospital formulary was strictly implemented. Doses of antibiotics consumed parentally were recorded before and after the regulations. All the data were analyzed using chi-square test at 5% level of significance. All the differences were tested at  $p < 0.05$  (two-tailed).

## RESULTS AND DISCUSSION

The study showed that the number of patients received antibiotics before and after the regulations were 2375 and 2425 respectively. Out of them 1364 received antibiotics parentally before the regulation whereas 1093 received after the regulation.

The total doses of different class of antibiotics consumed in parentral route before and after the regulations were summarized in Table-1. The total numbers of doses of antibiotics consumed parentally were 38181 and 26223 before and after the intervention respectively.

The number of patients received antibiotics parentally were reduced by 21% . The study had shown a significant reduction in the total number of doses of antibiotics (31%) used parentally. This may lead to decrease the risk of drug interactions, over use of drugs and make the treatment more economical.

During the pre-regulation period, third generations cephalosporins were the most commonly prescribed antibiotics. After the restriction, this was decreased by 11%. The prescription of second generation cephalosporins also reduced by 52%. The reduction in the use of second and third generation cephalosporins led to an increase in the use of first generation cephalosporins by 43%.

The broad spectrum antibiotic used during the period of study was Chloramphenicol which itself reduced by 70%.The use of carbapenams - the most expensive antibiotics used parentally- were decreased by 18%. Use of Lincosamides was reduced by 43%. Lincosamide and its combination with third generation cephalosporin were also decreased. The use of drugs belonging to oxazolidinone category was reduced to 37% .A reduction in the consumption of Sulphonamides, Penicillin and Imidazole derivatives were observed after the interventions justifying the rational use of antibiotics by parental route.

**Table – 1. Summary of the Number of Doses of Antibiotics Consumed (Parental Route)**

| Antibiotics  | Use of antibiotics         |                           | p-value* |
|--|----------------------------|---------------------------|----------|
|  | Before Regulation<br>n (%) | After Regulation<br>n (%) |          |
| <b>Total Number of Doses of Antibiotics Consumed (n)</b> | 38181                      | 26223                     |          |
| <b>Aminoglycosides</b>                                   | 2230(5.84)                 | 1662(6.34)                | <.0001   |
| <b>Broad Spectrum Antibiotics</b>                        | 75(0.20)                   | 16(0.06)                  | <.0001   |
| <b>Extended Spectrum Penicillins</b>                     | 3447(9.03)                 | 2866(10.93)               | <.0001   |
| <b>Carbapenams</b>                                       | 1538(4.03)                 | 870(3.32)                 | <.0001   |
| <b>Cephalosporins-First Generation</b>                   | 58(0.15)                   | 208(0.79)                 | <.0001   |
| <b>Cephalosporins-Second Generation</b>                  | 2101(5.50)                 | 695(2.65)                 | <.0001   |
| <b>Cephalosporins-Third Generation</b>                   | 19423(50.87)               | 11934(45.51)              | <.0001   |
| <b>Cephalosporins-Fourth Generation</b>                  | 241(0.63)                  | 66(0.25)                  | <.0001   |
| <b>Lincosamide + Cephalosporin-Third Generation</b>      | 148(0.39)                  | 95(0.36)                  | <.0001   |
| <b>Lincosamides</b>                                      | 1842(4.82)                 | 722(2.75)                 | <.0001   |
| <b>Macrolides</b>  | 86(0.23)                   | 207(0.79)                 | <.0001   |
| <b>Oxazolidinones</b>                                    | 116(0.30)                  | 50(0.19)                  | <.0001   |
| <b>Penicillins</b>                                       | 2135(5.59)                 | 1990(7.59)                | <.0001   |
| <b>Uredopenicillins</b>                                  | 1848(4.84)                 | 1873(7.14)                | <.001    |
| <b>Fluroquinolones- First Generation</b>                 | 1444(3.78)                 | 1946(7.42)                | <.0001   |
| <b>Fluroquinolones-Second Generation</b>                 | 788(2.06)                  | 613(2.34)                 | <.0001   |
| <b>Glycopeptides</b>                                     | 603(1.58)                  | 383(1.46)                 | 0.0005   |
| <b>Imidazole Derivatives</b>                             | 58(0.15)                   | 27(0.10)                  | 0.4395   |

\*p-value calculated using Chi-square test (two tailed,  $\alpha = 0.05$ ).

The use of Uredopenicillins was increased by 48% as it was considered as the drug of choice for the treatment of advanced pneumonia. Use of first generation fluoroquinolones

were increased whereas the consumption of the second generation fluoroquinolones did not alter much.

The use of extended spectrum antibiotics was increased in the post regulatory period (21%). Amoxicillin was the most commonly used drug of choice during the period of study for upper respiratory infections particularly for sinusitis. But increase in the resistance against this drug caused an increase in the use of amoxicillin combined with clavulanate. Use of macrolid antibiotics increased mainly due to the increase in the use of Azithromycin prescribed for the treatment of bronchitis. These increases in the use of extended spectrum penicillins and macrolids showed that all these antibiotics were found to be sensitive enough to treat the infections in place of drugs belonging to higher classes.

### CONCLUSION

The evaluation of restricted use of antibiotics utilization showed that the effective use of parenteral antibiotics was improved after regulation. The study emphasizes that strict regulation on the use of antibiotics is essential to promote rational use of antibiotics by parenteral route.

### REFERENCES

- [1] David L Paterson. Clinical Infectious Diseases 2006; 42: 590-595,
- [2] Viller EP, Co NM, Co BG, et al. Santo Tomas J Med 1994; 43(1):44-57.
- [3] Ghosh R, Neogi JN, Srivastava BS, et al. J Nep Med Assoc 2003; 42:346-49.
- [4] Shrishyla MV, Naga Rani MA, Venkararaman BV. Indian J Pharmacol 1994; 26: 282-7.
- [5] Don AB, Robert JA. Pharmacy Practice 2008; 3: 3-13.
- [6] Ghosh R, Neogi JN, Srivastava BS, et al. J Nep Med Assoc 2003; 42: 346-49.
- [7] Pradhan SC, Shewade DG, Tekur U, et al. Int J Ther Toxicol 1990; 28:339-43.
- [8] Ahkee S, Barzallo M, Ramirez J. Infect Med 1996; 13:800-2,823.
- [9] Dickerson LM, Mainous AG, Carek PJ. Pharmacotherapy 2000; 20:711-23.
- [10] Gyssens IC, Blok WL, Van den Broek PJ, et al. Eur J Clin Microbiol Infect Dis 1997; 16: 904-12
- [11] Berild D, Ringertz SH, Lelek M, et al. Scand J Infect Dis 2001; 1:63-7.
- [12] Ramirez JA, Vargas S, Ritter GW, et al. Archives of Internal Medicine 1999; 20: 2449-54.
- [13] Martinez MJ, Freire A, Castro I, et al. Pharm World Sci 2000; 2: 53-8.
- [14] Sevinc F, Prins JM, Koopmans RP, et al. J Antimicrob Chemother 1999; 4: 601-6.
- [15] Gould IM. J Antimicrob Chemother 1999; 4:459-65.
- [16] Gentry CA, Greenfield RA, Slater LN, et al. Am J Health Syst Pharm 2000; 3: 268-74.
- [17] South M, Royle J, Starr M. Med J Aust 2003; 5: 207-9.
- [18] Ruttimann S, Keck B, Hartmeier C, et al. Clin Infect Dis 2004; 3: 348-56.