



**ORIGINAL RESEARCH PAPER**

**Community Medicine**

**A STUDY OF INCIDENCE, DEMOGRAPHIC DISTRIBUTION AND THE COMMON PATHOGENS OF NEONATAL SEPSIS IN NICU:**

**KEY WORDS:** Incidence, Demography, Neonatal Sepsis, NICU.

**Dr Ravindra Y M**

Assistant Professor, Department of Community Medicine, Shridevi Institute of Medical Sciences and Research Hospital.

**Dr Raju Hanumant Patil\***

Assistant Professor, Prakash Institute of Medical Sciences and Research Centre, Urun-Islampur, Maharashtra. \*Corresponding Author

**ABSTRACT**

Detailed studies on the clinical manifestations and laboratory profile of neonatal septicaemia in rural India are uncommon. Good laboratory facilities, especially blood culture, are frequently unavailable in the rural healthcare setting, resulting in the non-availability of relevant data on culture-proven neonatal sepsis. Neonatal sepsis is a significant cause of morbidity and mortality. Appropriate clinical diagnosis and empirical antibacterial treatment in a given setting is crucial as pathogens of bacterial sepsis and antibiotic sensitivity pattern can considerably vary in different settings. As neonatal septicemia is life threatening emergency and delays in diagnosis and treatment may have immediate and longterm adverse consequences, antibiotic surveillance is needed. A sincere attempt is made to understand the Incidence and Demographic distribution of Neonatal Sepsis in NICU.

**Introduction:**

Neonatal Sepsis is the most important cause of morbidity and Mortality in developing countries. Neonatal sepsis is diagnosed when Generalized systemic features are associated with pure growth of bacteria from one or more sites.<sup>1</sup>

In developing countries, neonatal mortality (death in the first 28 days of life per 1000 live births) due to all causes is about 34 per 1000 live births, most of these deaths occur in the first week of life<sup>2,3</sup>.

In developing countries sepsis is the commonest cause of mortality responsible for 30% to 50% of 5 million neonatal deaths every year<sup>2</sup>.

It is important to remember that bacterial flora is dynamic, different from one place as compared to the other and it changes in the same place over a period of time. It is essential to closely monitor the bacterial flora of the NICU and the antibiotic sensitivity pattern of pathogens to evolve rational antibiotic policy, which is most suitable and specific for a particular NICU.<sup>1</sup>

Detailed studies on the clinical manifestations and laboratory profile of neonatal septicaemia in rural India are uncommon. Good laboratory facilities, especially blood culture, are frequently unavailable in the rural healthcare setting, resulting in the non-availability of relevant data on culture-proven neonatal sepsis<sup>4</sup>. Neonatal sepsis is a significant cause of morbidity and mortality. Appropriate clinical diagnosis and empirical antibacterial treatment in a given setting is crucial as pathogens of bacterial sepsis and antibiotic sensitivity pattern can considerably vary in different settings. As neonatal septicemia is life threatening emergency and delays in diagnosis and treatment may have immediate and longterm adverse consequences, antibiotic surveillance is needed<sup>8-13</sup>. A sincere attempt is made to understand the Incidence and Demographic distribution of Neonatal Sepsis in NICU.

**Aims and Objectives:**

To understand the Incidence and Demographic distribution of Neonatal Sepsis in NICU.

**Materials and Methods:**

**Design:** It is an observational cross sectional study.

**1. Source:** Shridevi Institute of Medical Sciences and Research Hospital.

**Period of Study:** April 2016 to March 2017.

**INCLUSION CRITERIA:**

Neonates were included when at least three of the following risk factors were present<sup>1</sup>:

1. Febrile illness in the mother during or within two weeks of delivery (more than 38° C, oral temperature).
2. More than 3 vaginal examinations during labor.

**EXCLUSION CRITERIA:**

1. Neonates with lethal congenital anomalies

**Results:**

Out of 2992 NICU admissions in the study period from April 2012 to March 2013, 419(14%) cases were taken up for the study considering inclusion and exclusion criteria. Out of 419 cases, blood culture was positive in 197 (47.016%) cases.

**Table no.1: SEX DISTRIBUTION AMONG CLINICAL AND BLOOD CULTURE POSITIVE SEPSIS**

Sex	CLINICAL SEPSIS		BLOOD CULTURE POSITIVE SEPSIS	
Male	262	p-value	94	p-value
Female	157		103	
Total	419		197	

Out of 419 cases of clinical sepsis 242(57.75%) were male neonates, 177(42.24%) were female neonates. Male neonates with clinical sepsis were admitted more frequently than female neonates which is statistically significant. (p-value 0.01)

Among 197 cases of proven sepsis 104(52.79%) were male neonates and 93(47.2%) were females neonates. There was no sex difference in blood culture positive sepsis (p-value 0.42)

**Table no. 2. BACTERIAL PROFILE IN PROVEN SEPSIS**

ORGANISM	NO OF PATIENTS
Gram negative	86(43.65%)
Klebsiella	64(32.48%)
E.coli	11(5.58%)
Pseudomonas	6(3.04%)
Proteus	4(2.03%)
Serratia	1(0.5%)
Gram positive	99(50.25%)
Coagulase positive Staphylococcus	50(25.38%)
CONS	41(20.81%)
Streptococcus pneumonia	8(4.06%)
Candida	12(6.09%)
Total	197

**Discussion:**

Among these 419 cases were suspected of septicemia and 197 cases were of proven septicemia. So, the incidence of clinical septicemia among the cases admitted in NICU, was 14% and the

incidence of proven septicemia was 6.58%.

According to NNPD (2002 - 2003) reports, the incidence varying from 0.1% to 4.5% from 18 hospitals across India<sup>4</sup>. The reported incidence of neonatal sepsis varies from 7.1 to 38 per 1000 live births<sup>5</sup>.

#### SEX:

Among 419 cases of clinical sepsis 242(57.75%) were male neonates, 177(42.24%) were female neonates, ratio being 1.3:1. Male neonates were admitted with clinical sepsis more frequently than female neonates. The ratio was similar in study done by **Muhammad Z et al (2010)**<sup>6</sup>. In the study done by **Waheed M et al(2003)** male to female ratio was 2.1:1.<sup>7</sup>

In developing countries, neonatal mortality (death in the first 28 days of life per 1000 live births) due to all causes is about 34 per 1000 live births, most of these deaths occur in the first week of life<sup>2,3</sup>.

In developing countries sepsis is the commonest cause of mortality responsible for 30% to 50% of 5 million neonatal deaths every year<sup>2</sup>.

It is important to remember that bacterial flora is dynamic, different from one place as compared to the other and it changes in the same place over a period of time. It is essential to closely monitor the bacterial flora of the NICU and the antibiotic sensitivity pattern of pathogens to evolve rational antibiotic policy, which is most suitable and specific for a particular NICU.<sup>1</sup>

Out of 419 clinical sepsis 197 (47.01%) were blood culture positive. Out of 197 organisms isolated in blood culture 99(50.25%) were gram positive, 86(43.65%) were gram negative and 12(6.09%) were fungal sepsis (candida). Gram positive and gram negative sepsis occurred in equal proportions in the present study.

Klebsiella (32.48%) was the most common organism in our study followed by coagulase positive Staphylococcus (25.38%) and CONS (20.81%). Among gram positive organisms most common were Coagulase positive Staphylococcus (25.38%), CONS (20.81%) followed by Streptococcus pneumoniae (4.06%). Among Gram negative neonatal sepsis most common are Klebsiella (32.48%), followed by E.coli (5.58%), Pseudomonas (3.04%), Proteus (2.03%) and Serratia (0.5%).

In the study done by **Viswanathan R et al(2012)**<sup>14</sup> among 216 cases of clinical sepsis, 100(46.3%) cases had blood culture positive, which was similar to our study. In their study gram negative infection were predominant (58/100 cases). Most common organism was Klebsiella followed by E.coli, Enterobacter sp.

In the study done by **Shrestha S et al(2013)** blood culture yield by conventional method was 44.13%<sup>7</sup>, which is similar to our study. Gram positive organisms were 39.36% in which Staphylococcus aureus most common followed by CONS. Gram negative organisms were 60.64%, amongst Klebsiella most common followed by Pseudomonas.

In the present study 78.6% cases were EOS and 21.3% were LOS. EOS was common presentation of proven sepsis, Klebsiella (37.4%) was common organism causing EOS. Whereas CONS (35.7%) was commonest in LOS. Similar observations were made by **Shrestha S et al(2013)**<sup>15</sup> in which 84.08% were culture proven early onset sepsis and 15.95% were late onset sepsis<sup>7</sup>.

#### Conclusion:

Klebsiella (32.48%) was the most common organism in our study followed by Coagulase positive Staphylococcus (25.38%) and CONS (20.81%)

#### Conclusion:

So, the incidence of clinical septicemia among the cases admitted

in NICU, was 14% and the incidence of proven septicemia was 6.58%.

#### References:

1. Meharban Singh; Care of the newborn, 7th Edition; 223–230
2. Ashok K. Deorari; Neonatal sepsis: Manageable daunting issue for India. Journal of Neonatology 2009, Vol.23, 1, 2009: 7–11.
3. Bryer J, Boschi - Pinto C, Shibuya K, Black RE; WHO Child Health Epidemiology Reference Group. WHO Estimate of the causes of death in children. Lancet 2005; 365: 1147–1152.
4. Viswanathan R, Singh AK, Ghosh C, Dasgupta S, Mukherjee S, Basu S. Profile of neonatal septicemia at a district-level sick newborn care unit. J Health Popul Nutr. 2012 Mar;30(1):41-8.
5. Al-Shamahy HA, Sabrah AA, Al-Robasi AB, Naser SM et al Types of Bacteria associated with Neonatal Sepsis in Al-Thawra University Hospital, Sana'a, Yemen, and their Antimicrobial Profile. Sultan Qaboos Univ Med J. 2012 Feb;12(1):48-54.
6. Muhammad Z, Ahmed A, Hayat U, Wazir MS, Rafiyatullah, Waqas H et al Neonatal sepsis: causative bacteria and their resistance to antibiotics. J Ayub Med Coll Abbottabad. 2010 Oct-Dec;22(4):33-6.
7. Waheed M, Laeeq A, Maqbool S. et al The etiology of neonatal sepsis and patterns of antibiotic resistance. J Coll Physicians Surg Pak. 2003 Aug;13(8):449-52.
8. Report 2002-2003. National Neonatal and Perinatal Database Network. New Delhi. National Neonatology Forum of India 2004
9. Explore simplified antimicrobial regimens for the treatment of neonatal sepsis. WHO/FCH/CAH/04/1/2002.
10. Deorari AK. For the investigators of National Neonatal perinatal Database: Changing pattern of bacteriologic profile in neonatal sepsis among intramural babies. J Neonatol 2006; 20: 8-15.
11. M Jeeva Sankar, Jhuma Sankar, Deepak Chawla, Sushma Nangia. Antibiotic usage in neonates – guidelines and current practices Journal Neonatology 2009; Vol-23 No.1: 68-77
12. Adams-chapman. Stoll BJ. Prevention of nosocomial infection in the neonatal intensive care unit. Curr Opin Pediatr. 2002; 14:157-164.
13. J.E. Colle, R.S. Miles, B.Watt. tests for the identification of bacteria Practical medical microbiology edited by Mackie and Mc. Cartney 14th Edition.
14. Viswanathan R, Singh AK, Ghosh C, Dasgupta S, Mukherjee S, Basu S. Profile of neonatal septicemia at a district-level sick newborn care unit. J Health Popul Nutr. 2012 Mar;30(1):41-8.
15. Shrestha S, Shrestha NC, Dongol Singh S, Shrestha RB, Kayestha S, Shrestha M, Thakur NK et al Bacterial Isolates and its Antibiotic Susceptibility Pattern in NICU. Kathmandu Univ Med J (KUMJ). 2013 Jan-Mar;11(41):66-70.