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3 **Prevalence of carbapenem-resistant hospital**
4 **acquired infections and their antimicrobial**
5 **susceptibility pattern in a tertiary care hospital**
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14 **ABSTRACT**

Background: The emergence of antimicrobial-resistant bacteria has become a public threat, creating a burden on medical care in hospitals. Carbapenem-resistant organisms are a source of both community-acquired and healthcare-acquired infection that poses a substantial hazard to public health. This study aimed to conclude the prevalence of carbapenem resistance gram-negative bacteria from a clinical specimen at Index Medical College -Indore.

Methodology: This study was conducted in the Department of Microbiology, at Index Medical College, Indore, between January 2020 and January 2022. The isolates were subjected to antimicrobial susceptibility tests by Kirby Bauer's disk diffusion test most of the isolates were resistant to beta-lactam drugs, cephalosporin's and aminoglycosides. These isolates were further confirmed by phenotypic detection using the Modified Hodge test, Modified carbapenem Inactivation, Combined disc diffusion test and Double Disk Synergy.

Results: The percentage distribution of health-associated infection show highest resistance in both urinary tract and respiratory tract infection, followed by skin & soft tissue infection and least in septicemia and other health associated infection. Highest percentage of resistance was seen in the age group between 21-30 and the least in less than 10 years with a statistical significance of p -value=.00001. The most common isolates recover was E.coli in Enterobacterials and in Non fermenter it was Pseudomonas aeruginosa and Acientobacter. The sensitivity of MHT, mCIM, CDDT, and DDST within CI 95% were 74%, 95%, 84%, and 95% respectively. The overall prevalence of carbapenem resistance is 17.75%.

Conclusion: The production of carbapenemase is the major mechanism underlying carbapenem resistance around the world and represents a great health concern. More knowledge is needed to control resistant genes and resistant organisms and their dissemination. There is an urgent need for global collaboration to plan valid strategies to prevent the spread of carbapenemase and the development of new antimicrobial molecules.

14
15 *Keywords: Carbapenem resistance; Metallo beta-lactamase; Antimicrobial susceptibility;*
16 *Phenotypic detection; Nosocomial Infections*
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20 **1. INTRODUCTION**

21 The most versatile family of beta-lactamase, are carbapenemase [1] Carbapenemase,
22 enzymes hydrolyze almost all beta-lactamase and does not work against inhibitor. [2, 3].

23 Carbapenem-resistant gram-negative bacteria are difficult to treat infections in hospitalized
 24 patients. It leads to high mortality [4] and is the last resort for salvage treatment of multidrug-
 25 resistant Gram-negative bacteria. Carbapenem becomes a life threatening to the survival
 26 of critically ill patients, with 50% mortality [5]. There is an increased alert of prevalence in
 27 multidrug-resistant organism which causes serious nosocomial infections. Globally
 28 Carbapenem-resistant gram-negative organisms are the main cause of nosocomial infection.
 29 [6]. This is a cross-sectional study performed to determine the prevalence of carbapenem-
 30 resistant Gram-negative bacilli isolated from patients admitted in wards tertiary care
 31 hospitals in Central India. We compared different methods for the detection of
 32 carbapenemase and Metallo beta-lactamase (MBL).

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35 2. MATERIAL AND METHODS

36 This study is a cross-sectional study, conducted in the the Bacteriology section of the
 37 microbiology laboratory of Index Medical College ,Hospital and Research Center, Indore
 38 (M.P.) from January 2020 to January 2022 A total of 246 clinical samples from patient
 39 admitted for more than 48 hours were collected from different clinical department.

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42 2.1 Antibiotic Susceptibility Testing

43 Antibiotic susceptibility testing of Gram-negative isolates was done by Kirby Bauer's disc
 44 diffusion method using the following antibiotics Imipenem, Meropenem, Ertapenem
 45 Ciprofloxacin, Amikacin, Piperacillin/tazobactam, Ceftazidime, Gentamicin, Ampicillin,
 46 Cefazolin, Tobramycin, Tigecycline Cefepime, Ceftriaxone, Amoxiclav, Cefotaxime, Colistin
 47 in Mueller Hinton Agar according to CLSI guidelines (7).

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50 2.2 Phenotypic detection of carbapenem and Metallo β lactamase production

51 The detection was done by the Modified Hodge test [8], Modified carbapenem Inactivation
 52 Method [8], for phenotypic detection of MBL, Combined disc diffusion test [9] and Double
 53 Disk Synergy test (10) was done.

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56 3. RESULTS

57 A total of 246 clinical samples were collected during the study period from Jan 2020 to Jan
 58 2022 from various clinical departments from which 129 (52.43%) were positive for different
 59 isolates. From the 129 positive isolates, 22 (8.94%) were gram-positive bacteria and 107
 60 (43.49%) were gram negative bacilli. From 107 gram-negative isolates 80.37% were
 61 *Enterobacteriales* and 19.62% were non- fermenter. *E.coli* (32.55%) followed by, *Klebseilla*
 62 *pneumoniae* (17.5%), *Citrobacter species* and *Pseudomonas aeruginosa* (8.52),
 63 *Acientobacter Species* 7.55% , *Proteus mirabilis* 4.56% and *Enterobacter species* 3.87%.
 64 On performing Kirby Bauer disc diffusion method 59.09% of male patients were sensitive
 65 and 63.15% were resistant to carbapenem where as 40.90% were sensitive and 36.84%
 66 were resistant to carbapenem. The statistical analysis of p-value = 0.00001 for both
 67 carbapenem-resistant and carbapenem sensitive as shown in Table 1

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72 **Table 1** Gender-wise distribution of Gram-Negative carbapenem-resistant and Gram-Negative carbapenem-
 73 sensitive isolates

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Gender	Carbapenem resistant	Carbapenem sensitive	Total	p-value
Male	12(63.15)	52(59.09)	64(59.81)	0.00001
Female	7(36.84)	36(40.90)	43(40.18)	0.00001
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Percentage distribution of carbapenem-resistant drug show 21% resistance by Imipenem and Meropenem followed by Ertapenem 11%. In coexisting resistance was seen as Meropenem plus Ertapenem, Imipenem plus Ertapenem 5% each, Imipenem plus Meropenem 11% and 26% were resistant to all three drugs shown in Fig 1. The prevalence of carbapenem resistance was 17.75%. The percentage of sensitivity of Imipenem was 79%, Meropenem 79%, and Ertapenem 89% respectively.

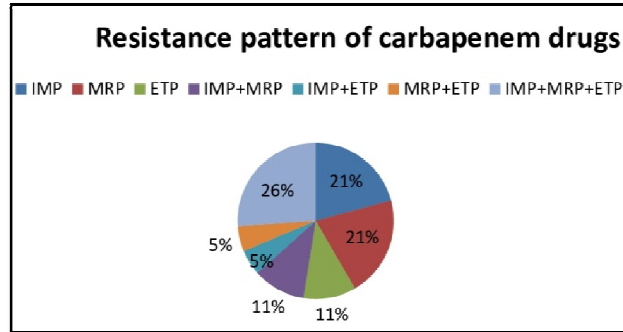


Fig 1 Resistance pattern of carbapenem drugs

The antimicrobial sensitivity pattern highest sensitivity of Colistin (89%) followed by Tigecycline (58%), Amikacin (21%), Ciprofloxacin and Amoxiclav (16%) Ampicillin, Cefazolin, Tobramycin, Piperacillin/Tazobactam, Levofloxacin (11%), Cefepime and Gentamicin (5.2%), Cefotaxime, Ceftriaxone and Ceftazidium with no sensitivity. Ceftriaxone, Cefotaxime and Ceftazidium show 100% resistance followed by Gentamicin and Cefepime (95%), Ampicillin, Cefazolin, Levofloxacin, Tobramycin, Piperacillin/tazobactam (89%), Ciprofloxacin and Amoxiclav (84%), and least resistance was observed in Tigecycline (42%) Fig.2

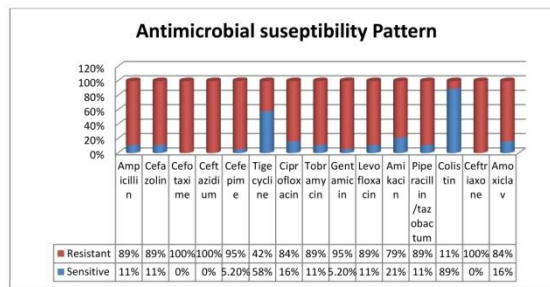


Fig 2. Antimicrobial susceptibility test

Highest percentage of nosocomial infection was seen in urinary tract infection (65.4%) followed by skin & soft tissue infection (15.88%), respiratory tract infection (10.28%), (4.67%) septicemia and least infection was seen in other health associated infection (3.73%). The percentage distribution of health associated infection has highest carbapenem resistance in urinary tract and respiratory tract infection account of (31.57%) followed by skin & soft tissue infection (15.78%), and least about (10.52%) was seen in septicemia and other health associated infection. Table 2

Table 2 Types of hospital acquired infection in relation to gram-negative carbapenem-resistance and gram-negative carbapenem-sensitive

Type of HAI	GNR N(%)	CRGNR N(%)	p-value
Urinary tract infection	70(65.4)	6(31.57)	.000001
Skin & Soft tissue infection	17(15.88)	3(15.78)	.000943
Septicemia	5(4.67)	2(10.52)	.004893
Respiratory tract infection	11(10.28)	6(31.57)	.001058
Other HAI's	4(3.73)	2(10.52)	.024271

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GNR-Gram Negative , CRGNR-Carbapenem Resistant Gram Negative

Age wise distribution accounts for sample collected from patients have highest percentage of 23.17% age group 41-50yrs, followed by 21.95% , 31-40 yrs 19.1%, 21-30 year ,15.44% more than 50years, 32 (13%) and 7.31% age less than 10year. In overall age group positive isolates were observed in age group 21-30 year , 28.68%, followed by 18.6%% (31-40yrs), 17.05%(10-20yrs), 15.5% more than 50yrs, 12.43%(41-50yrs) and 7.75% less than 10 yrs. The overall gram negative bacilli were highest in age group between 21-30 (28.9%) followed by 31-40 (20.5%), 10-20 (15.8%), more than 50 (15.3%), 41-50 (13.0%), less than 10 years (6.5%) respectively. Overall carbapenem resistance is highest in age group 21-30 (42.1%) followed by 10-20 age group (21.05), more than 31 (10.52%) and less than 10 yrs (5.26%) Overall p-value= .000857. Table 3

Age group (in Years)	Total sample N=256(%)	GNR N=107 (%)	CRGNR N=19(%)	p-value
less than 10	18(7.31)	7 (6.5)	1(5.26)	.016157
10_20	32(13)	17(15.8)	4(21.05)	.000538
21-30	47(19.1)	31(28.9)	8(42.1)	.00001
31-40	54(21.95)	22(20.5)	2(10.52)	.00001
41-50	57(23.17)	14(13.0)	2(10.52)	.0001214
more than 50	38(15.44)	16(15.3)	2(10.52)	.000159

Table 3 Age wise distribution of isolates

Gram negative bacteria were isolated from the different department; the maximum isolates were medicine 51% followed by surgery 21%, intensive care unit 14%, obstetrics & gynaecology 7% and pediatric 6%. Among the different departments the maximum carbapenem-resistant gram-negative bacteria was present in medicine 31.57% which was followed by surgery 26.31% , intensive care unit 21.05% and obstetrics & gynecology and Pediatric showing same percentage of resistance 10.52% .Among the gram-negative bacteria the maximum bacteria were isolated from Urine 65.4% followed by Pus 15.8%, Endotracheal aspirate 6.5%, Blood 4.6%, Sputum and other health associate infection [HAI's] 3.7%. Among them maximum carbapenem resistant GNR were isolated from Urine 31.5% followed by Endotracheal aspirate 21%, Pus 15.7% and 10.5% Blood, Sputum and other health associate infection (Table 4) The statistical significant difference of p-value=.00001.

Department	Blood	Urine	Pus	Sputum	ET	Other HAI's	GNR
Medicine	0	53	0	2	0	0	55
Surgery	0	8	12	1	0	2	23
Obstetrics & Gynaecology	0	4	3	0	0	1	8
Pediatric	4	0	1	0	0	1	6
Intensive care unit	1	5	1	1	7	0	15
Total GNR	5(4.6)	70(65.4)	17(15.8)	4(3.7)	7(6.5)	4(3.7)	107
CRGNR	2(10.5)	6(31.5)	3(15.7)	2(10.5)	4(21.0)	2(10.5)	19

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Table 4 Department wise distribution of GNR and CRGNR
ET- Endotracheal Aspirate, HAI- Hospital associated infection,
GNR-Gram negative rod,
CRGNR- Carbapenem resistant gram negative Rods

Among the nosocomial infection, the maximum number of organisms were isolated from urinary tract infection which include *E.coli* 423% followed by *Klebseilla pneumoniae* 20%, *Citrobacter species* 13%, *Pseudomonas aeruginosa* 10%, *Enterobacter species* 8%, *Proteus species* 6 % and *Acientobacter species* 1%. The second highest infection is caused by skin soft tissue infection *E.coli* contribute the highest prevalence in causing soft & skin tissue infection species and *Pseudomonas aeruginosa* 6% with no isolates from *Enterobacter species* and *Acientobacter species* isolated. Among the respiratory tract infection accounts for the third highest infection, responsible organism include *Acientobacter species* 63% followed by *Klebseilla pneumoniae* 18%, *E.coli* and *Proteus species* each 9%. Fourth highest nosocomial infection was septicemia and other hospital-associated infection, which was caused by 60%, *Klebseilla pneumoniae* followed by *E.coli* and *Pseudomonas aeruginosa* 20%, respectively. Among the least nosocomial findings were from other hospital-associated infections in which 50% of isolates were from *Pseudomonas aeruginosa* and *Acientobacter Species*. Carbapenem resistance in *E.coli* and *Klebseilla pneumoniae* 21.5% was the highest among the *Enterobacterials*. The least carbapenem resistance was seen among *Enterobacter species* and *Proteus species* 5.26%. Carbapenem resistance among non-fermenter *Pseudomonas aeruginosa* and *Acientobacter species* were the same about 15.7%.

Among the phenotypic methods for the detection of Carbapenem were done by MHT and mCIM. For Metallo beta lactamase detection was done by CDDT and DDST. The sensitivity of MHT, mCIM, CDDT, and DDST within CI 95% were 74%, 95%, 84%, 95% respectively. The overall prevalence of carbapenem resistance is 17.75%.

5. DISCUSSION

The overall prevalence of carbapenem-resistant in this study was 17.75%. Similarly, the study was found with the prevalence of 18%, 17%, 17.32% respectively [11, 12, and 13]. In contrast Mulla S et al [14] and Mate et al. [15] show a prevalence of 30%. The prevalence of carbapenem-resistance in different parts of India varies from 14-69%. The prevalence of carbapenem-resistant from various countries was found to be around 36% in Egypt [16], 13.6% and 37.9% in Iran [17, 18], 56% in Pakistan [19], 24.6% in China (20), 19% in Algeria [21], 2.82% in Turkey [22], 86.3% in Tunis [23], 5.99% in Morocco [24] and 2.9% in Ghana [25], 14.6%, 65%, 30%, in India [26] 27.1% in Ethiopia (27), 0.22% in Germany [28].

On performing Kirby Bauer disc diffusion method 59.09% of male patient were sensitive and 63.15% were resistant to carbapenem where as 40.90% were sensitive and 36.84% were resistant to carbapenem with the statistical significance of *p*-value 0.00001 ; which was similar to T.V. Parimala 50.90% [29] Satyajeet K. Pawar et al., 65.3% [30] and Namitha Thomas [31]

In this study overall carbapenem resistance is highest in age group 21-30 (42.1%) similar to Namitha Thomas (31) reported highest carbapenem resistance in age group between 21-40 years (36.25%). Overall *p*-value= .000857. In contrast the study by Pawar SK et al.[32] and Monika Saini et al [33] found highest percentage in age 41-50 years.

In our study the maximum organisms were isolated from urinary tract infection, followed by skin soft tissue infection, respiratory tract infection, septicemia and least nosocomial findings were from other hospital associated infections. Similar findings were from Urinary tract infection from [34, 35]

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243 Carbapenem resistance in *E.coli* and *Klebseilla pneumoniae* (21.5%) were highest among
244 the *Enterobacterials*. Least carbapenem resistance was seen among *Enterobacter species*
245 and *Proteus species* (5.26%). Carbapenem resistance among non- fermenter *Pseudomonas*
246 *aeruginosa* and *Acientobacter Species* were same about 15.7%. A similar study was found
247 which show *E.coli* and *Klebseilla pneumoniae* to be the commonest cause of
248 infection.[36,37] Incontrast the most prevalent bacteria reported were Klebsiella species, *A.*
249 *baumanii*, followed by *E. coli* and *P. aeruginosa* [17].There is a statistical significant
250 difference of $p < 0.05$ in the prevalence of gram negative bacteria and carbapenemase-
251 resistant isolates recovered from different types of specimens.

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253 Among the different departments the maximum carbapenem-resistance were present is
254 medicine 31.57% which was followed by surgery 26.31% , intensive care unit 21.05% and
255 obstetrics & gynecology and pediatric showing the same percentage of resistance 10.52%.
256 In contrast to our study it show the maximum number of isolates from the Surgery ward
257 22% followed by 19% medicine ward, 18% orthopedic ward, 15% MICU and 12% Pediatric
258 ward , 14% isolates obtained from other wards and ICUs. [38, 39]

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260 Among maximum carbapenem-resistant GNR were isolated from Urine 31.5% followed by
261 Endotracheal aspirate 21%, Pus 15.7% and 10.5% Blood, Sputum and other health
262 associate infection. A similar study by Lim et al. [40] it show high percentage of resistance in
263 urine 25.9%, lower respiratory tract 14.3% and blood 17%. In the study of Pano Pardo et al
264 [41] and Seibert et al. [42] show these bacteria are commonly isolated from bronchial
265 alveolar lavage (BAL), urine, and blood.

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267 Our study it shows the highest sensitivity of Colistin 89% and 3rd generation cephalosporine
268 show 100% resistance. A study showed that many carbapenemase producers are
269 susceptible in vitro to the glycylicycline group (Tigecycline), but there is rapid increase in
270 resistance to this drug [43]. Morrill et al [44], reported monotherapy is not effective against
271 infection caused by carbapenem-resistant bacteria.

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273 In our study the highest sensitivity was found in DDST 95% followed by mCIM 95%, , CDDT
274 84%, and MHT 74%, within CI 95% according to Naim H et al(45) also found CDDT
275 84.81%. In contrast the study by Naim H,et al [45] found sensitivity of MHT, and DDST as
276 97.41% and 84.81%. In a study by Cury et al. (46), it shows least percentage of 35.5% MHT.

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279 6. CONCLUSION

280 Our study illustrates the emergence of carbapenem-resistant Gram-negative hospital
281 acquired infection from patients. To identify the responsible agent that leads to infection in
282 healthcare settings. The common agents were *E.coli*, *Klebseilla pneumoniae*, *Citrobacter*
283 *species*, *Pseudomonas aeruginosa*, *Enterobacter species*, *Proteus species* and
284 *Acientobacter species*. The recovered isolates show the prevalence of 17.75% carbapenem
285 resistance to Imipenem, Ertapenem and Meropenem drugs. , the carbapenem resistance is
286 considered as the global alarm for pandemic resistance. As these carbapenem were the last
287 resorts to combat from multidrug-resistant organism. Warning signal of MDR, XDR and PDR
288 left with few drugs like Colistin and Tigecycline. In few studies we can see these drugs are
289 also showing resistance.

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292 COMPETING INTERESTS

293 Authors have declared that no competing interests exist.

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ETHICAL APPROVAL

The study was approved by Ethics committee MU/MM/BNS/2020/51(a)

REFERENCES

1. Queenan AM, Bush K. Carbapenemases: The versatile beta lactamases. *Clin Microbiol Rev.* 2007;20:4:40-58.
2. Livermore DM, Woodford N. Carbapenemases: A problem in waiting. *Curr Opin Microbiol.* 2000;3:489-95.
3. Nordmann P, Boulanger AE, Poirel L. NDM , Metallo β lactamase with increased carbapenemase activity from *Escherichia coli*. *Antimicrob Agents Chemother.* 2012;56:2184-6.
4. Tzouveleki LS, Markogiannakis A, Psychogiou M, Tassios PT, Daikos GL. Carbapenemases in *Klebsiella pneumoniae* and other Enterobacteriaceae: An evolving crisis of global dimensions. *Clin Microbiol Rev.* 2012;25:682-707.
5. Bourafa N., et al., Molecular characterization of carbapenem-resistant Gram-negative bacilli clinical isolates in Algeria. *Infection and drug resistance.*2018;11:735.
6. Karuniawati A., Saharman Y.R., and Lestari D.C., Detection of carbapenemase encoding genes in Enterobacteriaceae, *Pseudomonas aeruginosa*, and *Acinetobacter baumannii* isolated from patients at Intensive Care Unit Cipto Mangunkusumo Hospital in 2011 *Acta Med Indones.*2013;45(2):101–6.
7. Clinical and Laboratory Standards Institute. Performance Standards for Antimicrobial Susceptibility Testing; CLSI document M100-S28. Wayne PA: Clinical and Laboratory Standards Institute; 2018
8. Clinical and Laboratory Standards Institute. Performance Standards for Antimicrobial Susceptibility Testing; CLSI document M100-S28. Wayne PA: Clinical and Laboratory Standards Institute; 2018.
9. Pournaras S, Zarkotou O, Poulou A, Kristo I, Vrioni G, Themeli-Digalaki K, et al. A combined disk test for direct differentiation of carbapenemase-producing enterobacteriaceae in surveillance rectal swabs. *J Clin Microbiol.* 2013;51(9):2986-90.
10. Dheepa Muthusamy & Appalaraju Boppe. Phenotypic Methods for the detection of various Betalactamases in Carbapenemase resistant isolates of *A.baumannii* in a Tertiary care Hospital in a south India. *India. J of Clin Diagnostic Research.* 2012;6(6):970-73.
11. Haji SH, Aka STH, Ali FA. Prevalence and characterisation of carbapenemase encoding genes in multidrug-resistant Gram-negative bacilli. *PLoS One* 2021; 16: e0259005. *Microbiology and Applied Sciences* 5(12): 18–22.
12. Sreeja K. Vamsi,Rama S. Moorthy, Mary N. Hemiliamma, , Rama B. Chandra Reddy, Deepak J. chanderakant. Phenotypic and genotypic detection of carbapenemase production among gram negative bacteria isolated from hospital acquired infections. *Saudi Med J.* 2022; 43(3):236-243
13. Gupta E, Mohanty S, Sood S, et al. Emerging resistance to carbapenems in a tertiary care hospital in north India. *Indian J Med Res.*2006;124(1):95– 98.
14. Mulla S, Charan J, Panvala T. Antibiotic sensitivity of Enterobacteriaceae at a tertiary care center in India. *Chron Young Sci.* 2011;2:21:4-18.
15. Mate, P. H., Devi, K. S ., Devi, K. M., Damrolien, S., Devi, N. L., & Devi, P. P. Prevalence of Carbapenem Resistance among Gram - Negative Bacteria in a Tertiary Care Hospital in North - East India. *IOSR Journal of Dental and Medical Sciences.*2014;13(12), 56-60.

- 346 16. Makharita R.R., et al., Antibiogram and genetic characterization of carbapenem-
347 resistant gram-negative pathogens incriminated in healthcare-associated infections.
348 Infection and drug resistance.2020;3:3991.
- 349 17. Jalalvand K., et al., Evaluation of phenotypic and genotypic characteristics of
350 carbapnemases-producing enterobacteriaceae and its prevalence in a referral
351 hospital in Tehran city. Iranian journal of pathology.2020;15(2):86.
- 352 18. Shokri D., et al., Resistotyping, phenotyping and genotyping of New Delhi metallo- β -
353 lactamase (NDM) among Gram-negative bacilli from Iranian patients. Journal of
354 medical microbiology.2017;66(4):402–411.
- 355 19. Ain N.U., et al., High frequency and molecular epidemiology of metallo- β -lactamase-
356 producing gram negative bacilli in a tertiary care hospital in Lahore, Pakistan.
357 Antimicrobial Resistance & Infection Control.2018;7(1):1–9.
- 358 20. Jin C., et al., Molecular Characteristics of Carbapenem-Resistant *Enterobacter*
359 *cloacae* in a Tertiary Hospital in China. Infection and drug resistance.2020;13:1575.
- 360 21. Bourafa N., et al., Molecular characterization of carbapenem-resistant Gram-negative
361 bacilli clinical isolates in Algeria. Infection and drug resistance.2018;11:735.
- 362 22. Karabay O., et al., The carbapenem-resistant Enterobacteriaceae threat is growing:
363 NDM-1 epidemic at a training hospital in Turkey. Annals of clinical microbiology and
364 antimicrobials. 2016; 15(1):1–6.
- 365 23. Kollenda H., et al., Screening for carbapenemases in ertapenem-resistant
366 Enterobacteriaceae collected at a Tunisian hospital between 2014 and 2018.
367 European Journal of Microbiology and Immunology, 2019. 9(1): p. 9–13.
368 <https://doi.org/10.1556/1886.2018.00033>.
- 369 24. Mahrach Y., et al., Phenotypic and molecular study of carbapenemase-producing
370 Enterobacteriaceae in a regional hospital in northern Morocco. J Clin Med Sci.2019.
371 3: p. 113.
- 372 25. Codjoe, F.S., Detection and characterisation of carbapenem-resistant gram-negative
373 bacilli infections in Ghana. 2016, Sheffield Hallam University.
374 <http://shura.shu.ac.uk/id/eprint/15577>
- 375 26. Kaur MGS, Kaur T. Detection of carbapenem-resistant gram-negative bacteria in
376 clinical isolates from a tertiary care hospital. J Bacteriol Mycol Open Access.
377 2016;2(1):00011.33
- 378 27. Devi P. Incidence of carbapenem-resistant nonfermenting gram-negative bacilli from
379 patients with respiratory tract infections among intensive care units. Int J Res Med
380 Sci. 2017;3(6):4.
- 381 28. Mate PDK, Devi K, Damrolien S, Devi P. Prevalence of carbapenem resistance
382 among Gram-negative bacteria in a tertiary care hospital in north-east India. IOSR J
383 Dent Med Sci. 2014;13(12):56–60.
- 384 29. T.V. Parimala. (2017). Screening of Carbapenem Resistant Enterobacteriaceae
385 among Nosocomial Isolates: A Study from South India. Int.J.Curr.Microbiol.App.Sci
386 6(4): 460-465.
- 387 30. Satyajeet K Pawar et al. Carbapenem resistant Enterobacteriaceae: Prevalence and
388 bacteriological profile in a tertiary teaching hospital from rural western India. Indian J
389 Microbiol Res. 2018; 5(3): 342-347.
- 390 31. Namitha Thomas and Tarana Sarwat. 2019. Prevalence of Carbapenem Resistant
391 Enterobacteriaceae in A Tertiary Care Hospital. Int.J.Curr.Microbiol.App.Sci. 8(11):
392 1418-1424.
- 393 32. Pawar SK, Mohite ST, Shinde RV, Patil SR, Karande GS. Carbapenem-resistant
394 Enterobacteriaceae: Prevalence and bacteriological profile in a tertiary teaching
395 hospital from rural western India. Indian J Microbiol Res. 2018;5(3):342-347.
- 396 33. Monika Saini, Aditya Mishra, Sweta Gupta. Prevalence of Carbapenem Resistance in
397 Gram Negative Bacilli Isolates and Their Antimicrobial Susceptibility Pattern. Int J
398 Med Res Prof. 2016; 2(3):28-32.

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34. Deepak K Tempe, Jyotsna Agarwal, Kapil Chaudhary, Parin Lalwani, Madhu Sudan Tudu, Upendra Hansdah, Bibhavati Mishra. Carbapenem Resistance Patterns in General Intensive Care Unit of a Tertiary Care Hospital in India. *MAMC Journal of Medical Science*. 2015;1(2) : 85-91.
 35. Nair PK. Prevalence of carbapenem resistant Enterobacteriaceae from a tertiary care hospital in Mumbai, India. *J Microbiol Infect Dis*. 2013;03(04):207–10.
 36. Habibi S, Wig N, Agarwal S, Sharma S, Lodha R, Pandey R and Kapil A. Epidemiology of nosocomial infections in medicine intensive care unit at a tertiary care hospital in northern India. *Tropical Doctors*. 2008; 38: 233–5.
 37. Rimrang B., et al., Emergence of NDM-1-and IMP-14a-producing Enterobacteriaceae in Thailand. *Journal of antimicrobial chemotherapy*.2012;67(11): 2626–2630. .
 38. Dipak M. Panjwani, Sucheta J. Lakhani, Sanjay J. Mehta, Kunjan M. Kikani and Khushboo Madaan. A Study of Hospital Acquired Bacterial Infections and its Antimicrobial Susceptibility Pattern in a Teaching Hospital of Gujarat, India. *Int.J.Curr.Microbiol.App.Sci*. 2020;9(02): 1399-1408.
 39. Nagaraj S, Chandran SP, Shamanna P, Macaden R (2012). Carbapenem resistance among *Escherichia coli* and *Klebsiella pneumoniae* in a tertiary care hospital in south India. *Indian J Med Microbiol*.30(1): 93-95.
 40. Lim, Y.J.; Park, H.Y.; Lee, J.Y.; Kwak, S.H.; Kim, M.N.; Sung, H.; Kim, S.H.; Choi, S.H. Clearance of carbapenemase-producing Enterobacteriaceae (CPE) carriage: A comparative study of NDM-1 and KPC CPE. *Clin. Microbiol. Infect*. 2018, 24:1104.
 41. Pano Pardo, J.R.; Serrano Villar, S.; Ramos Ramos, J.C.; Pintado, V. Infections caused by carbapenemase-producing Enterobacteriaceae: Risk factors, clinical features and prognosis. *Enferm. Infec. Microbiol. Clin*. 2014;32 (Suppl. S4):41–48.
 42. Seibert, G.; Hörner, R.; Meneghetti, B.H.; Righi, R.A.; Dal Forno, N.L.; Salla, A. Nosocomial infections by *Klebsiella pneumoniae* carbapenemase producing enterobacteria in a teaching hospital. *Einstein*. 2014;12:282–286.
 43. Sader H., Farrell D., Flamm R., Jones R. Variation in potency and spectrum of tigecycline activity against bacterial strains from U.S. medical centers since its approval for clinical use (2006 to 2012). *Antimicrob Agents Chemother*.2014;58:2274–2280.
 44. Morrill HJ, Pogue JM, Kaye KS, LaPlante KL. Treatment Options for Carbapenem-Resistant Enterobacteriaceae Infections. *Open Forum Infect Dis*. 2015 May 5;2(2):050
 45. Naim H, Rizvi M, Gupta R, Azam M, Taneja N, Shukla I, et al. Drug resistance and molecular epidemiology of carbapenem resistant gram-negative bacilli isolates. *J Global Infect Dis*. 2018;10,13
 46. Cury AP, Andreazzi D, Maffucci M, Caiiffa Junior HH, Rossi F. The modified Hodge test is a useful tool for ruling out *Klebsiella pneumoniae* carbapenemase. *Clinics (Sao Paulo)* 2012.67,1427-31.