http://jmscr.igmpublication.org/home/ ISSN (e)-2347-176x ISSN (p) 2455-0450 crossref DOI: https://dx.doi.org/10.18535/jmscr/v10i5.09



Journal Of Medical Science And Clinical Research

### Study of Bacteriological Profile and Antimicrobial Susceptibility Pattern of Nosocomial Infection in Intensive Care Unit-In a Tertiary Care Hospital

Authors

Dr Bijoyeta Das<sup>1</sup>, Dr Subhrendu Sekhar Sen<sup>2</sup>

<sup>1</sup>Post Graduate Trainee, <sup>2</sup>Associate Professor Department of Microbiology, Silchar Medical College & Hospital Corresponding Author **Dr Subhrendu Sekhar Sen** 

#### Introduction

Compared with an average patient, an ICU patient has five to seven folds higher risk of nosocomial infection and ICU infections contributes to 20% to 25% of all nosocomial infections in a hospital.

The patterns of microorganisms causing various infections and their antibiotic resistance pattern vary widely from one country to another; as well as from one hospital to another and even among ICUs within one hospital.

Having proper knowledge regarding the antibiotic resistance pattern in hospital settings will help in implementing proper antibiotic policy for the hospital and hence benefit the community in this region.

#### Aims & Objectives

The objectives of the present study were the following

- 1) To determine the prevalence of nosocomial infections in patients admitted in different Intensive Care Units of Silchar Medical College.
- 2) To study the bacteriological profile, their drug sensitivity and resistance

pattern from the isolates of the patients.

#### Materials & Methods

**Study Design:** Observational study (cross sectional study)

#### Place of study:

- 1) Department of Microbiology, Silchar Medical College & Hospital, Silchar
- Different Intensive Care Units (ICUs) of Silchar Medical College & hospital, Silchar

**Study Period:** One year from June 2018 to May 2019.

#### **Clinical Specimen**

- The different types of the samples were-Blood, Urine, Catheter tip, Endo-tracheal tube, Endo-tracheal aspirate, Pus from surgical site.
- Different departments from where samples were collected are Neonatal Intensive Care Unit (NICU), Pediatric Intensive Care Unit (PICU), Medicine ICU, Obstetrics &

Gynecology ICU, etc. and samples from patients of any age and both the sexes were included.

#### Collection

Under strict aseptic condition samples were immediatelv transferred collected and to Bacteriology section of Department of Microbiology, Silchar Medical College & Hospital for processing.

#### Processing of the sample

First samples were inoculated in culture media for primary isolation of bacteria. The media used are as follows:

- a) 5% sheep blood agar medium and MacConkey's agar for all specimens.
- b) Cysteine Lactose Electrolyte Deficient Media which was used for urine

#### Isolation and identification of isolates

The isolates were identified by Colony morphology and cultural characteristics, Gram stain, Motility, Biochemical tests, Sugar fermentation test.

# Antibiotic susceptibility testing by disc diffusion method (CLSI 2018):

- Mueller Hinton agar plates were made till depth of 4mm.
- Pure culture was selected for preparation of inoculums.3-4 similar colonies were selected and transferred into peptone water. Incubated at 35°C for 2-8 hours till light moderate turbidity was achieved. The turbidity was adjusted to Mac Farlands standard  $0.5(1.5 \times 10^8$  CFU/ml).
- A sterile cotton swab was dipped in the

suspension

- The soaked swab was rotated firmly against the upper inside wall of thetube to remove excess fluid.
- It was streaked evenly into entire agar surface of the plate three times, turning the plate 60° between each streaking.
- The inoculums is dried 5-15 minutes with lid in place.
- Commercially prepared antibiotic discs were applied using aseptic technique.
- The discs were placed with centres at least 24 mm apart.
- Incubated immediately and examined after 14-16 hours.
- The zone of inhibition was measured in millimetres with standard chart provided through CLSI Guidelines 2018.
- Quality control of Antimicrobial susceptibility testing is done by using reference strain i.e. *Staphylococcus aureus* ATCC- 25923, *Escherichia coli* ATCC- 25922, *Klebsiella pneumoniae* ATCC- 700603 as per standard CLSI guideline.

#### **Results & Observations**

The following observations were made in the study:

#### 1. Prevalence of Nosocomial infection

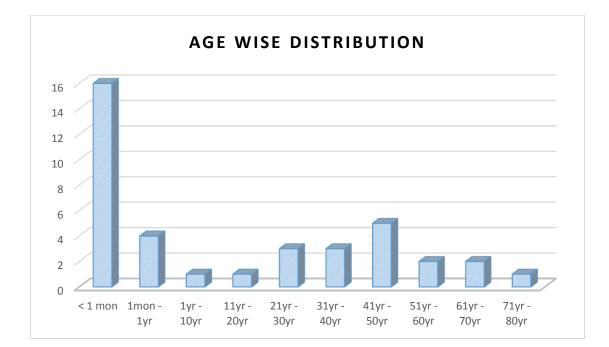
During the period of the study, a total of 160 samples were analyzed. Of these, 38 patients (23.75%) were found to develop Nosocomial infection. Type of samples & their positivity pattern on culture are shown in **table** 

Type of Sample	Total number	Positivity (%)
Urine, Catheter tip	52	21 (40.38%)
Blood, Peripheral venous line etc.	68	08 (11.76%)
Tracheal aspirate, Endotracheal tube	30	08 (26.66%)
Pus from surgical site, Drain tip	10	01 (10%)

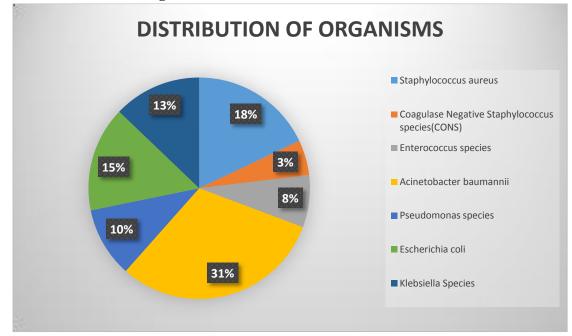
- 2. Demographic characteristics of Nosocomial infection:
  - A) Isolation of the samples were from various ICUs, & showed that more number of specimens were isolated from NICU 29.31% (17 out of 58) followed by SNCU 26.66% (4 out of 15), ICU 26.08% (6 out of 23), PICU 19.04% (4 out of 21), SICU 20% (1 out of 5), MICU 18.51% (5 out of

27), HDU 9% (1 out of 11) as follows in Table / chart 4

- B) Patients who developed nosocomial infection, among them 59 (37%) were male and 101 (63%) were female (P value 0.039)
- C) The isolates belonged to different age groups. Prevalence of nosocomial infectionamong different age group is depicted in table.



3. Distribution of isolated organisms in nosocomial infection



# **4.** Antibiotic Susceptibility pattern: Antibiotic susceptibility pattern of the isolates of our as follows—

- In our study we noted that non-fermenting Acinetobacter species ranks top in the list clinical isolates from among ICU. Antimicrobial susceptibility pattern was analyzed among these isolates and it was observed that most of NF-GNB were multi drug resistant organisms (MDRO) being resistant to three or more class of antibiotics. High rates of resistance was noted to even carbapenems and aminoglycosides.
- Second most common organism in our study was *Staphylococcus aureus*. 42.8% of *Staphylococcus aureus* were found to be MRSA and the proportion of VRE was 0%.

2022

• Members of the Enterobacteriacae family are the third in the list of most common clinical isolates. *Escherichia coli* and *Klebsiella* spp were most common clinical isolates in this family. In both of these GNB, high rates of non susceptibility was noted against quinolones, cephalosporins and beta lactam inhibitor group of drugs. Resistance to carbapenems was also significantly higher.

Antimicrobial Agent	Acinetobacter species (12)	Pseudomonas species (4)
Ceftazidime	11 (91.6%)	2 (50%)
Cefoperozone+ Salbactam	3 (25%)	1 (25%)
Meropenem	10 (83.3%)	2 (50%)
Gentamicin	10 (83.3%)	3 (75%)
Amikacin	10 (83.3%)	2 (50%)
Ciprofloxacin	10 (83.3%)	3 (75%)
Levofloxacin	9 (75%)	1 (75%)
Tetracycline	5 (41.6%)	
Piperacillin+Tazobactam		1(25%)

#### [Table/Fig-7]: Antimicrobial resistance pattern of non-fermenting GNB.

\*% indicates resistance to corresponding antimicrobial agent.

[Table/Fig-8]: Antimicrobial resistance pattern of gram positive cocci.

Antimicrobial agent	Staphylococcus. aureus (7)	Enterococcus species(3)	CoNS (1)
Penicillin	6 (85.7%)		0 (0.00%)
Ampicillin		2 (66.6%)	
Amoxycillin+	6 (85.7%)		
Clavulanic acid			
Gentamicin	2 (28.5%)		
Ciprofloxacin	4 (57.1%)		0 (0.00%)
Erythromycin	3 (42.8%)		
Clindamycin	1 (14.2%)		1 (100%)
High level Gentamicin		1 (33.3%)	
Vancomycin	0 (0.00%)	1 (33.3%)	0 (0.00%)
Cefoxitin	3 (42.8%)		1 (100%)
Linezolid	0 (0.00%)		

\*% indicates resistance to corresponding antimicrobial agent.

Antimicrobial agent	Escherichia coli (6)	Klebsiella species (5)
Ceftriaxone	5 (83.3%)	4 (80%)
Ceftazidime	5 (83.3%)	4 (80%)
Cefoperazone+ Sulbactam	1 (16.6%)	1 (20%)
Meropenem	2 (33.3%)	2 (40%)
Gentamicin	4 (66.6%)	3 (60%)
Amikacin	2 (33.3%)	3 (60%)
Ciprofloxacin	5 (83.3%)	4 (80%)

[Table/Fig-9]: Antimicrobial resistance pattern of Enterobacteriacae

\*% indicates resistance to corresponding antimicrobial agent

So, it can be concluded from our study that overall MDRO (resistant to 3 or more class of antimicrobial agents) isolates were 59.2%, MRSA isolates were 42.8%, MR CoNS isolates 100%, VRE 0.00%

#### Conclusion

From this prospective cross-sectional study, it can be concluded that Nosocomial infection is a significant problem in this region.

The risk of the development of nosocomial infections is directly related to the duration of ICU stay and the duration of the use of the use of the use of the indwelling catheters/ tubes. The prolonged use of indwelling devices needs careful prophylactic standards of microbiological monitoring.

The empirical and the indiscriminate usage of antibiotic should be avoided in order to curtail the emergence and spread of drug resistance among nosocomial pathogens.

A sustained co-ordination between the clinician and the clinical microbiologist is essential not only for improving clinical outcome but also for optimizing resource utilization. Use of higher antibiotics like carbapenems results in increased healthcare associated cost and burden, also contributing to spread of drug resistance among nosocomial pathogens

Improper detection and reporting of Nosocomial infection may lead to major clinical and epidemiological consequences. Antibiotic policies, effective surveillance and scrutiny of epidemiological trends of the infections are need of the hour for better management of ICU infections with resistant organisms. As the limitation of our study is small sample size, further studies on resistance pattern with larger sample size, and also samples from the health care providers would provide a clearer picture of the prevalence in this hospital.

#### Bibliography

- Dugad AG et al. Prevalence of nosocomial infection; Int J Res Med Sci.2015 Dec;3(12): 3514-3518.
- 2. Apurva Sastry; Essentials of Hospital infection control; Introduction to Hospital acquired infection:p1
- Shehabi, A.A., Baadran, I. 1996. Microbial infection and antibiotic resistance patterns among Jordanian intensive care patients. Eastern mediterranean Health J.,2(3): 515-520
- Ducel, G., Fabry, J., Nicolle, L.2002. editors. Prevention of Hospital acquired infections: A Practical guide.2<sup>nd</sup> ed. Geneva, World Health Organization
- Krishna, Prakash, S. Nosocomial infection: An overview New Delhi: Maulana Azad Medical College.13P
- Harrison's Principle of internal Medicine, Hospital acquired infections, 16<sup>th</sup> edition Vol 1 Chapter 92, page 442
- Günserena, F., Mamıkogʻlua, L., Ztürkb, S., Yücesoyc, M., Biberogʻluc, K., Yulugʻc, N., *et al.* 1999. A surveillance study of antimicrobial resistance of Gramnegative bacteria isolated from intensive care units in eight hospitals in Turkey. *J. Antimicrob. Chemother.*, 43: 373–378.

- Barai, L., Fatema, K., AshrafulHaq, J., Omar Faruq, M., Areef Ahsan, A.S.M., Golam Morshed, M.A.H., *et al.* 2010. Bacterial profile and their antimicrobial resistance pattern in an intensive care unit of a tertiary care hospital in Dhaka. *Ibrahim Med. Coll. J.*, 4(2): 66-69.
- 9. Topley & Wilson
- HAI: Guidelines to laboratory methods, edited by M.T. Parker WHO regional Publications, European series no.4,1978
- Garner JS et al. CDC definitions for nosocomial infections, 1988. Am J infect control, 1988, 16: 128-140
- 12. Centres for disease control & prevention. " National Healthcare Safety Network (NHSN) Patient Safety component Manual. CDC, Atlanta" (2019)
- Benett JE, Dolin R, Blaser MJ. Mandell Douglas and Benett's Principle & Practice of infectious diseases, 8<sup>th</sup> edition. Philadelphia: Elsevier;2014
- 14. Damani N. Manual of infection prevention & control. Oxford: OUP Oxford; 2011 Dec 22
- 15. Dudeck MA, Edwards JR, Allen-BridsonK, et al. NHSN 2013 HAI Data: National Healthcare Safety Network report, data summery for 2013, deviceassociated module. Am j infect control. 2015; 43(3): 206-21
- 16. Singh S, Chakravarthy M, Chaya V, et al. Analysis of a multicentric pooled healthcare associated infection data from India: New insights. J Natl Accredit Board Hosp HealthcProvid. 2014; 1:39
- 17. Mehta Y, Jaggi N, Rosenthal VD, et al. Device-Associated Infection Rates in 20 cities of India, Data Summery for 2004-2013: Findings of the International Nosocomial infection Control Consortium. Infect Control Hosp Epidemiol. 2016; 37(2): 172-81
- 18. Kalanuria AA, Zai W, Mirski M. VAP pathogenesis: Ventilator-associated

pneumonia in the ICU. Crit care.2014;18(2):208.

- Weinstein RA, Bonten MJ, Kollef MH, et al. VAP risk factors: Risk factors for ventilator-associated pneumonia: from epidemiology to patient management. Clin Infect Dis.2004;38(8):1141-9
- 20. Singh S, Chakravarthy M, Rosenthal VD, et al. Surgical site infection rates in six cities of India: findings of the International Nosocomial Infection Control Consortium (INICC). Intern health. 2014;7(5):354-9
- 21. Singh R, Singla P, Chaudhary U. Pathogenesis: Surgical site infections: classification, risk factors, pathogenesis and preventive management. Int J Pharm Res Health Sci.2014;2:203-14
- Cui P, Fang X. Pathogenesis of infection in surgical patients. Curropin crit care. 2015;21(4):343
- 23. Vincent JL, Bihari DJ, Suter PM, Bruining HA, White J, Nicolas-Chanoin MH, Wolff M, Spencer RC, Hemmer M: The prevalence of nosocomial infection in intensive care units in Europe. Results of theEuropean Prevalence of infection in Intensive Care (EPIC) Study. EPICInternational Advisory Committee. JAMA 1995;274:639-644.
- 24. Vincent JL, Rello J, Marshall J, Silva E, Anzueto A, Martin CD, Moreno R, Lipman J, Gomersall C, Sakr Y, Reinhart K. The Extended Prevalenceof Infection in the ICU Study: EPIC II. JAMA 2009;302:2323-2329.
- 25. Esen S, Leblebicioglu H. Prevalence of nosocomial infections at Intensive care units in Turkey: a multicentre 1-day point prevalencestudy. *Scan J Infect Dis* 2004;36:144-8.
- 26. Legras A, Malvy D, Quinioux AI, Villers D, Bouachour G, Robert R, Thomas R. Nosocomial infections: prospective survey of incidencein five French intensive care units. *Intensive Care Med* 1998;24:1040-6.

2022

- Malacarne P, Boccalatte D, Acquarolo A, Agostini F, Anghileri A, Giardino M, Giudici D, Langer M, Livigni S, Nascimben E, Rossi C,Bertolini G. Epidemiology of nosocomial infection in 125 Italianintensive care units. *Minerva Anestesiol*2010;76:13-23.
- 28. Kallel H, Dammak H, Bahloul M, Ksibi H, Chelly H, Ben Hamida C, Rekik N, Bouaziz M. Risk factors and outcomes of intensive care unitacquiredinfections in a Tunisian ICU. *Med Sci Monit*2010;16:PH 69-75.
- 29. Ding JG, Sun QF, Li KC, Zheng MH, Miao XH, Ni W, Hong L, Yang JX, Ruan ZW, Zhou RW, Zhou HJ, He WF. Retrospective analysis ofnosocomial infections in the intensive care unit of a tertiary hospitalin China during 2003 and 2007. *BMC Infect Dis* 2009;9:115.
- 30. Richards MJ, Edwards JR, Culver DH, Gaynes RP. Nosocomial infections in medical intensive care units in the United States.National Nosocomial Infections Surveillance System. *Crit Care Med*1999;27:887-92.
- 31. Vincent JL, Sakr Y, Sprung CL, Ranieri VM, Reinhart K, Gerlach H, Moreno R, Carlet J, Le Gall JR, Payen D: Sepsis in European intensivecare units: Results of the SOAP study. *Crit Care Med* 2006;34:344-53.
- 32. Siegel T, Mikaszewska-Sokolewicz M, Mayzner-Zawadzka E. Epidemiology of infections at the intensive care unit. *Pol MerkurLekarski*2006;20:309-14.
- 33. 34) de Oliveira AC, Kovner CT, da Silva RS. Nosocomial infection in an intensive care unit in a Brazilian university hospital. *Rev Lat AmEnfermagem*2010;18:233-9.
- 34. Agarwal R, Gupta D, Ray P, Aggarwal AN, Jindal SK. Epidemiology, risk factors and outcome of nosocomial infections in a Respiratoryintensive Care Unit in North India. *J Infect* 2006;53:98-105.

- 35. Pampita Chakraborty, Sukumar Mukherjee. A study on the Prevalence and Microbiological Profile of Nosocomial Infections in the ICU of a Tertiary Care Hospital in Eastern India. Int. J. Curr.Microbiol.App.Sci (2016) 5(5): 920-925
- 36. Ambanna Gowda Durgad, Sudarshan Varadarajan. Prevalence of nosocomial infections in the intensive care unit. Int J Res Med Sci. 2015 Dec; 3(12): 3514-3518
- 37. Zaveri Jitendra R, Patel Shirishkumar M, Nayak Sunil M, Desai Kanan, Patel Parul. A study on bacteriological profile and drug sensitivity & resistance pattern of isolates of the patients admitted in intensive care units of a tertiary care hospital in Ahmedabad. Natl J Med Res 2012 2(3): 330-334
- 38. Shalini S, Kranthi K, Gopalkrishn Bhat K. The Microbiological Profile of Nosocomial infections in the Intensive care Unit. J Clin Diag Res 2010 Oct [ cited: 2010 Oct 31]; 4: 3109-3112
- 39. Akash Deep, R. Ghildiyal, S. Kandian and N. Shinkre. Clinical and microbiological profile of nosocomial infections in the pediatric intensive care unit (PICU). Indian Pediatr. 2004 Dec;41(12): 1238-46
- 40. Dileep Kumar Sharma, Yogendra Kumar Tiwari, Nitya Vyas, Rakesh Kumar Maheshwari. An investigation of the incidence of Nosocomial infection among patients admitted in the intensive care unit of a tertiary care hospital in Rajasthan, India.Int, J.Curr.Microbiol. App.Sci (2013) 2(10):428-435
- 41. Sugata Dasgupta, Soumi Das, Neeraj S. Chowan, AvijitHazra. Nosocomial infections in the Intensive care unit: Incidence, risk factors, outcome and associated pathogens in a public tertiary teaching hospital of Eastern India.

Dr Bijoyeta Das et al JMSCR Volume 10 Issue 05 May 2022

2022