Evaluating the Trends of Bloodstream Infections by Nonfermenting Gram Negative Bacilli among the Patients in a Tertiary Care Hospital of Western Part of India and its Antibiogram

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ABSTRACT

Non-fermenting gram-negative bacilli (NFGNB) are an emerging problem in Blood stream infections. A major concern is multi-drug resistance which severely limits treatment options. Earlier it was believed to be non pathogenic but recently they are more frequently isolated as primary pathogen. Usually they cause hospital acquired infection (HAI). A prospective study was conducted to isolate the NFGNB from blood samples, to identify the risk factors leading to blood stream infections and to determine the antibiotic susceptibility pattern of them. The study was conducted in a tertiary care hospital, over a period of 2 years. Identification of NFGNB was done by biochemical tests and by VITEK 2. Antibiotic susceptibility was determined by disc diffusion method. Extended-spectrum β-lactamases (ESBLs) and metallo-β-lactamases (MBLs) production were detected by the combined disc diffusion test. Out of 2021 blood samples, blood culture positive was in 32.7% of patients of whom the cause was NFGNB. Acinetobacter baumannii was the most common organism, 27.69% followed by Stenotrophomonas maltophilia, next to it was Pseudomonas aeruginosa Acinetobacter lwoffiiëte. The most common risk factors for colonization BSIs with NFGNB was comorbid conditions, such as diabetes mellitus, cardiovascular diseases, hypertension, tuberculosis and chronic renal disease patients on haemodialysis. In general, the isolates of NFGNB revealed pretty much good sensitivity to carbapenem (imipenem, ertepenam), colistin and aminoglycosides (amikacin, gentamicin), where as cephalosporin group revealed a low susceptibility rate. ESBL and MBL producer NFGNB were identified and the isolation rate is very alarming. The trend of increasing numbers of cases of NFGNB in Blood stream infections compounded by MDR is of great concern. It is necessary to administer antibiotics judiciously, strengthen surveillance and laboratory services in intensive care units, and re-evaluate treatment guidelines for management of infection by these organisms.

Keywords
Gram-Negative Non-Fermenting Bacilli (NFGNB), Blood Stream Infections (BSIs), Multi-drug resistance

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Introduction

The non-fermenting organisms are comprised of gram negative rod shaped bacilli.\(^1\) The non-fermenting gram negative bacilli (NFGNB) are taxonomically group of aerobic non spore forming bacilli that either do not utilize carbohydrates as the source of energy or degrade them through metabolic pathways other than fermentation.\(^2\) They are widely distributed in nature as saprophyes, found in soil, water, sewage or as commensals on human skin or in the human gut and some of them found in hospital environment.\(^1, 3, 4\) These nonfermenters are unfortunately the by-product of medical and surgical advances in health care system of serious ill patients.\(^5\) Recently, these NFGNB are emerging problem in sepsis, which is associated with significant mortality and morbidity. A major concern is multi-drug resistance which severely limits treatment options.

The predominant species of concern among NFGNB are *Pseudomonas aeruginosa*, *Acinetobacter baumannii*, *Stenotrophomonas maltophilia* and, less so, members of the *Burkholderia cepacia* group.\(^3\) Except *P. aeruginosa* the NFGNB are most often cause nosocomial infections in immune-compromised patients like urinary tract infections (UTI), Bloodstream infections (BSIs), ventilator associated pneumonia (VAP) and surgical site infections (SSI).\(^1\)

Bloodstream infections (BSIs) are the significant causes of morbidity and mortality for many patients.\(^6\) BSIs are defined as the presence of viable infectious microorganism in the bloodstream causing clinical illness.\(^7\) The term bloodstream infection and bacteremia are synonymously used, which generally refer to the significant growth of a microorganism in a blood culture obtained from the patient with clinical signs of infection.\(^8\) Bacteremia may range from self-limiting infections to septicaemia which is life threatening and needs rational antimicrobial treatment.\(^9\) In the developing countries, like India lack of standard antimicrobial guidelines, emergence of antimicrobial resistance, paucity of good diagnostic facilities and poor hospital environment, poor quality of hand hygiene are major denominators for surge in BSI associated morbidity and mortality.\(^10\)

Materials and Methods

This was a prospective study. The study was conducted in the Microbiology Department of Dr. D.Y. Patil Medical College, Hospital and Research Centre, over a period of 2 years (i.e. July 2012 to September 2014). A total 2021 blood samples from the suspected patients of sepsis were collected in the adult and paediatric patients. Bloods were collected aseptically in brain heart infusion broth (BHI) or in BACT/ALERT 3D system. In case of neonates 2 ml blood, children 3-5 ml blood and for the adults 10 ml blood were taken. The samples were taken from the suspected patients, admitted to different wards and various intensive care units (ICU) of this hospital. The study was approved by the Ethical Committee of our institute.

Blood samples were processed for culture by standard conventional methods. Identification of Nonfermenters were carried out by Gram staining (gram negative bacilli/ gram negative coccobacilli), cell and colony morphology, pigment production, catalase test, p citrate test, triple sugar iron (alkaline slant/ no change butt), oxidase test and by motility test. Further identification was done by Hugh and Leifson oxidative-fermentative test (O-F) for glucose, sucrose, lactose, mannitol; gelatin liquefaction, nitrate reduction test, Decarboxylation of arginine, lysin and ornithine and growth at 35\(^\circ\)C and at 42\(^\circ\)C for 18-24 hours on two tubes of trypticase soy
agar (TSA). The final identification and confirmation was done by the Vitek 2 system.2

Identification of pigment production by King’s A and King’s B medium11

King’s A medium11: Pyocyanin, a blue phenazine derivative characteristic of *P. aeruginosa* was diffusible and its production was enhanced by growth in “King A (Fig. 1)”11.

King’s B medium11: Fluorescent *Pseudomonas* were characterised by production of water soluble pigment, which diffused freely in the media and fluoresce brightly under U.V ray. The organisms produced this pigments were *P. aeruginosa, P. putida, P. fluorescens, P. chlororaphis* etc. and was manifested in low iron containing media.6 “King B” medium was the universally use medium for the production of fluorescent pigment.11

Antibiotic susceptibility testing was determined by Kirby - Bauer disc diffusion method2,12

Muller-Hinton agar media was used. Commercially available Himedia discs were used. The strength of the discs used and their zone size interpretation were carried out by National Committee for Clinical Laboratory Studies (NCCLS) guideline. The antibiotics, which were tested, Piperacillin (10mcg/disc), Carbenicillin (100mcg/disc), Ampicillin (10mcg/disc), Cefotaxim (30mcg/disc), Ceftriaxone (30mcg/disc), Ceftazidime (30mcg/disc), Cotrimaxazole (25 mcg/disc), Ciprofloxacin (5 mcg/disc), Norfloxacain (10 mcg/disc) Gentamicin (10mcg/disc), Amikacin (30mcg/disc), Imipenem (10mcg/disc), Chloramphenicol (30 mcg/disc) Tobramycin (10mcg/disc), Ofloxacin (5mcg/disc), Amoxicillin/Clavulanic acid (20/10mcg/disc), Piperacillin/Tazobactam (100/10mcg/disc), Tigecycline (15mcg/disc), Colistin (10mcg/disc) and Ertepenem (10mcg/disc).

Detection of extended spectrum β-lactamases production12,13

The Combine disk diffusion test (CDDT) was used to determine the prevalence of extended spectrum β-lactamases (ESBL) production. Muller-Hinton agar media was used. One Ceftazidime (CAZ) (30μg) disc was placed on a lawn culture of test isolates and at the distance of 15 mm on both side of CAZ disc, a combination disc of Ceftazidime/Tazobactam (30/10 μg) and Ceftazidime / Clavulanic acid (30/10 μg) were placed. A≥ 5 mm increased in a zone diameter for either antimicrobial agent tested in combination with Clavulanic acid or Tazobactam versus the zone diameter of the agent when tested alone = ESBL producer (Fig. 2).10,13

Detection of metallo β-lactamases production

Muller-Hinton agar media was used. One Imipenem (10μg) disc was placed on a lawn culture of isolates and at the distance of 15 mm a combination disc of 10μg of Imipenem and 100μl of EDTA disc was placed. Then it was incubated at 35°C for 18 - 24 hours. An increase in zone size ≥ 7 mm around the Imipenem -EDTA disc as compared to Imipenem disc alone was recorded as positive (Fig. 3).10,13

Results and Discussion

In this study, out of 2021blood samples, total number of culture positive isolates were 661 (32.7 %) among which 445 (67.32%) were gram positive cocci (GPC) and 216 (32.68%) were gram negative bacilli (GNB). Out of 216 GNB, 65 (30.1%) were non-fermenting gram
negative bacilli (NFGNB). Out of the total 65 isolates, highest number of isolates (23%) were obtained from male surgical ward, followed by Medicine Intensive Care Unit (MICU) (10.8%) next to it was male medicine ward (9.6%) (Fig. 4). While discussing about the gender distribution, in this study male (69.23%) outnumbered the female (30.77%) (Fig. 5). In our study the patients were divided into ten age groups. The majority of the patient belongs to 41 to 50 years, accounting for 27%, followed by the age group of 31 to 40 years comprises 18%, next to this is the age group of 11 to 20 years accounting for 9.23% (Fig. 6).

The highest number of isolates were Acinetobacter boumanna, comprises 27.69% followed by Stenotrophomonas maltophilia (previous designation: Pseudomonas maltophilia) 21.53%, next to it was Pseudomonas aeruginosa (13.84%), Acinetobacter lwoffii (6.15%), Pseudomonas fluroscence (4.61%), Acinetobacter boumanna complex (ABC) (4.61%), Burkholderia cepacia (4.61%) (previous designation: Pseudomonas cepacia), Sphingomonas paucimobilis (3.07%) (previous designation: Pseudomonas paucimobilis), Pseudomonas stutzeri (3.07%), Pseudomonas putida (3.07%) and each one isolates of Acinetobacter radioresistance, Acinetobacter calcoaceticus, Acinetobacter haemolyticus, Burkholderia multivorans and Moraxella osloensis (Table 1).

In this study we have analyzed the risk factors for colonization BSIs with NFGNB. Prolonged hospitalization, mechanical ventilation, indwelling foreign devices (especially orthopedic implants, in-situ canula), unjustified antimicrobial therapy and comorbidities, have identified as risk factors which are predisposing to acquisition BSIs by NFGNB. In this study 29.23% isolates were obtained from the patients who had comorbid conditions, such as diabetes mellitus, cardiovascular diseases, hypertension, tuberculosis and chronic renal disease patients on haemodialysis. Around 24.61% isolates were obtained from the patients, who were on indwelling intravascular catheters or orthopedics implants in situ, followed by18.46% of isolates from those patients who have admitted in this hospital for a long tenure, next to it was 15.38% isolates from those patients who were on mechanical ventilators and 12.31% isolates were yield from the patients who had prolonged history of hospitalization (Fig. 7).

The isolates of Pseudomonas aeruginosa revealed 100 % sensitivity to Colistin and also revealed good susceptibility to Ertepenam (90.8%) followed by Imipenem (86.77%), Tobramycin (66.66%) next to it, was Amikacin (64.02%) (Fig. 8). The isolates of Acinetobacters showed 60% were sensitive to Imipenem. In this study we have reported 52.2% susceptibility to chloramphenicol and 48.9% to gentamicin. Close to it, in this study amikacin and norfloxacin each comprises of 47.8%. In this study Ceftazidime shows a bit low sensitivity pattern, accounting for 37.8 % (Fig. 9).

The isolates of Stenotrophomonas maltophilia showed 100 % sensitivity to Colistin revealed good susceptibility to Ertepenam (96.65%), Ofloxacin (94.12%), Ceftazidime (94.12%) followed by Ciprofloxacin (88.23%) (Fig. 10). Among the total 65 isolates of NFGNB, 20 isolates (30.77%) were multidrug resistance (MDR). However, amidst these 20 isolates 11 (55%) were ESBL- producers and rest (45%) were MBL- producers.

S. maltophilia showed a good sensitivity to eretpenam (96.65%), ofloxacin (94.12%), ceftazidime (94.12%) and ciprofloxacin (88.23%)
Table 1 Distribution of non-fermenting gram negative bacilli in different clinical samples (n=65)

| Name of the organism                      | Number of isolates (%) |
|-------------------------------------------|------------------------|
| Pseudomonas aeruginosa                    | 9 (13.84%)             |
| Pseudomonas fluroscence                   | 3 (4.61%)              |
| Pseudomonas putida                        | 2 (3.07%)              |
| Pseudomonas stutzeri                      | 2 (3.07%)              |
| Acinetobacter baumannii                   | 18 (27.69%)            |
| Acinetobacter baumannii complex (ABC)     | 3 (4.61%)              |
| Acinetobacter lwofii                      | 4 (6.15%)              |
| Acinetobacter radioreistance              | 1 (1.53%)              |
| Acinetobacter calcoaceticus               | 1 (1.53%)              |
| Acinetobacter haemolyticus                | 1 (1.53%)              |
| Burkholderiacepacia                       | 3 (4.61%)              |
| Burkholderiamultivorans                   | 1 (1.53%)              |
| Stenotrophomonasmaltophilia               | 14 (21.53%)            |
| Sphingomonas paucimobilis                 | 2 (3.07%)              |
| Moraxella osloensis                       | 1 (1.53%)              |
| Total                                     | 65                     |

Fig. 1 Kings B medium under U-V ray

Fig. 2 ESBL producer
Fig. 3 MBL producer

Fig. 4 Ward wise distribution of different clinical samples (n=65)

Fig. 5 Gender distribution of the patients (n=65)
Fig. 6 Age distribution of the patients (n=65)

Fig. 7 The incidence of infection due to gram negative nonfermenting organisms

Fig. 8 Antibiotic susceptibility pattern of Pseudomonas aeruginosa (n=9)
Bloodstream infections by NFGNB remained a challenge for the clinician and microbiologists due to the limited facilities in the laboratories to identify NFGNB, changing bacterial etiology and emergence of antimicrobial resistance. Early detection of NFGNB and determination of its antimicrobial susceptibility can reduce the occurrence of BSI and can also decrease the rate of emergence of MDR isolates. Our study evaluates the incidences of bloodstream infections by NFGNB, risk factors underlying and antimicrobial susceptibilities among the paediatric and adult group of patients.

The non-fermenting gram negative bacilli are found in nature as inhabitants of soil, water and also the commensals of human and animal mucous membranes. Recently these organisms are gaining importance as the frequently isolated primary pathogen in patients with prolonged hospitalization.
NFGNB have the ability to adapt well in hospital environment as they can survive on dry surfaces, in antiseptic solution and distilled water for many days. They can easily have transmitted to human body by sources like indwelling intravascular catheters, drain tubes from surgical site, surgical intervention and from other inanimate objects like bed rails, bedside tables, ventilators, air humidifiers and sinks and from these the NFGNB is transmitted to the patients.

In this study a total of 2021 blood samples were processed. In this study, overall incidence of bloodstream infection by NFGNB was based on significant bacterial growth in the blood cultures obtained from suspected patients was 732.7%. Comparatively, in 2013 a study done in Eastern India had revealed 201 non-fermenters were isolated from 1650 clinical samples, accounting for an isolation rate from blood culture is 16.41%. 14Where as another study in Gujrat by Patel et al., isolated 2397 (23.93%) NFGNB, out of total 20721 various clinical samples, accounting for isolation rate of blood culture is 6.96%.15

Infection due to NFGNB can occur at any age. Bloodstream infections by NFGNB varied significantly within age groups, where the highest prevalence was recorded among patients at the 41 to 50. Similarly, only few studies suggest a correlation between the infection due to NFGNB and age. A study, done in Eastern part of India in 2013 revealed that majority of the patients (45%) were adults and above 45 years, which is similar to this current study.14

The highest number of isolates were Acinetobacter boumannii, comprises 27.69%. Acinetobacter boumannii has emerged as an important opportunistic pathogen in healthcare systems. As it hard to desiccate, so difficult to eradicate and has numerous intrinsic and acquired mechanisms of drug resistance. Thus this organism possesses a great threat to the clinician as well as to microbiologists. These organisms found extensively in nature and are able to alive in environment. They can stay alive within disinfectants and can create problem in health care facilities spreading by cross contamination and causing to blood stream infections.16

Stenotrophomonas maltophilia was the second common isolates (21.53%). Stenotrophomonas maltophilia is water borne organisms and recently emerged as an important opportunistic pathogen in debilitated host. They are enraging as a known cause of infection in the nosocomial settings.

The isolates of this emerging pathogen from blood is quite difficult to interpret as primary pathogen. However if this isolate yields from a site which is supposed to be sterile, such as from blood, drain tip or CVP tip, then this isolate represents as true or primary pathogen. Muder et al., report same kind of study where he was reported a series of 91 patients with Stenotrophomonas maltophilia bacteraemia, among them 56% did not reveal any clinically apparent portal of entry but 84 % of these individuals had central venous catheter in place.17 In 2007 Gautam et al., isolated 22 Stenotrophomonas maltophilia. Out of which 13 were from the blood samples of bacteraemia patients and 9 were from respiratory isolates.18

In this study, Pseudomonas spp was another common organism causing BSIs. Pseudomonas are ubiquitous in nature as saprophytes. Earlier it is believed to be non pathogenic. But recently they have emerged as primary opportunistic pathogens in hospitalized patients as well as immunocompromised patients and
responsible for causing variant infections including BSIs. They are very hard to desiccate, difficult to eradicate and has numerous intrinsic and acquired mechanisms of drug resistance. They can stay alive within disinfectants and can create problem in health care facilities spreading by cross contamination. The abuse and the unjudicial practice of antibiotics are responsible for the burgeoning resistance of commonly used antibiotics towards Pseudomonas. More over the multidrug resistance among these organisms makes the treatment of this infection difficult and expensive.\textsuperscript{19}

*Burkholderia cepacia* complex (BCC) found in many niches of both natural and clinical environments BCC is emerging as an important cause of morbidity and mortality in hospitalized patients because of high intrinsic antibiotic resistance, such as aminoglycosides, chloramphenicol and polymyxins. An upsure of septicaemia due to BCC is documented in various studies.\textsuperscript{18}

In our study from 65 NFGNBs we have isolated 3 isolates of *B. cepacia* and one isolate of *Burkholderia multivorans* from the blood taken in BACT/ALERT 3D SYSTEM bottle. The patients was diagnosed with sepsis and admitted in the ICU and the central venous line was in situ. Similarly, in 2006-2007 Gautam et al., isolated 39 isolates of BCC from various specimens. Out of these 39 total isolates, 30 isolates of BCC were obtained from 8601 blood cultures, accounting for 0.35%.\textsuperscript{18}

In this current study we have yielded 2 isolates of *Sphingomonas paucimobilis* from blood samples. These isolates were obtained from the blood cultures of two young patients who were admitted in ICU and female medical ward for a long tenure with the diagnosis of septicaemia. We have isolated only one isolates of *Moraxella* group from the the central venous tip of a young female, admitted in ICU with the diagnosis of septicaemia.

The risk factors associated with this pathogen are intensive care admission, prolonged hospitalization, on mechanical ventilation, presence of central venous catheter, indwelling catheters, orthopaedic implants, unjudicial use of broad spectrum, antibiotics and comorbid conditions. These predisposing factors accelerate the occurrence of the blood stream infection due to these organisms.

These NFGNB are posing a great threat to human race as they are resistant to routinely used antibiotics. The abuse and the unjudicial practice of antibiotics are responsible for the burgeoning resistance of commonly used antibiotics towards NFGNB. The resistance to antimicrobials is increasing in recent years and almost resistance to all commonly used antibiotics. More over the multidrug resistance among these organisms makes the treatment of this infection caused by NFGNB difficult and expensive.

*Pseudomonas aeruginosa* shows a good sensitivity to Imipenem (86.77\%) which is almost similar to the study by Patel et al., who reported 94\% sensitivity to this drug.\textsuperscript{15} A study by Rit et al., reported that *P. aeruginosa* were highly susceptible to Colistin (100\%), Imipenem (91.8\%) and Amikacin (69.3\%).\textsuperscript{14} In my study similarly Colisti (100\%), Imipenem (86.77\%) and Amikacin (64.02\%) revealed the same findings. The isolates of *P. aeruginosa* were sensitive to and Ciprofloacin (57.67\%), in comparison to this study another study by Patel et al., revealed a very low susceptibility rate to Amikacin (39.6\%) and Ciprofloacin (16.53\%).\textsuperscript{15} Here we found a good sensitivity to Gentamicin (57.14\%) unlike this current study, Rit et al., reported only 23.76\% of susceptibility to Gentamicin.\textsuperscript{14} In this study 61.37\% was
susceptible to Piperacillin, similarly a study by Juyal et al., revealed 52.13% sensitive to this drug. Ciprofloxacin and Ceftazidime both accounting for 57.67%. Unlikely, a study by Patel et al., who reported only 24.6% susceptibility rate to Ceftazidime. Where as Carbenicillin (44.44%) and Ceftazidime-tazobactam (17.98%) reveals quite a low sensitivity to this organism. In comparison to my study by Juyal et al., revealed 69.15% sensitivity to Piperacillin-tazobactam. 

Imipenem (88%) show the highest sensitivity to Pseudomonas flurosence, similarly a study by Rit et al., reported 100 % sensitivity to Imipenem. In this current study Amikacin and Ceftazidime each of them show 66.7% sensitivity to Pseudomonas flurosence. Similar to this study, Rit et al., revealed 66.66% sensitivity to Amikacin. However Rit et al., revealed a low sensitivity rate to Gentamicin (33.33%) and Ciprofloxacin (33.33%) where as in this study Gentamicin and Ciprofloxacin accounting for 71.4% and 61.9% susceptibility. Here Piperacillin accounting for (71.4%).

Imipenem (89.65%) shows the highest sensitivity for Pseudomonas putida, almost similar to this study, a study by Patel et al., revealed 100% susceptibility to Imipenem. The other isolates of P.putida show a moderate susceptibility pattern towards Amikacin (68.96%) and Ciprofloxacin (62.06%), where as in comparison to them study by Patel et al., revealed 100% sensitivity to Amikacine and Cefipime. Gentamicin show 58.62%, Ciprofloxacin, Ceftazidime and Piperacillin an reveal a susceptibility rate of, 62.1% 55.17% and 51.72% respectively.

Imipenem (95.23%) shows the highest sensitivity for Pseudomonas stutzeri, Gentamicin reveals 90.47%. Ceftazidime shows 85.71%, Ciprofloxacin and Amikacin reveal 80.95% individually.

The isolates of A.boumannii showed 60% were sensitive to Imipenem. Almost similar susceptibility of Imipenem (68.06%) was reported by Juyal et al., In comparison to this Rit reported a good sensitivity to Imipenem (90%). Where as another study by Parimal et al., revealed 72.9% sensitivity to Imipenem. In this study we have reported 52.2% susceptibility to Chloramphenicol and 48.9% to Gentamicin. In contrast to this study, another study by Rit revealed low susceptibility to Chloramphenicol (28%) and Gentamicin (24%). Where as Juyal et al., reported exactly similar sensitivity to Gentamicin (48.61%). Close to it, in this study Amikacin and Norfloxacin each comprises of 47.8%, in comparison to this study Rit et al., reported 62% susceptibility to Amikacin. In comparison to this Parimal et al., revealed a low sensitivity to Amikacin (38.8%). However Juyal et al., reported a good sensitivity to Amikacin, accounting for 73.61%. In this study Ceftazidime shows a bit low sensitivity pattern, accounting for 37.8%.Similarly a study by Rit K, reported 28 % sensitivity to Ceftazidime.

Stenotrophomonas maltophilia revealed 100% sensitivity to Colistin, followed by Ertepenam 96.65%, next to it was Ceftazidime and Ofloxicin, both accounting for 94.12%, Ciprofloxacin (88.23%), Gentamicin (84.12%). Whereas, a study by Juyal et al., reported only 16.67% susceptibility to Gentamicin and almost resistant to Ceftazidime. However another study by Rit et al., reported a good susceptibility to Ceftazidime (66.7%) In this study 70.58% susceptibility rate was for Cefotaxim. In this current study Amikacin and Imipenem reveal a good sensitivity, accounting for72.36% and 79.71% sensitivity respectively, where as a study done by Juyal et al., reported almost resistant to Imipenem and Amikacin (33.33%).
Resistance to antimicrobials is common and has increased over the years among NFGNBs. Multidrug resistance among these organisms makes the treatment of infections caused by them, difficult and expensive. A large scale use of the third- generation Cephalosporins like Cefotaxime, Ceftriaxone, and Ceftazidime has led to the evolution of newer betalactamases such as the ESBLs. ESBLs are plasmid mediated enzymes that hydrolyze the oxyimino β lactams and the Monobactams (Aztreonam) but have no effect on the Cephamycins (Cefoxitin, Cefotetan) and the Carbapenems (Imipenem). \textsuperscript{21, 22, 23} Being plasmid mediated, they can be easily transferred from one organism to another. \textsuperscript{23} A rapid detection of ESBL and MBL positive isolates is necessary to control infection and to prevent their dissemination. In this study we have performed Combined Disc Diffusion Test (CDDT) to However in this study among the total 65 isolates of NFGNB, 20 isolates (30.77%) were multidrug resistance (MDR). However, amidst these 20 isolates 11 (55%) were ESBL- producers and rest (45%) were MBL-producers. Among the total 65 isolates of NFGNB, 20 isolates (30.77%) were multidrug resistance (MDR). However, amidst these 20 isolates 11 (55%) were ESBL producers and rest (45%) were MBL-producers.

In conclusion a large number of NFGNB are isolated as primary pathogen, which has a potential to cause BSIs. The remarkable thing about all these isolates is that these isolates obtained from the typical cases of HAI. These organisms have possibly come from inanimate objects like ventilator, humidifier, wash basin and from diluted disinfections. Most of the patients had high risk factors, like comorbid conditions (DM, hypertension, post renal dialysis, cardiac disease, tuberculosis etc) prolonged hospitalization, indwelling catheters and orthopaedics implants in situ and unjusticial use of antibiotics. Although BSI rates declined over time, but BSI had high mortality and these NFGNB pathogens exhibited substantial antimicrobial resistance. Most effective antibiotics are colistin, imipenem, ertepenam, amikacin and gentamicin. There is a wide spread variability of antibiotic profile in common hospital for these pathogens. the antibiotic susceptibility can change from hospital to hospital set up and there may be a gross geographical variation. So, it is imperative that every hospital should monitor a proper antibiogram profile for these isolates from time to time to serve as a basic empirical therapy to prevent the development of MDR cases. Treating these pathogen should be based on the laboratory data after identifying the proper causative agents and antibiotic susceptibility result. Minimized the use and abuse of antimicrobial agents, proper surveillance of antibiotic panel, strict infection control measures and even simple yet proper hand washing method and by using disinfections of inanimate objects, can prevent the emergence \textit{Acinetobacter} and can reduce the rate of MDR strains.

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