Incidence of ventilator associated pneumonia

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ABSTRACT

Introduction: The development of VAP is also associated with greater hospital mortality rates and longer lengths of stay in Intensive Care Unit (ICU) and hospitals. A prospective study was conducted to find out the incidence of VAP and to identify the most prevalent pathogens causing VAP. Materials and Methods: The study was conducted during the period of September 2007 to March 2008 at ICU. The endotracheal aspirates were collected and transported to microbiology laboratory within 15 minutes, where they were processed and cultured as per the standard protocol and standard microscopic examinations were conducted. Statistical analysis was performed using SPSS 11.0 and Systat 8.0. Results: Of the 100 patients studied, 29 were found to have VAP. Among these patients, 32% were reported to have hypertension, 29% were reported to have diabetes and 12% had both diabetes and hypertension. Gram negative organisms were predominant among the isolates accounting for 89%. The rest were found to be gram positive organisms. Among gram negative organisms, Pseudomonas species, Klebsiella species and E.coli were responsible for highest number of VAP infections. Conclusion: The range of bacteria that cause VAP and their susceptibility patterns vary widely among hospitals and selection of initial antimicrobial therapy need to be tailored to each institution’s local patterns of antimicrobial resistance.

Key words: Intensive care unit; Ventilator associated pneumonia; Endotrachial aspirates.

INTRODUCTION

Incidence of Hospital acquired infection has drastically reduced with proper implementation of preventive guidelines in the hospital environment. However, several such (Nosocomial) infections are still encountered in various departments of a Hospital. Pneumonia results from microbial invasion of the normally sterile lower respiratory tract and lung parenchyma caused by a defect in host defenses, challenge by a particularly virulent microorganism, or an overwhelming inoculum.

Nosocomial pneumonia is the second most among them which is associated with the highest case/fatality ratio. Among different types of nosocomial infections, ventilator associated pneumonia (VAP) continues to complicate the course of 8-28% of patients receiving mechanical ventilation (MV). The mortality rate associated with VAP ranges from 24-50% and can escalate up to 76% based on specific settings and host-pathogen relationship. The development of VAP is also associated with greater hospital mortality rates and longer lengths of stay in Intensive Care Unit (ICU) and hospitals. Microorganisms responsible for VAP may differ according to the population of patients in the ICU, the duration of hospital and ICU stays, and the specific diagnostic method(s) used. The high rate of respiratory infections due to Gram negative bacilli in this setting has been repeatedly documented. Several studies have reported that more than 60% of VAP is caused by aerobic Gram negative bacilli. More recently, some investigators have reported that gram-positive bacteria have become increasingly more common in this setting with S.aureus being the predominant gram-positive isolate. Although the incidence of VAP and the contribution of various micro-organisms to this have been studied extensively around the world, no such work has been carried out.
in south India. A prospective study was conducted at Narayana Medical College and General Hospital to find out the incidence of VAP and to identify the most prevalent pathogens causing VAP.

MATERIALS AND METHODS

The study was conducted during the period of September 2007 to March 2008 at ICU of our Medical college attached superspeciality hospital. A total of 100 patients connected with mechanical ventilators for more than 48 hours were studied. A proforma was filled in detail for each patient with regard to his/her name, age, sex, IP no, ward/unit, Clinical diagnosis, indications on artificial ventilation, nature of sample collected for VAP, new symptoms after ventilator connected (after 48hrs) and empirical treatment given.

Sample collection:
The endotracheal aspirates were collected by using a sterile 12 gauge endotracheal suction catheter tube. Suction catheter is connected to the suction pump and passed through endotracheal tube. In case of purulent secretion the Endo Tracheal suction catheter tips were collected using sterile scissors into the sterile container (screw capped). In case of thick or dry secretions with clinical diagnosis of pneumonia, sterile normal saline was pushed into endotracheal tube and then suction was done for the aspirate. The samples were transported to microbiology laboratory within 15 minutes. Specimens were processed and cultured as per the standard protocol as soon as they were obtained and standard microscopic examinations were conducted. Antimicrobial susceptibility was analyzed using Mueller–Hinton agar (hi-media) using, Kirby-Bauer disk diffusion method. The isolated colony was sub cultured in peptone water and compared with 0.5 Mac. Farlands turbidity. Plates were inoculated with a sterile cotton swab dipped in the standardized suspension of organism.

After inoculum has dried, specified antibiotic discs were placed 2 cm apart from each other with forceps and was incubated for 16-18 hours at 37°C aerobically. The zone size was interpreted according to the reference charts provided by manufacturer. The statistical software namely SPSS 11.0 and Systat 8.0 were used for the analysis of the data and Microsoft Word and Excel have been used to generate graphs, tables etc.

RESULTS

Of the 100 patients studied, 29 were found to have VAP. Further among these patients, a higher percent of patients were reported to have early onset defined as development of VAP before 5 days of MV than late onset defined as development of VAP after 5 days of MV (Table 1).

Association of diabetes and hypertension
Among the patients studied, 32% of patients were reported to have hypertension, 29% were reported to have diabetes and 12% were reported to have both diabetes and hypertension. Further VAP was found to be higher among patients with hypertension (Fig.1).

Microorganisms isolated
The causative organisms were isolated and identified. Gram negative organisms were predominant among the isolates accounting for 89%. The rest were found to be gram positive organisms. Among gram negative organisms, Pseudomonas species, Klebsiella species and E.coli were responsible for highest number of VAP infections (Fig.2).

Table 1: Comparison of Early and Late onset VAP

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<td>Total number of cases studied</td>
<td>100</td>
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<tr>
<td>Number of VAP cases identified</td>
<td>29</td>
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<tr>
<td>Early onset (&lt;5days) cases</td>
<td>16(55.17%)</td>
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<tr>
<td>Late onset (&gt;5days) cases</td>
<td>13(44.82%)</td>
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Prolonged (more than 48 hours) MV is the most important factor associated with nosocomial pneumonia. However, VAP may occur within the first 48 hours after intubation. It is usual to distinguish early-onset VAP, which occurs during the first 5 days of MV, from late-onset VAP, which develops 5 or more days after initiation of MV.

The incidence of the current study was 29% which was compared with 28% of incidence in a study by Fagon JY, et al.[1]. The onset of VAP in the current study was 55.17% in early onset which was comparable with 45% of early onset in a study by Prod’hom G et al.[6]. Not only are the causative pathogens commonly different but the disease is usually less severe and the prognosis better in early-onset than late-onset VAP. Tullu MS, et al. reported that E.coli, Klebsiella species and Pseudomonas species were isolated as the commonest organism colonizing the endotracheal tube leading to ventilator associated pneumonia. The current study also reported similar trend in isolates responsible for VAP. The frequency of isolation of Pseudomonas species (27.58%), Acinetobacter (3.44%), Klebsiella species (27.58%), E.coli (20.68%), Citrobacter (6.89%), Proteus species (3.44%), S.aureus (6.89%) and Enterococci (3.44%) were found to be comparable with that of study done by Chastre J et al.[1] where the frequency of isolates of Pseudomonas species (24.4%), Acinetobacter (7.9%), Klebsiella species (15.6%), E.coli (24.1%) and Citrobacter.
National nosocomial infections surveillance (NNIS) report stated that *Pseudomonas aeruginosa* have become class I cephalosporinase producers and are resistant to piperacillin and ceftazidime. *Klebsiella species* and other *Enterobacteriaceae* strains are also increasingly being recognized as producers of transferable extended spectrum beta-lactamases, which confer resistance to third generation cephalosporins. However, in current study *Pseudomonas species* is sensitive to cefoperazone+sulbactum, piperacillin, carbenicillin and imipenem and most of the *Enterobacteriaceae* members are sensitive to amikacin, ciprofloxacin, cefepirazone+sulbactum. However, because the range of bacteria that cause VAP and their susceptibility patterns vary widely among hospitals in the same or different countries, selection of initial antimicrobial therapy need to be tailored to each institution’s local patterns of antimicrobial resistance.

**REFERENCES**


