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Prevalence of Acinetobacter baumannii in Diyala Governorate from Different Clinical Sources and its Antibiotics Profile Analysis

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

Acinetobacter baumannii from various clinical sources (Blood, urine, CSF, Ear swab, burn swab, wound swab, sputum and vaginal swab) were isolated from patients were attending to Baquba Teaching Hospital and Al-Betool Teaching Hospital. A total of (27) isolates of A. baumannii bacteria out of (200) samples were isolated from various clinical sources. Diagnoses all *Acinetobacter baumannii* bacteria isolates and AST test by Vitek 2 advice. The maximum isolates were obtained from blood samples 10(37.03%), urine samples 4(14.8%), CSF samples 4 (14.81%), burn swab

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(14.81%), sputum swab 2(7.4%), while 1(3.7) for wound swab, Ear swab, and vaginal swab. All isolates were tested for their resistance to (23) different antibiotics and the results showed that resistant to Amoxicillin, Amoxicillin/Clavulanic acid and Trimethoprim were (96.2%) but lesser to Ticarcillin, Ticarcillin / Clavulanic Acid, Piperacillin/Tazobactam, Ceftriaxone, Imipenem, Ciprofloxacin and Levofloxacin were 88.8%.

Keywords: Acinetobacter baumannii; antibiotic susceptibility test; vitek 2 system and antibiotic.

1. INTRODUCTION

A.baumannii is a Gram-negative bacterium, aerobic and non-motile rod. This opportunistic bacterium has emerged as a significant concern in human health, responsible for a variety of infections, including pneumonia, meningitis, septicemia, and urinary tract infections. It possesses several virulence factors, such as the ability to form biofilms, adhere to surfaces, invade host cells, acquire iron, and induce host cell death [1]. A. baumannii easily attaches to both biological and non-biological surfaces, where it can form biofilms. This characteristic is significant for many pathogens as it aids in the colonization of prosthetic materials and plays a role in drug resistance and evasion of the host immune system in vivo [2]. A. baumannii has developed resistance to many of the most effective antimicrobial agents, leading to a significant increase in morbidity and mortality rates, particularly in intensive care units across various countries [3]. A. baumannii is a significant multidrug-resistant (MDR) opportunistic pathogen commonly associated with nosocomial infections, largely due to its remarkable ability to develop resistance to various groups antibiotics. Multiple of mechanisms contribute to the acquisition of this multidrug resistance [4]. Resistance in A. baumannii strains exhibiting the MDR phenotype is attributed to a diverse array of genes responsible for antibiotic resistance. encompassing both intrinsic and acquired mechanisms [5]. There are three mechanisms of antibiotic resistance in A. baumannii: First, resistance can arise by reducing membrane permeability or increasing antibiotic efflux, thereby preventing access to the target. Additionally, bacteria may employ genetic changes or post-translational modifications to protect the drug target. Finally, antibiotics can be inactivated through hydrolysis directly or modification processes [6]. This pathogen has developed multidrug resistance (MDR) in recent vears, mainly due to the widespread overuse of and antibiotics inadequate antibiotic management. Prolonged hospital stays, use of

mechanical ventilation catheters, and are associated with MDR isolates, and invasive infections are more likely to occur in immunocompromised individuals and those who are critically ill [7,8]. The present study therefore aimed to determine the prevalence of Acinetobacter baumannii in Divala governorate from different clinical source and to determine the sensitivity and resistance of isolates to antibiotics.

2. MATERIALS AND METHODS

In the current study, 27 A. baumannii isolates emploved. Samples were gathered were between January and July of 2024. Blood. urine. CSF, burns, wounds, blood cultures, and sputum were among the items that were taken from Baquba Teaching Hospital and Al-Betool Teaching Hospital. All obtained samples, with the exception of blood, were grown immediately on blood agar and MacConkey agar and incubated for twenty-four hours at 37°C. The blood specimen must be placed in blood culture bottles that include nutrients that encourage the growth of aerobes bacteria and then cultured on blood agar and MacConkey agar and then incubated for further 24 hours at 37°C.All bacterial isolates were tested by Vitek 2 system. Bacterial identification based on the biochemical reactions between the bacterial suspention and the culture media existed in the GN-ID Cards loaded manually into the VITEK system. The susceptibility of A. baumannii isolates against 23 antibiotics were determined by Vitek 2 system by using Antibiotics Susceptibility Gram-negative (AST-GN).

3. RESULTS

In this study a total of (200) samples were obtained from from (Blood, urine, CSF, Ear swab, burn swab, wound swab, sputum and vaginal swab as shown in Table (1).

In this study most of *A. baumannii* isolates were highly resistant to the tested antibiotics as shown in Table (2). The high resistance of *A*. *baumanniii* isolates to antibiotics shown in this study, 26 isolates were resistance 3 antibiotics (96.2%), 25 isolates were resistance 4 antibiotics with the rate (92.5%), while 7 isolates were resistance 24 antibiotics with the rate (88.8) as shown in Table(2).

Resistance to broad-spectrum antimicrobial agents in *A. baumannii* is now an emerging issue globally. In the present study, Tigecycline was found the most effective antimicrobial against isolated *A. baumannii* strains, with a 40.7% sensitivity rate, followed by trimethoprim-sulfamethoxazole bv 37.03%. Minocycline and Colistin whth a 33.3% while showed а high resistance rate to Amoxicillin , Amoxicillin/ Clavulanic acid and Trimethoprim by (96.2%), Cefuroxime, Cefixime, Cefotaxime and Ceftazidime by (92.5%), Ticarcillin, Ticarcillin / Clavulanic Acid, Piperacillin, Ceftriaxone, Ciprofloxacin and Levofloxacin (88.8%), and Cefepime, Imipenem, Meropenem, Gentamicin, Piperacillin, Tobramycin which exhibited the low level of antimicrobials effectiveness against A. baumannii strains.

Type of specimen	Total specimen	Number of isolates	%
blood	30	10	37.03
Urine	55	4	14.81
CSF	15	4	14.81
Ear swab	5	1	3.7
Brun swab	30	4	14.81
Wound swab	45	1	3.7
Sputum swab	15	2	7.4
vaginal swab	5	1	3.7
Total	200	27	100

Table 2. Antibiotic susceptibility test test on A. baumannii isolates

AB	Acinetobacter baumannii bacteria (27 isolated)	
	S(100%)	R(100%)
Amoxicillin	1(3.7)	26(96.2)
Amoxicillin/Clavulanic acid	1(3.7)	26(96.2)
Ticarcillin	3(11.1)	24(88.8)
Ticarcillin / Clavulanic Acid	3(11.1)	24(88.8)
Piperacillin	3(11.1)	24(88.8)
Piperacillin/Tazobactam	6(22.2)	21(77.7)
Cefuroxime	2(7.4)	25(92.5)
Cefixime	2(7.4)	25(92.5)
Cefotaxime	2(7.4)	25(92.5)
Ceftriaxone	3(11.1)	24(88.8)
Ceftazidime	2(7.4)	25(92.5)
Cefepime	4(14.8)	23(85.1)
Imipenem	3(11.1)	24(88.8)
Meropenem	4(14.8)	23(85.1)
Gentamicin	5(18.5)	22(81.4)
Ciprofloxacin	3(11.1)	24(88.8)
Levofloxacin	3(11.1)	24(88.8)
Tigecycline	11(40.7)	16(59.2)
Minocycline	9(33.3)	18(66.6)
Colistin	9(33.3)	18(66.6)
Tobramycin	6(22.2)	21(77.7)
Trimethoprim	1(3.7)	26(96.2)
Trimethoprim/Sulfamethoxazole	10(37.03)	17(62.9)





Fig. 1. Increase Imipenem resistance of A. baumannii clinical isolates in Iraq



Fig. 2. Increase Colistin resistance of A. baumannii clinical isolates in Iraq

These local investigations show that multidrug resistant *A. baumannii* is becoming more common in Iraqi hospitals. One such example is Imipenem (Fig. 1) and the other example is Colistin (Fig. 2); the reason for these discrepancies in the results might be the overuse of antimicrobial medications in the hospitals over the past few years (8).

4. DISCUSSION

The results shown in Table (1) appeared that from the total 27 isolates 10(37.03%) from blood, 4(14.81%) isolated from Urine, CSF, Brun ,2(7.4%) from sputum respectively. The lowest number and percentage of *A. baumannii* was Wound, Ear, vaginal swabs 1(3.7%). Urine is often only detected when an indwelling urinary catheter is present. While it is infrequently linked to pyelonephritis and urosepsis, it is seldom invasive and typically just affects the lower urinary tract [9,10]. This result was compared to the result obtained by [11,12] and [13] have isolated A. baumannii from urine rate reaching to (16,10% and 11.5%) respectively. The results of this study shows that A. baumannii bacteria can be isolated from burn and this result agreed with previous study [14], who has isolated these bacteria from burn samples at rate of (17%). Around the world, burns are a serious public health problem and one of the most prevalent and destructive types of trauma [14,15]. The presence of an ample amount of oxygen in the respiratory tract, especially the lower part of it, encourages bacteria to settle and invade the region because it is a forced wind [16]. A bacteriological study stated that more than 60% of hospital-acquired pneumonia infections are caused by Negative bacteria including A. baumannii the reason for this may be attributed to their ability to blind adhesion to epithelial cells in the respiratory tract as a result of inhibitory substances such as bacteriocins, was as the presence of mucous matter [17].

Because A. baumannii may form biofilms in the skin and cause infections in soft tissues, it is present in wounds and occlusive dressings [7]. Because many bacterial species have particular receptors for these chemicals, bacteria can readily colonize the surfaces of burn wounds. The reason for the difference in the isolation ratio is due to the number of samples taken the time of collection of the samples, the environment from which the samples were isolated, the health conditions in which the patients lived, and the length of time they stayed in Hospitalization, indiscriminate use of antibiotics excessively and the difference in the number of samples taken for a study and variance. Numerous dangerous nosocomial infections, including as urinary tract infections, meningitis, wound infections, bloodstream infections, and ventilator-associated pneumonia, can be caused by A. baumannii. individuals in the high-risk groups include individuals who are mechanically ventilated, have indwelling foreign devices, are debilitating patients, and are hospitalized to intensive care units (ICUs) [18,19].

The results of the current study demonstrated that the highest resistance to almost all β-lactam antibiotic classes under study was as follows cephalosporins (Amoxicillin, Amoxicillin/ Clavulanic acid .Cefuroxime. Cefixime Cefotaxime, Ceftriaxone, Ceftazidime, Cefepime, Piperacillin ,Oxacillin) These findings were close to local studies related to A. baumannii isolates by Adnan S et al. [20,21], who found that (100%) of A. baumannii isolates in Iraqi hospital environment resisted Oxacillin, Ceftazidime and Cefepime. Another study in Iran hospital by Maryam R et al. [22] found (100% resistance rate to (Cefoxitin, Cefoxitin, Oxacillin, Ceftazidime, Cefepime). The three main types of resistance mechanisms are: (1) enzymes that deactivate antimicrobials; (2) bacterial targets are less accessible; or (3) mutations that alter targets or cellular activities. Regarding the first group, a variety of β-lactamases found in Acinetobacter species hydrolyze and provide resistance to cephalosporins, carbapenems, and penicillins. Chromosome-encoded AmpC cephalosporinases provide resistance to broad-spectrum cephalosporins [23].

As the isolates of bacteria *A.baumannii* resistance to Gentamicin in the percentage

(81.4%). The results are consistent with [24]. which was reached by the resistance rate (62%). While the resistance to isolates of the antibiotic Imipenem (88.8%), Meropenem (85.1%). This close with the results of the current study, which gave a100% resistance to this antibiotic.

The results of Ciprofloxacin and Levofloxacin are agreed with [25] research, which was reached by the resistance rate (85%), and also agreed with [26] research. The resistance rate was (80%), Another local study revealed that A. baumannii clinical isolates were 100 % resistant to Ciprofloxacin [27]. resistance of A. baumannii for Tobramycin, were (77.7%) consistent with the study of [28]. that showed 100% resistance for Tobramycin.

Because of its low rates of resistance, tigecycline could be the treatment of choice for managing infections at this facility. Although The results of resistance to trimethoprim /sulphandmethoxole also had a low resistance rate in our present study, were close to the results reached by the researcher in the research [24] and also close to the results of the research of [29] who recorded that the Acinetobacter baumannii resist to a wide range of antibiotic including trimethoprim / sulphandmethoxole in the percentage (81%). fewer patients were treated by this antibiotic, as its side effects of nephrotoxicity and neurotoxicity [30].

These bacteria are resistant to nearly all antibiotics for a variety of reasons. Among these, the most significant one is the existence of efflux pump families [31]. There are several families of efflux pumps in A. baumannii, but the most significant family is the existence of Resistance. These families allow us to exclude antibiotics. heavy metals, and a host of other substances. The three pump groups that make up the nodulation cell division (RND) family are each in charge of resisting a particular kind of antibiotics. Clinical isolates of A. baumannii were reported to be 37% resistant to colistin in a previous investigation conducted in Sudan in 2015 [32]. Previous study in Iraqi in 2016 found that A. baumannii clinical isolates were 20% resistant to colistin [33]. Previous study has reported that A. baumannii from clinical isolates were 96% resistant to colistin [34]. Colistin-resistant A. baumannii may become resistant to colistin due to modifications of the outer membrane which may increase the permeability to other cell wall antimicrobial agents. Previous studies have reported that colistin-resistant A. baumannii strains were more susceptible to other antimicrobial agents than colistin-susceptible strain [35].

Previous study in Iraq in 2012 shown that resistance to Imipenem was 57.1% [36]. Previous study in Irag in 2014 shown that low resistance to Imipenem 30% [37]. Other study also in Iraq shown that One (10%) isolate was observed to be imipenem and meropenem resistant [38]. Iraqi study in 2017 found that A. baumannii clinical isolates were 68.7% and 50% Ceftazidime and Imipenem resistant to respectively [39]. A. baumannii clinical isolates in previous study were 68.7% and 90.9% resistant to Imipenem [40,41].

In present study appeared to have resistance to Cefotaxime (92.5%) compare to previous study shown that all isolates appeared to have resistance to Cefotaxime (60%) [37]. previous study has reported that most isolates of A. baumannii are considered to be resistant to Cefotaxime in rate reach to (80%) and (60%) respectively [42,43]. Other previous study found that (12.2%) of isolates were resistant to Cefotaxime [44].

5. CONCLUSION

A.baumannii has demonstrated a high level of resistance to the majority of tested antibiotics, earning it the reputation of being the most therapeutically significant Acinetobacter species globally. It has become a significant nosocomial opportunistic pathogen in hospital infection epidemics.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc) and text-to-image generators have been used during writing or editing of manuscripts.

CONSENT

It's not applicable.

ETHICAL APPROVAL

It's not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- 1. Kunz AN, Brook I. Emerging resistant Gram-negative aerobic bacilli in hospitalacquired infections. Chemotherapy. 2010; 56(6):492-500.
- Lee HW, Koh YM, Kim J, Lee JC, Lee YC, Seol SY, Cho DT. Capacity of multidrugresistant clinical isolates of Acinetobacter baumannii to form biofilm and adhere to epithelial cell surfaces. Clinical microbiology and infection. 2008;14(1):49-54.
- 3. Moulana Z, Babazadeh A, Eslamdost Z, Shokri M, Ebrahimpour S. Phenotypic and genotypic detection of metallo-betalactamases in Carbapenem resistant Acinetobacter baumannii. Caspian journal of internal medicine. 2020;11(2):171.
- 4. Da Silva GJ, Domingues S. Insights on the horizontal gene transfer of carbapenemase determinants in the opportunistic pathogen Acinetobacter baumannii. Microorganisms. 2016;4(3):29.
- 5. Fallah A, Rezaee MA, Hasani A, Barhaghi MH, Kafil HS. Frequency of bap and cpaA virulence genes in drug resistant clinical isolates of Acinetobacter baumannii and their role in biofilm formation. Iranian journal of basic medical sciences. 2017; 20(8):849.
- Wong D, Nielsen TB, Bonomo RA, Pantapalangkoor P, Luna B, Spellberg B. Clinical and pathophysiological overview of Acinetobacter infections: a century of challenges. Clinical microbiology reviews. 2017;30(1):409-47.
- 7. Pamu J. Prevalence and antimicrobial susceptibility pattern of methicillin-resistant staphylococcus aureus at port moresby general hospital, Papua New Guinea: A Retrospective Study (Doctoral dissertation, Auckland University of Technology); 2019.
- 8. Hussein N, Al-Mathkhury H, Sabbah M. Imipenem-Resistant Acinetobacter baumannii isolated from patients and hospitals environment in Baghdad. Iraqi Journal of Science. 2013;54(4):803-12.
- 9. Kb L. Incidence and risk factors for acquiring nosocomial urinary tract infection in the critically ill. J Crit Care. 2002; 17:50-7.
- 10. Avecillas JF, Mazzone P, Arroliga AC. A rational approach to the evaluation and treatment of the infected patient in the intensive care unit. Clinics in Chest Medicine. 2003;24(4):645-69.

- Rodríguez-Baño J. Cisneros JM. 11. Fernández-Cuenca F, Ribera A, Vila J, Pascual A, Martínez-Martínez L, Bou G, Pachón J, Grupo de Estudio de Infección Hospitalaria. Clinical features and epidemiology of Acinetobacter baumannii colonization and infection in Spanish hospitals. Infection Control & Hospital Epidemiology. 2004 Oct;25(10):819-24.
- 12. Rungruanghiranya S, Somboonwit C, Kanchanapoom T. Acinetobacter infection in the intensive care unit.Journal of Infectious Diseases and Antimicrobial Agents.2005;22(2):77-92.
- 13. Yaman A, Aksungur P. Antibiotic resistance in Acinetobacter species in nosocomial and outpatient infections.
- 14. Qader AR, Muhamad JA. Nosocomial infection in sulaimani burn hospital, IRAQ. Annals of burns and fire disasters. 2010;23(4):177.
- Rezaei E, Safari H, Naderinasab M, Aliakbarian H. Common pathogens in burn wound and changes in their drug sensitivity. Burns. 2011;37(5):805-7.
- Lopes BS, Gallego L, Amyes SG. Multidrug resistance profiles and the genetic features of Acinetobacter baumannii isolates from Bolivia. The Journal of Infection in Developing Countries. 2013;7(04):323-8.
- 17. MacFaddin JF. Biochemical tests for identification of medical bacteria, williams and wilkins. Philadelphia, PA. 2000;113(7).
- Akrami F, Namvar AE. Acinetobacter baumannii as nosocomial pathogenic bacteria. Molecular Genetics, Microbiology and Virology. 2019;34:84-96.
- Lerner AO, Abu-Hanna J, Carmeli Y, Schechner V. Environmental contamination by carbapenem-resistant Acinetobacter baumannii: the effects of room type and cleaning methods. Infection Control & Hospital Epidemiology. 2020; 41(2):166-71.
- 20. Adnan S. Identification of Acinetobacter baumannii Isolated from different infections and study the prevalence of antibiotic resistance in patients of baquba City. Academic Science Journal. 2024;2(2):91-104.
- 21. Al-Kadmy IM, Ali AN, Salman IM, Khazaal SS. Molecular characterization of Acinetobacter baumannii isolated from Iraqi hospital environment. New microbes and new infections. 2018; 21:51-7.

- 22. Maryam R, Golnaz YZ, Mojgan O, Malihe T, Nour A. Identification of five phylogenic groups of carbapenemase (bla OXA-23, 24, 51, 58,143) in Acinetobacter baumannii strains isolated from clinical samples in Iran by multiplex PCR.
- Tien HC, Battad A, Bryce EA, Fuller J, Mulvey M, Bernard K, Brisebois R, Doucet JJ, Rizoli SB, Fowler R, Simor A. Multidrug resistant A cinetobacter infections in critically injured Canadian forces soldiers. BMC infectious diseases. 2007; 7:1-6.
- 24. Lin MF, Lan CY. Antimicrobial resistance in Acinetobacter baumannii: From bench to bedside. World Journal of Clinical Cases: WJCC. 2014;2(12):787.
- 25. Alrifai SB, Mahmood WS, Jasem NH. Surveillance of Multidrug-resistant Iraqibacter Isolated from Patients with Urinary Tract Infection at a Baghdad Urology Center.
- Ghaima KK. Study OXA beta-lactamase Genes in Clinical Isolates of Multidrug Resistant Acinetobacter baumannii. University of Baghdad. University of Baghdad, Baghdad (PhD's thesis); 2016.
- Al-Khafaji SM. Study on capsule of Acinetobacter baumannii and its effect on Immune Response (Doctoral dissertation, PH. D. Thesis. Biology department. College of Science. Al-Mustansiriya University)
- 28. Al-Kadmy IM, Ali AN, Salman IM, Khazaal SS. Molecular characterization of Acinetobacter baumannii isolated from Iraqi hospital environment. New microbes and new infections. 2018;21:51-7.
- 29. Angoti G, Bandehpour M, Goudarzi H, Hajizadeh M, Zarringhalam Moghaddam M, Kouchaki A. Detection of efflux pump genes (adeA, adeB, adeC and abeM) in Acinetobacter baumannii isolated from hospitalize patients, North-west of Iran. Infection Epidemiology and Microbiology. 2016;2(4):8-11.
- 30. Loho T, Dharmayanti A. Colistin: an antibiotic and its role in multiresistant Gram-negative infections. Acta Medica Indonesiana. 2015;47(2).
- Monem S, Furmanek-Blaszk B, Łupkowska A, Kuczyńska-Wiśnik D, Stojowska-Swędrzyńska K, Laskowska E. Mechanisms protecting Acinetobacter baumannii against multiple stresses triggered by the host immune response, antibiotics and outside-host environment.

International Journal of Molecular Sciences. 2020;21(15):5498.

- 32. Omer MI, Gumaa SA, Hassan AA, Idris KH, Ali OA, Osman MM, Saleh MS, Mohamed NA, Khaled MM. Prevalence and resistance profile of Acinetobacter baumannii clinical isolates from a private hospital in Khartoum, Sudan. Am J Microbiol Res. 2015;3(2):76-9.
- Al-Samaree MY, Al-Khafaji ZM. Antibiogram of Acinetobacter baumannii isolated from Baghdad Hospitals. Int. J. Adv. Res. Biol. Sci. 2016;3(4):238-42.
- 34. Narjis MA, Mahdi MS. Isolation and identification of multi-drug resistance Acinetobacter baumannii isolated from clinical samples at Baghdad, Iraq. Journal of Applied and Natural Science. 2023; 15(2):663-71.
- Li J, Nation RL, Owen RJ, Wong S, Spelman D, Franklin C. Antibiograms of multidrug-resistant clinical Acinetobacter baumannii: promising therapeutic options for treatment of infection with colistinresistant strains. Clinical infectious diseases. 2007;45(5):594-8.
- Shali AA. Identification of Multidrug-Resistant Genes in" Acinetobacter baumannii" in Sulaimani City-Kurdistan Regional Government of Iraq. Asian Journal of Medical Sciences. 2012;4 (5):179-83.
- Jabur MH. Isolation of Acinetobacter baumannii from different clinical source and study some antibiotic resistant and βlactamase production. Med J Babylon. 2014;11(2):456-64.

- AL-Harmoosh RA, Jarallah EM, AL-Shamari AM. Coexistence of the blaIMP and blaSIM genes in clinical isolates of Acinetobacter baumannii IN Babylon Hospitals-Iraq. International Journal of PharmTech Research. 2016;9(7):257-64.
- 39. Al-Dulaimi AA, Al-Taai HR, Al-Bajlany SM. Virulence Factors of Acinetobacter baumannii isolated from different clinical specimens in Baquba.
- 40. Ibrahim ME. Prevalence of Acinetobacter baumannii in Saudi Arabia: risk factors, antimicrobial resistance patterns and mechanisms of carbapenem resistance. Annals of clinical microbiology and antimicrobials. 2019;18(1):1.
- 41. Mahmood NH, Al-Brefkani AM. Detection and Characterization of Carbapenem Resistant Acinetobacter baumannii Isolated from Different Clinical Specimens in Duhok Province–Iraq. History of Medicine. 2023;9(1):783-94.
- 42. Kumari AM, Routray A, Yadav D, Madhavan R. Imipenem resistance and biofilm production in Acinetobacter. drug invention today. 2013;5(3):256-8.
- 43. Nazmul MH, Jamal H, Fazlul MK. Acinetobacter species-associated infections and their antibiotic susceptibility profiles in Malaysia. Biomed Res-India. 2012;23(4):571-5.
- 44. Shareek PS, Sureshkumar D, Ramasubramanian V, Ghafur KA, Thirunarayanan MA. Antibiotic sensitivity pattern of blood isolates of Acinetobacter species in a tertiary care hospital: A retrospective analysis. American Journal of Infectious Diseases. 2012;8(1):65.

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