



ISSN: 0975-833X

Available online at <http://www.journalcra.com>

INTERNATIONAL JOURNAL  
OF CURRENT RESEARCH

International Journal of Current Research  
Vol. 10, Issue, 12, pp.76488-76491, December, 2018

DOI: <https://doi.org/10.24941/ijcr.33479.12.2018>

## RESEARCH ARTICLE

# BACTERIOLOGICAL PROFILE AND ANTIBIOTIC SENSITIVITY PATTERN OF NEONATAL SEPTICAEMIA USING AUTOMATED BLOOD CULTURE AND VITEK-2 SYSTEMS IN A TERTIARY CARE HOSPITAL, BENGALURU

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### ARTICLE INFO

#### Article History:

Received 22<sup>nd</sup> September, 2018  
Received in revised form  
16<sup>th</sup> October, 2018  
Accepted 04<sup>th</sup> November, 2018  
Published online 31<sup>st</sup> December, 2018

#### Key Words:

Neonatal sepsis, Early onset sepsis(EOS), Late onset sepsis(LOS), BacT/ALERT and Vitek 2 systems.

### ABSTRACT

**Background:** Neonatal sepsis refers to systemic infection of the newborn. It is characterized by nonspecific symptoms, and documented by positive blood culture. An estimated 1.6 million deaths occur due to neonatal infections worldwide, 40% of them being limited to developing countries. The incidence of neonatal sepsis in our country is around 30 per 1000 live births. These children are more likely to have neuro developmental side effects. Therefore it is important to diagnose and treat the condition as early as possible. **Objective:** The aim of the present study is to determine the microbiological profile of neonates with sepsis and to determine their antibiotic susceptibility pattern using automated systems-BacT/ALERT and Vitek 2 systems. **Materials and Methods:** This study is a Retrospective study conducted between December 2017 and February 2018 at NICU, Vani Vilas hospital, BMC&RI, Bengaluru. All admitted neonates with clinical signs and symptoms of sepsis at the time of admission or who developed sepsis during their hospital stay were included in the study. Blood samples were collected aseptically from neonates suspected with neonatal sepsis. Samples were inoculated into pediatric automated blood culture bottle "pediatric" (yellow, 30 ml, BacT/ALERT PF). Cultures with positive results were streaked on blood and Mac Conkey agar. Identification and antibiotic sensitivity were done using Vitek-2 systems. **Results:** In this study blood samples were collected from 253 neonates admitted to NICU of Vanivilas hospital, 72 samples were found to be culture positive. The most common organism isolated from both Early onset sepsis(EOS) and Late onset sepsis(LOS) was *Klebsiellapneumoniae* 57% and 53% respectively. Most of the *Klebsiella* organisms were sensitive to Piperacillin-tazobactam, Amikacin and Imepenem. The next most common organism isolated was *Staphylococcus aureus* in EOS and *Enterococcus* species in LOS. The most important risk factors found were preterm and low birth weight, with respiratory distress being the most frequent presentation. **Conclusion:** Appropriate identification of sepsis source, prompt antibiotic prescription and aggressive management can effectively prevent adverse events following neonatal sepsis. There is requirement of simple and sustainable intervention to reduce the burden of multidrug resistant pathogens causing sepsis. The implementation of hand washing, barrier nursing and antibiotic stewardship policies is need of time.

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**Citation:** Dr. Manjula C., Dr. Jyoti S Kabbin and Dr. Ambica R. 2018. "Bacteriological profile and antibiotic sensitivity pattern of neonatal septicaemia using automated blood culture and vitek-2 systems in a tertiary care hospital, Bengaluru", *International Journal of Current Research*, 10, (12), 76488-76491.

## INTRODUCTION

Neonatal sepsis refers to systemic infection of the newborn. It is characterized by nonspecific symptoms, and documented by positive blood culture. The knowledge of common bacterial agents causing septicemia and their antibiogram is useful in deciding empirical antimicrobial therapy. The pattern of organisms causing sepsis differs from place to place and can change in the same place over a period of time (Renuka Anegundi, 2017).

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Early-onset neonatal sepsis(EOS) refers to the presence of a confirmed infection in the blood of patients younger than 72 hours of life, and late-onset neonatal sepsis (LOS) refers to the onset of such infection between 72 hours and 90 days old (Reyes, 2015). An estimated 1.6 million deaths occur due to neonatal infections worldwide, 40% of them being limited to developing countries. The incidence of neonatal sepsis in our country is around 30 per 1000 live births (Chacko et al., 2005). Infants surviving neonatal sepsis can have significant neurologic sequelae as a consequence of central nervous system involvement, septic shock or hypoxemia secondary to severe parenchymal lung disease. Early treatment with appropriate antibiotics would minimise the risk of severe

morbidity and mortality besides reducing the emergence of multidrug resistant organisms in intensive care units by rational antibiotic use (Viswanathan *et al.*, 2012). Routine use of broad spectrum antibiotics alters the microbial flora resulting in emergence of resistant organisms and fungal infections (Vinodkumar *et al.*, 2008). Therefore it is important to diagnose and treat the condition as early as possible. The spectrum of microbial etiology of neonatal sepsis varies from region to region and even varies in different hospitals of the same region.

### Objectives of the study

- To identify the microbiological profile and antibiotic sensitivity pattern of neonatal septicemia using automated blood culture systems.
- To assess various factors contributing to neonatal sepsis

### MATERIALS AND METHODS

This study is a Retrospective study conducted between December 2017 and February 2018 at NICU, Vani Vilas hospital, BMC&RI, Bengaluru. Ethical clearance for the study was obtained from the institutional ethical committee. All admitted neonates with clinical signs and symptoms of sepsis at the time of admission or who developed sepsis during their hospital stay were included in the study. Blood samples were collected aseptically from neonates suspected with neonatal sepsis. Samples were inoculated into paediatric automated blood culture bottle "paediatric" (yellow, 30 ml, BacT/ALERT PF).

Cultures which flagged positive in the automated blood culture system were streaked on Blood and Mac Conkey agar plates. Identification and antibiotic sensitivity was done using Vitek-2 systems (CLSI, 2016).

### RESULTS

In this study blood samples were collected from 253 neonates admitted to NICU of Vani Vilas hospital. 72(28%) blood samples were found to be culture positive. Out of 253 neonates, 156(62%) were male and 97(38%) were female babies. Early onset sepsis (EOS) was suspected in 190 neonates and Late onset sepsis (LOS) in 97 neonates. 53(28%) blood samples yielded positive cultures in EOS and 19(30%) in LOS. The most common organism isolated from both Early onset sepsis (EOS) and Late onset sepsis (LOS) was *Klebsiellapneumoniae* which was 57% and 53% respectively.

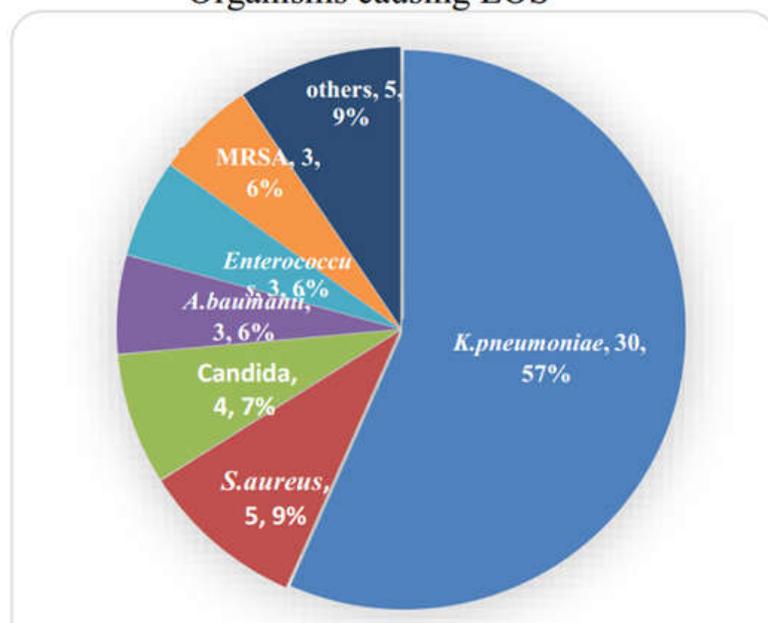
Most of the *Klebsiella* organisms were sensitive to Amikacin, Piperacillin-tazobactam, and Imepenem. The next most common organism isolated was *Staphylococcus aureus* in EOS and *Enterococcus* species in LOS. The most important risk factors found were preterm and low birth weight, with respiratory distress being the most frequent presentation.

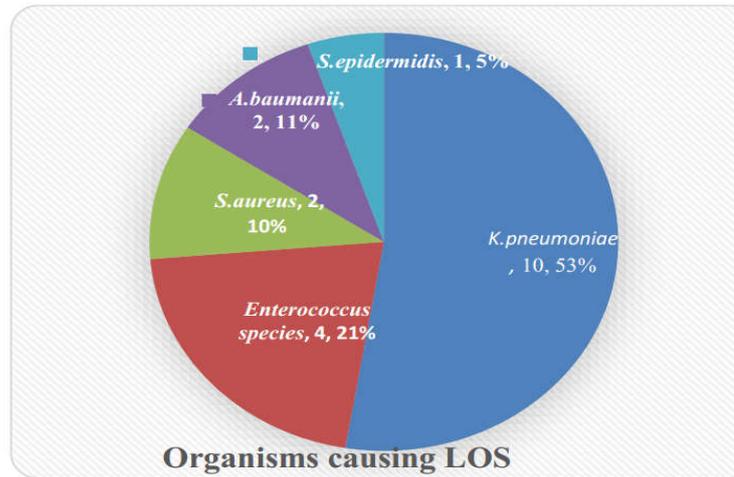
### DISCUSSION

In this study blood culture positivity was 28%. Culture positivity in EOS and LOS was 28% and 19.58% respectively. This is lower than in the studies of Renuka Anegundi *et al* and Chacko B *et al* where the culture positivity was 46.8% and 55.4%. *Klebsiellapneumoniae* was the most common organism isolated in our study similar to the study of Vishwanath R *et al*.

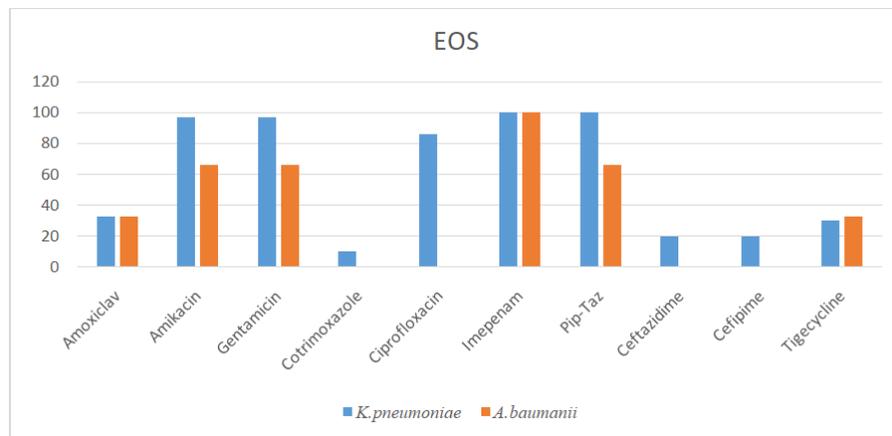
| Sl No | Studies                              | Percentage of Isolates | Most Common Organism         |
|-------|--------------------------------------|------------------------|------------------------------|
| 1.    | Renuka Anegundi <i>et al.</i> (2017) | 46.8%                  | MR CONS, <i>K.pneumoniae</i> |
| 2.    | Chacko <i>et al.</i> (2005)          | 41.7%                  | <i>P.aeruginosa</i>          |
| 3.    | Viswanathan <i>et al.</i> (2012)     | 46.3%                  | <i>K.pneumoniae</i>          |
| 4.    | Present study                        | 28%                    | <i>K.pneumoniae</i>          |

### Organisms causing EOS

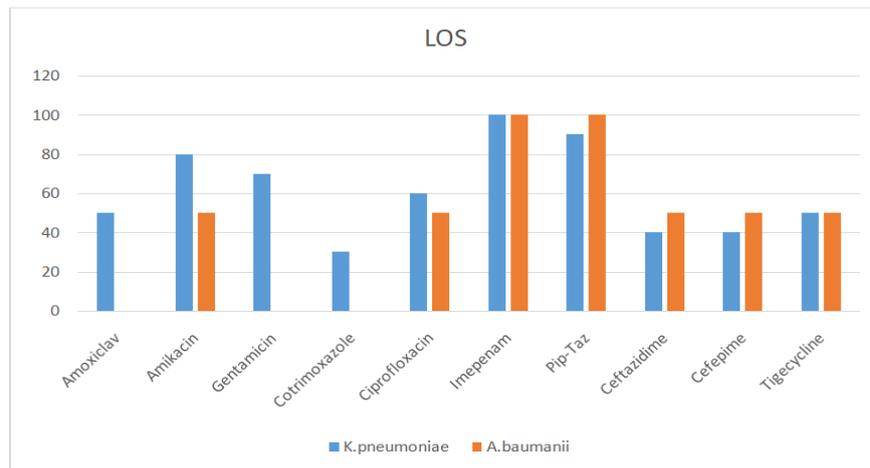




**AST of organisms causing EOS**



**AST of organisms causing LOS**



**Risk Factors Contributing To Early and Late Onset Neonatal Septicemia**

| RISK FACTORS                    | EOS    | LOS   |
|---------------------------------|--------|-------|
| Preterm                         | 68%    | 74.3% |
| Low Birth Weight                | 62.04% | 68%   |
| Respiratory Distress            | 65%    | 54.2% |
| Meconium Stained Amniotic Fluid | 34.2%  | 10%   |
| Premature Rupture Of Membranes  | 6.7%   | -     |
| Birth Asphyxia                  | 23.2%  | 12.8% |
| Teenage Pregnancy               | 2%     | -     |
| Pre- Eclampsia                  | 1.2%   | -     |
| Multiple Gestation              | 2.6%   | 1.2%  |
| Congenital Anomaly              | -      | 0.64% |

The study by Viswanathan *et al.*, (2008) also showed 6.6% of emerging carbapenem resistance among gram negative isolates. In the same study, Gram-negative organisms showed 100% resistance to ampicillin, cefotaxime, and gentamicin. According to Vinodkumar *et al.*, (2008), majority of gram negative organisms showed resistance to one or more antibiotics. In our study majority of the isolates were sensitive to imepenam and Piperacillin-tazobactam.

### Conclusion

Appropriate identification of sepsis source, prompt antibiotic prescription and aggressive management can effectively prevent adverse events following neonatal sepsis. There is requirement of simple and sustainable intervention to reduce the burden of multidrug resistant pathogens causing sepsis. The implementation of hand washing, barrier nursing and antibiotic stewardship policies is need of time.

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