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## SUPPLEMENTARY MATERIAL

**Table 1.** Characteristics of included articles.

Publication year	n (%)
2023	5 (10.6)
2022	14 (29.8)
2021	9 (19.2)
2020	7 (14.9)
2019	12 (25.5)
<b>Journals</b>	
Antimicrobial Agents and Chemotherapy (1-6)	6 (12.8)
Antibiotics-Basel (7-10)	4 (8.5)
International Journal of Antimicrobial Agents (11-14)	4 (8.5)
Journal of Antimicrobial Chemotherapy (15-17)	3 (6.4)
Critical Care (18, 19)	2 (4.3)
Infection and Drug Resistance (20, 21)	2 (4.3)
Journal of Infection and Chemotherapy (22, 23)	2 (4.3)
Pharmaceutics (24, 25)	2 (4.3)
Acta Anaesthesiologica Scandinavica (26)	1 (2.1)
Antibiotics (27)	1 (2.1)
British Journal of Clinical Pharmacology (28)	1 (2.1)
Clinical Microbiology and Infection (29)	1 (2.1)
Drug Design, Development and Therapy (30)	1 (2.1)
European Journal of Clinical Pharmacology (31)	1 (2.1)
Frontiers in Pharmacology (32)	1 (2.1)
International Journal of Critical Illness and Injury Science (33)	1 (2.1)
International Journal of Infectious Diseases (34)	1 (2.1)
Jac-Antimicrobial Resistance (35)	1 (2.1)
Journal of Clinical Pharmacology (36)	1 (2.1)
Journal of Clinical Pharmacy and Therapeutics (37)	1 (2.1)
Journal of Critical Care (38)	1 (2.1)
Journal of Intensive Care (39)	1 (2.1)
Journal of Pharmacy Practice (40)	1 (2.1)
Journal of Trauma and Acute Care Surgery (41)	1 (2.1)
Medicina (Lithuania) (42)	1 (2.1)

Neurocritical Care (43)	1 (2.1)
Pharmacology Research and Perspectives (44)	1 (2.1)
Saudi Pharmaceutical Journal (45)	1 (2.1)
Surgical Infections (46)	1 (2.1)
Translational Andrology and Urology (47)	1 (2.1)
<b>Study Type</b>	
Retrospective observational	19 (40.4)
Prospective observational	13 (27.7)
Population pharmacokinetic	10 (21.3)
Randomized-controlled	3 (6.4)
Prospective pharmacokinetic	2 (4.3)
<b>Study Population</b>	
Critically ill	33 (70.2)
Hospital-acquired/Ventilator-associated pneumonia	3 (6.4)
Burn	3 (6.4)
Mixed	3 (6.4)
Cardiac surgery	1 (2.1)
Hemorrhagic stroke	1 (2.1)
Acute brain injury	1 (2.1)
Trauma	1 (2.1)
Sepsis	1 (2.1)
<b>Patient Characteristics</b>	
Total number of patients	6193 (100)
Median [IQR] number of patients per study	70 [32.5-167.5]
Range of number of patients per study	8-531
Male patients	3838 (62)
Female patients	2355 (38)
Range of mean/median age of the patients, years	29-88

## ARTICLES

- 1) Aréchiga-Alvarado NA, Medellín-Garibay SE, Milán-Segovia RDC, Ortiz-Álvarez A, Magaña-Aquino M, Romano-Moreno S. Population pharmacokinetics of amikacin administered once daily in patients with different renal functions. *Antimicrob Agents Chemother.* 2020;64(5):e02178-19. [\[CrossRef\]](#)
- 2) Carrié C, Delzor F, Roure S, Dubuisson V, Petit L, Molimard M, et al. Population pharmacokinetic study of the suitability of standard dosing regimens of amikacin in critically Ill patients with open-abdomen and negative-pressure wound therapy. *Antimicrob Agents Chemother.* 2020;64(4):e02098-19. [\[CrossRef\]](#)

- 3) Heffernan AJ, Sime FB, Kumta N, Wallis SC, McWhinney B, Ungerer J, et al. Multicenter population pharmacokinetic study of unbound ceftriaxone in critically Ill patients. *Antimicrob Agents Chemother*. 2022;66(6):e0218921. [\[CrossRef\]](#)
- 4) Kumta N, Heffernan AJ, Cotta MO, Wallis SC, Livermore A, Starr T, et al. Plasma and cerebrospinal fluid population pharmacokinetics of meropenem in neurocritical care patients: a prospective two-center study. *Antimicrob Agents Chemother*. 2022;66(8):e0014222. [\[CrossRef\]](#)
- 5) Layios N, Visée C, Mistretta V, Denooz R, Maes N, Descy J, et al. Modelled target attainment after temocillin treatment in severe pneumonia: Systemic and epithelial lining fluid pharmacokinetics of continuous versus intermittent infusions. *Antimicrob Agents Chemother*. 2022;66(3):e0205221. [\[CrossRef\]](#)
- 6) Ollivier J, Carrié C, d'Houdain N, Djabarouti S, Petit L, Xuereb F, et al. Are Standard dosing regimens of ceftriaxone adapted for critically Ill Patients with augmented creatinine clearance? *Antimicrob Agents Chemother*. 2019;63(3):e02134-18. [\[CrossRef\]](#)
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- 8) Scharf C, Paal M, Schroeder I, Vogeser M, Draenert R, Irlbeck M, et al. Therapeutic drug monitoring of meropenem and piperacillin in critical illness-experience and recommendations from one year in routine clinical practice. *Antibiotics (Basel)*. 2020;9(3):131. [\[CrossRef\]](#)
- 9) Zhao J, Fan Y, Yang M, Liang X, Wu J, Chen Y, et al. Association between augmented renal clearance and inadequate vancomycin pharmacokinetic/pharmacodynamic targets in Chinese adult patients: A prospective observational study. *Antibiotics (Basel)*. 2022;11(7):837. [\[CrossRef\]](#)
- 10) Zhao S, He N, Zhang Y, Wang C, Zhai S, Zhang C. Population pharmacokinetic modeling and dose optimization of vancomycin in Chinese patients with augmented renal clearance. *Antibiotics (Basel)*. 2021;10(10):1238. [\[CrossRef\]](#)
- 11) Nicolau DP, De Waele J, Kuti JL, Caro L, Larson KB, Yu B, et al. Pharmacokinetics and pharmacodynamics of ceftolozane/tazobactam in critically Ill patients with augmented renal clearance. *Int J Antimicrob Agents*. 2021;57(4):106299. [\[CrossRef\]](#)

- 12) Tournayre S, Mathieu O, Villiet M, Besnard N, Brunot V, Daubin D, et al. Factors associated with meropenem pharmacokinetic/pharmacodynamic target attainment in septic critically ill patients treated with extended intermittent infusion or continuous infusion. *Int J Antimicrob Agents*. 2023;62(2):106868. [\[CrossRef\]](#)
- 13) Vu DH, Nguyen DA, Delattre IK, Ho TT, Do HG, Pham HN, et al. Determination of optimal loading and maintenance doses for continuous infusion of vancomycin in critically ill patients: Population pharmacokinetic modelling and simulations for improved dosing schemes. *Int J Antimicrob Agents*. 2019;54(6):702-8. [\[CrossRef\]](#)
- 14) Yu Z, Liu J, Yu H, Zhou L, Zhao Y, Zhong L, et al. Should the trough concentration of vancomycin be abandoned in therapeutic drug monitoring? A multicentre, retrospective study of critically ill patients without any form of dialysis. *Int J Antimicrob Agents*. 2023;61(6):106812. [\[CrossRef\]](#)
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- 16) Dreesen E, Gijsen M, Elkayal O, Annaert P, Debaveye Y, Wauters J, et al. Ceftriaxone dosing based on the predicted probability of augmented renal clearance in critically ill patients with pneumonia. *J Antimicrob Chemother*. 2022;77(9):2479-88. [\[CrossRef\]](#)
- 17) Fratoni AJ, Mah JW, Nicolau DP, Kuti JL. Imipenem/cilastatin/relebactam pharmacokinetics in critically ill patients with augmented renal clearance. *J Antimicrob Chemother*. 2022;77(11):2992-9. [\[CrossRef\]](#)
- 18) Carrié C, Chadefaux G, Sauvage N, de Courson H, Petit L, Nouette-Gaulain K, et al. Increased β-Lactams dosing regimens improve clinical outcome in critically ill patients with augmented renal clearance treated for a first episode of hospital or ventilator-acquired pneumonia: a before and after study. *Crit Care*. 2019;23(1):379. [\[CrossRef\]](#)
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- 22) Ishigo T, Ibe Y, Fujii S, Kazuma S, Aigami T, Kashiwagi Y, et al. Effect of renal clearance on vancomycin area under the concentration-time curve deviations in critically ill patients. *J Infect Chemother.* 2023;29(8):769-77. [\[CrossRef\]](#)
- 23) Mikami R, Imai S, Hayakawa M, Sugawara M, Takekuma Y. Clinical applicability of urinary creatinine clearance for determining the initial dose of vancomycin in critically ill patients. *J Infect Chemother.* 2022;28(2):199-205. [\[CrossRef\]](#)
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- 25) Troisi C, Cojutti PG, Rinaldi M, Laici C, Siniscalchi A, Viale P, et al. Measuring creatinine clearance is the most accurate way for calculating the proper continuous infusion meropenem dose for empirical treatment of severe Gram-negative infections among critically Ill patients. *Pharmaceutics.* 2023;15(2):551. [\[CrossRef\]](#)
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- 37) Zhou Q, Zhao F, Wang M. An individualized administration model of vancomycin in elderly patients with sepsis and factors influencing augmented renal clearance. *J Clin Pharm Ther.* 2020. [\[CrossRef\]](#)
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- 40) Oswalt A, Joseph AC, Sima A, Kurczewski L. Evaluation of intravenous vancomycin pharmacokinetic parameters in patients with acute brain injury. *J Pharm Pract*. 2019;32(2):132-8. [[CrossRef](#)]
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